

CORRECTIONS/REVIEWS OF 58 NORTH AMERICAN BUTTERFLY BOOKS

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Abstract. Corrections are given for 58 North American butterfly books. Most of these books are recent. Misidentified figures mostly of adults, erroneous hostplants, and other mistakes are corrected in each book. Suggestions are made to improve future butterfly books. Identifications of figured specimens in Holland's 1931 & 1898 Butterfly Book & 1915 Butterfly Guide are corrected, and their type status clarified, and corrections are made to F. M. Brown's series of papers on Edwards' types (many figured by Holland), because some of Holland's 75 lectotype designations override lectotype specimens that were designated later, and several dozen Holland lectotype designations are added to the J. Pelham Catalogue. Type locality designations are corrected/defined here (some made by Brown, most by others), for numerous names: aenus, artonis, balder, bremnerii, brettoides, brucei (Oeneis), caespitatis, calanus, callina, carus, colon, colorado, coolinensis, comus, conquista, dacotah, damei, dumeti, edwardsii (Oarisma), elada, epixanthe, eunus, fulvia, furcae, garita, hermodur, kootenai, lagus, mejicanus, mormo, mormonia, nilus, nympa, oreas, oslari, philetas, phylace, pratincola, rhena, saga, scudderi, simius, taxiles, uhleri. Five first reviser actions are made (albihalos=austinorum, davenporti=pratti, latalinea=subaridum, maritima=texana [Cercyonis], ricei=calneva). The name c-argenteum is designated nomen oblitum, faunus a nomen protectum. Three taxa are demonstrated to be invalid nomina nuda (blackmorei, sulfuris, svilhae), and another nomen nudum (damei) is added to catalogues as a "schizophrenic taxon" in order to preserve stability. Problems caused by old scientific names and the time wasted on them are discussed.

Introduction

Many books have been published on butterflies. Unfortunately, most of the authors of those books do not get experts to review their work before they publish it, so mistakes go uncorrected. Other mistakes inevitably appear because of production errors, the advance of knowledge, simple lapses, inaccessibility or absence of experts, and every other conceivable reason. Some mistakes were published long ago and were never corrected, and were later repeated in nearly every book, because authors tend to repeat what prior books wrote, even if it was wrong. Many old erroneous hostplants have been repeated endlessly for 100 or even 200 years. And certain charismatic theories spring up, and are repeated endlessly, even if they are wrong (such as eastern N.A. *Lycaena phlaeas* being an introduction from Europe, or *Hesperia juba* adults overwintering).

This work examines butterfly books that are popular or recently published, and reviews them, and especially attempts to correct mistakes, in particular misdeterminations of illustrations of butterflies, and errors in their hostplants, behavior, distribution, etc. Sometimes I suggest names of species and subspecies that I believe to be correct instead of the ones used. Subspecies names are sometimes difficult to apply, and for some subspecies I have examined few--sometimes zero--examples, so that some of my opinions here are subjective and may change. Progress marches on, and the names of many of our butterflies will change because of new research, and because of the relentless historical trend of splitting of genera and subspecies and even species. But I do my best here to review these books and correct errors, to aid those butterfly fanciers who want to know the truth and are trying to find it.

I did not have the time to find all the mistakes in these books, so I undoubtedly missed very many or even most of them. I tried hardest to check the identification of illustrated butterflies, and hostplants. I seldom read verbal descriptions of adults or verbal descriptions of ranges.

I haven't reviewed most of the "Butterfly Watching" books, which have exploded in number lately. Those books have very little new information of use to scientist-lepidopterists, as most "butterfly watchers" do not get new hostplants or discover and report new scientific information in a usable manner, because their hands-off ideology causes uncertainty of identification, and they mostly lack the desire to contribute scientific discoveries. Some of the major watcher books are reviewed though.

A Study in Lectotypes: 1) THE BUTTERFLY BOOK. W. J. HOLLAND. Revised Edition, 1931. Doubleday & Co., NY. 424 p., 77 pl. 2) THE BUTTERFLY BOOK. W. J. HOLLAND. 1898. Doubleday & McClure Co., NY. 382 p., 48 pl. 3) THE BUTTERFLY GUIDE. W. J. HOLLAND. 1915. Doubleday, Page & Co., N.Y. 237 p. Correct Identifications of Specimens on Plates and Determination of Type Status of Illustrated Specimens in all Three Books.

The Butterfly Book, by W. J. Holland, was an important book for most of the 1900s. It was first published in 1898, and was completely revised with 29 new plates in 1931. Page ix claims that over 65,000 copies were sold between those dates!, an incredible amount if true. It continued to sell into the 1960s, an amazing longevity for butterfly books. This longevity was evidently due to the color plates which illustrated numerous butterflies in pleasing fashion, and the lack of competing books, because the text contained very little information and was replete with errors and speculations. The best thing about the book is the little stories Holland wrote scattered through the text (listed as “Digression and Quotations” on p. xii), which are fascinating and often humorous (see p. 113, etc.--I think Holland had greater talent as a fiction writer than as a scientist.) However, the 1931 plates are important from a historical perspective, because many of the illustrations are of type specimens named by Henry Skinner, Herman Strecker, Holland, etc., and especially including type specimens of butterflies named by William Henry Edwards in the Carnegie Museum, and from other museums. Holland paid for the publication of vol. 3 of Edwards’ *Butterflies of North America*, in exchange for Edwards’ collection. Holland then stored the collection in his home cellar--where many specimens molded--and eventually deposited it in the Carnegie Museum, where Holland was the Director.

This section attempts to correct the identifications of all the specimens on the plates, and properly determine their type status (as holotype, lectotype, paralectotype, paratype, syntype, not a syntype, etc.). Often I was aided by clues provided by F. Martin Brown in his series of papers on William Henry Edwards’ types (in *Trans. Amer. Ent. Soc.* vols. 90-113, from 1964 to 1987). Brown studied Edwards’ specimens, which are mostly in the Carnegie Museum after being purchased by Holland, so many of those Edwards specimens were used by Holland for the plates in his three books. Lionel Paul Grey corrected the *Argynnis* (*Speyeria*) identifications for me in the early 1960s. In some cases, an examination of the actual specimen would be necessary to be certain of the identification (in some of the *Erynnis*, an examination of the genitalia would be necessary), but I have done my best to identify them, and have given Holland the benefit of the doubt in those cases in which the specimen could be either the species Holland thought it was or another very similar species.

This project was undertaken along with a study of Brown’s papers concerning the types of butterflies named by William Henry Edwards (papers published in the *Trans. Amer. Ent. Soc.* from vol. 90 to 113)(some of the later papers were coauthored by Lee D. Miller and a few parts were authored by John C. Downey and Paul A. Opler & W. Reinthal). I found some errors and corrections in Brown’s papers, including about a dozen corrections of type localities, which are listed below after the corrections of Holland’s plates.

In identifying these specimens and redoing the plate legends, we encounter a big problem: the “types”. Many specimens in the plates were called “types” by Holland in the text or plate legends of his book, and many of those “types” are actually not types, and thus were called “pseudotypes” by Brown (here I call them “not syntypes”, because the word “pseudotype” is not used in the ICZN Code). This is rather technical. According to the Code, a “syntype” is a member of a type series only if a holotype was not designated [if a holotype was designated, then the other specimens should be called paratypes, not syntypes], and when a lectotype is designated from a syntype then the other syntypes become paralectotypes and can no longer be called syntypes. Thus my use of the word “syntype” here is sometimes a bit loose, as many of the specimens I have called syntypes are now properly called paralectotypes, although they may have been syntypes back in 1931 before a lectotype was designated. If a specimen was called a paratype, and a lectotype has been designated, I changed the wording to paralectotype. (I note that most lepidopterists still do not use the word paralectotype.) But other specimens are called paratypes, even though the Miller/Brown checklist states that only syntypes (and not a holotype) exist, which would mean that those “paratypes” are properly called syntypes, but I left those as “paratypes” while indicating that they may be syntypes due to the lack of a holotype; I worded them that way because of the uncertainty that many of those might not actually have been part of the original type series and thus may not be real syntypes or paratypes, and changing the word from paratype to syntype would imply a level of certainty that does not exist.

This problem among Edwards’ specimens and names—“types” that are really not types--arose because Edwards labeled his specimens poorly. He failed to place data labels on individual specimens, and instead merely placed a label at the head of a series of pinned specimens in a drawer. And Edwards would not label individual specimens as “types”. As a consequence, after Holland purchased the collection, Holland took labels off specimens that Holland thought looked “typical” and sent them to Edwards to have Edwards write the word “type” on them in red ink, and then Edwards sent the labels back to Holland, who placed them on the specimens. Sometimes Holland wrote a “type” label on specimens in Edwards’ collection, loosely based on correspondence from Edwards, and unfortunately sometimes Holland placed “type” labels on specimens that were not the real types. Edwards made other mistakes. Edwards sometimes forgot what he had named, and in one case he named the same species three times, having forgotten twice that he had already named it! His second description of *nokomis* wing pattern was of a much different ssp. than his original description. His handwriting

contained letters that were all highly slanted to the right, and all the letters in his handwriting looked rather similar, making his handwriting illegible to me (Brown loaned me a copy of Edwards' handwritten notebooks in the 1980s, but I could not read them)(maybe Edwards had trouble reading his own notes too?). All this sloppiness inevitably caused errors and uncertainty, which Brown attempted to unravel and correct.

I have attempted to indicate the correct "type" status of the specimens illustrated, at least for the names proposed by W. H. Edwards, as suggested by Brown's research. But for other authors (Herman Skinner, etc.), numerous specimens are called "type" on Holland's plate explanations or in his text (frequently, the text and plate explanations list different specimens as types), and I have no way of knowing whether most of those specimens are actual syntypes (one would have to study the actual specimens and their labels, etc.), so for names not authored by Edwards I have mostly just repeated Holland's designations of "types", some of which may actually not be syntypes. In some cases the text for the species provides clues as to the status of a "type". Page x of the 1931 book states that Holland borrowed many types from other museums, so many of these "types" on the plates may actually be syntypes. P. 337 says that Holland borrowed many Skinner types from Philadelphia Acad. Nat. Sci. P. 197 & 200 say that some figures of Gibson types were reproduced from the originals. Holland evidently used specimens from ANSP and USNM also. So perhaps half or more of the specimens that Holland called "types" are truly syntypes. John Rawlins told me that he has isolated and studied all the types of Hermann Strecker in the Field Museum, which will help when he publishes the study.

Another important point (a nest of worms, some would say) about Holland's "types" has come to light. Holland made no deliberate lectotype designations in his book (the word lectotype did not enter the ICZN code until 1958-1960). However, the ICZN Code (articles 74.5, 74.6) states that if an author singles out a single syntype specimen by using words such as "the type" to refer to it, that his words constitute a lectotype designation. Quoting the code: "74.5. Lectotype designations before 2000. In a lectotype designation made before 2000, either the term "lectotype", or an exact translation or equivalent expression (e.g. "the type"), must have been used or the author must have unambiguously selected a particular syntype to act as the unique name bearing type of the taxon. ..." This sentence provides two alternate ways to designate a lectotype, other than by using the term lectotype. So, in those cases in which Holland singled out just one specimen as the "type", and that specimen is a syntype, his action actually constitutes a valid lectotype designation (when he designates two or more specimens as type in text and plates, then the specimens may be syntypes but none are a lectotype designation). For example, Holland clearly designated the female on pl. LIX fig. 21 as the lectotype of *callina* (see his p. 131-2)(it is actually the holotype, see below). And Holland's lectotype designation of the female (of the pair of Boisduval syntypes) *sylvinus* on LXXVI fig. 15 trumps the later male lectotype (the male of the pair) of Emmel Emmel & Mattoon. Holland clearly indicates on pl. LXI fig. 28-30 that the fig. 28 youngi (*Erebia*) "type" is the lectotype, as he called the other two specimens allotype and paratype, thus the subsequent designation of the same male in fig. 28 as lectotype by dos Passos was unnecessary and invalid. Holland designated the *ivalda* female on LXII fig. 4 as lectotype, so the later designation of the male as lectotype is invalid. Holland designated the valid lectotype of *howardi* on p. 391, thus Gatrell's later lectotype is invalid. Holland named a number of butterflies himself, and evidently he almost never declared holotypes for his own names, and just called them types. Evidently, with Holland's sloppiness, even if he calls a figure "type" in text or plates of his book, that specimen should not be automatically considered to be a lectotype designation, because Brown showed that many of those specimens are not even syntypes (a lectotype cannot be designated from a non-syntype, that would be considered a neotype designation, and recent Codes say that neotypes must only be designated during revisionary studies). As an example, consider the messy case of *Coenonympha inornata*: Holland's text says that pl. XXV fig. 29 is "type", so we could take that as a lectotype designation, even though it is not called type on the plate; Brown called it a pseudotype, but he said the specimen was missing so obviously there was no evidence on the specimen's labels actually proving it was a pseudotype; however Brown did find another *inornata* specimen possessing Edwards' "type" label that matched the O.D. which Brown designated as lectotype, and Brown insisted fig. 29 was a pseudotype so maybe he thought it didn't match the O.D. well; and because the fig. 29 specimen is missing, the odds of it being a real syntype are maybe only 50% (judging by Holland's poor record of correctly labeling Edwards' specimens as "types"), so based on those miserable odds I cannot accept the missing fig. 29 bug as the lectotype. So, it will take more study of actual specimens to make sure they are real syntypes before being certain about all of Holland's lectotype designations. Of course, when Holland himself named a bug, I consider his "type" designations in this book to be correct designations of lectotype or holotype (the O.D. must be consulted to see whether he designated a holotype, which he seldom did).

dos Passos & Grey (1947, Amer. Mus. Novit. #1370 p. 2-3) did not use any of Holland's lectotype designations in their Catalogue of *Speyeria* (I find that Holland validly designated lectotypes of ten *Argynnis* [*Speyeria*]), because of sloppiness in Holland's book and the book's lack of precise terminology such as "lectotype designation", but they could not find any rule in the Code that says that Holland's lectotypes are invalid. The current Code also seems to indicate that Holland's lectotype designations are valid, and there is no rule stating that they are invalid. The Pelham Catalogue has listed 51 Holland lectotype designations and accepted them as correct, and I accept 75. We also have to realize that the word lectotype (and neotype etc.) did not even appear in the ICZN Code until 1958/1960!, even though by then it had been in usage in the literature for about 50 years! So sloppiness in usage of the various words for type is understandable, because precise terminology was not supported by the ICZN at that time. Working on those types involves slogging through the

old-name sewer, and dealing with imprecise papers, nonexistent and weak ICZN rules, the failure to properly label specimens, deliberate mislabeling of specimens (by Herbert Morrison most notably), shifting museum location of specimens, dermestid beetles and shaky hands causing specimen damage and loss, and human foibles at every turn. Perfection is only a dream in the Old-Name Sewer.

ICZN Code art. 74.1.3. indicates that the valid designation of a lectotype prevents any of the remaining syntypes (which are now paralectotypes) from being named as another lectotype under art. 74.1., which means that the original lectotype is valid even if it is now lost and a 2nd (invalid) lectotype was designated (this applies to *howardi*, *rosa*, *ines*). So if the original is lost, someone cannot designate another. Thus Holland's *ines* & *howardi* lectotypes are still valid even though probably lost. Valid Holland 1898 and 1915 and 1931 lectotypes therefore mean that later ones, such as dozens proposed by F. M. Brown in his papers on Edwards' types, are invalid. Brown still did valuable research however, including finding which specimens are syntypes and which are not, which helps us now decide whether any attempted lectotype designation is valid. And most of Brown's invalid lectotypes were the specimens that Holland already validly designated as lectotype. I indicate in the Figure Legends below the specimens that Holland called types but Brown decided were not syntypes (Brown called them "pseudotypes").

Summary of lectotypes. I started this Holland book project in 2005, using the Miller/Brown checklist for reference. The recent J. Pelham Catalogue of the Butterflies... (which I received April 10, 2008) has helped with some of the types. I found 75 lectotype designations by Holland in his three books. The Pelham Catalogue lists 51 of Holland's 1931 "type" designations (I accept 61) and 2 of Holland's 1898 "type" designations (I accept 8 different ones) and none of Holland 1915 "type" designations (I accept 6) as valid designations of lectotypes. I went through Holland 1931 and Holland 1898 and Holland 1915 again, using the Pelham Catalogue to help the effort, and found that I agreed with the Catalogue treatment of type status for most names, but I found a few discrepancies between the Catalogue and my conclusions. I and the Catalogue agree on 41 names that are Holland 1931 lectotype designations (*alaskensis* [Erebia], *apama*, *arcticus* [Papilio], *arizonensis* [Papilio], *atossa*, *attalus*, *behrii* [Colias], *bremneri*, *brettoides*, *brucei* [Papilio], *butleri*, *cassus*, *coyote*, *dion*, *dorus*, *drusius* [Thorybes], *erinna*, *gibsoni* [Oeneis], *hippolyta*, *hirsuta*, *hollandii*, *idaho*, *johnsoni*, *lena*, *macaria*, *melissa*, *moschus*, *mysie*, *nitra*, *opis*, *pallida* [Colias hecla], *pima*, *pittacus*, *rosa*, *silvius*, *siris*, *sterope*, *straton*, *texana* [Cercyonis], *ulrica*, *wheeleri*). Three names are considered to be Holland 1931 lectotype designations in Pelham Catalogue that are not valid designations (*aliaska*, *fulvia*, *hyantis*) because the specimen is not a syntype for two of these and was not singled out as a type in Holland's book for the third (there might be a few more of these in the Catalogue, as I have not examined every name in the Catalogue for mentions of Holland's book). I add 20 names that are valid Holland 1931 lectotype designations, which are not listed as such in the Catalogue (*brucei* [Oeneis], *byssus*, *cabelus*, *clara*, *colon*, *columbia*, *golovinus*, *hanhami*, *howardi*, *iduna*, *ines*, *ivallda* female, *liliana*, *mardon* [the Catalogue wrongly calls this a Holland 1898 lectotype designation], *morrisoni* (Stinga), *semplei*, *simius* [the Catalogue wrongly calls this a Holland 1898 lectotype designation], *sylvinus* female, *youngi* [Erebia], *yukonensis* [Coenonympha] [Pelham Catalogue mistakenly calls this specimen *yukonensis* holotype, but O.D. failed to designate a holotype]). I also add eight more names as Holland 1898 lectotype designations (*aemilia*, *barnesi*, *beani* [Euphydryas], *carolina* [Amblyscirtes], *duryi*, *ethela*, *itys*, *platina*), three of which Pelham wrongly considered to be a Holland 1931 designation (*aemilia*, *barnesi*, *duryi*). I add six more names as Holland 1915 lectotype designations (*atlantis*, *dymas*, *flora*, *macounii*, *perse*, *wittfeldi*), two of which Pelham wrongly considered to be Holland 1931 lectotype designations (*dymas*, *perse*). For these names, Holland singled out a single specimen in the book as "type". One name (*floridensis* [Eurytides]) would be a Holland 1898 lectotype designation, except the figured specimen is the holotype by monotypy; a second name (*pseudodorippus*) would be a Holland 1931 lectotype designation except it is an aberration and thus is surely the holotype by monotypy; and another name (*yehl*) would be a Holland 1898 lectotype designation except the specimen is the holotype in FMNH. The details of each lectotype designation by Holland are given below in the corrected plate legends of the book. For one of these names, *ines*, the Catalogue states that it is a valid Holland designation if the type can be found, but as noted above a lectotype designation remains valid even if the lectotype is lost. The names *thekla* & *inornata* cannot be proven to have Holland 1931 lectotypes because the specimens are lost, and the case of the name *wrighti* is too dubious to consider it to have a Holland 1931 lectotype. There are many other names that were singled out as type by Holland and thus would be Holland lectotype designations, except those taxa already have holotypes (usually the specimen Holland singled out as "type") (lectotypes can only be designated of syntypes, in order to take the place of a holotype that was never designated in the original description), including the names *oslari* (Phaeostrymon), *mojave*, *albicincta*, & *leussleri*, and notably including these five names: The *jamaicensis* [Pyrrhocalles] bug was stated to be the borrowed type on p. 392 thus it is evidently holotype (but if there was no holotype designated in OD, then this is Holland's lectotype). The *gunderiae* bug can't be a lectotype because it is an aberration that is a unique specimen that is holotype by monotypy according to O.D. The *callina* bug has been called a lectotype, but it is the holotype as Higgins wrote (see pl. LIX legend below). But the following bugs are messier. The *colorado* (Hesperia) bug (pl. LII fig. 1) is very doubtfully lectotype of *colorado* because the figure is H. *nevada*, so I think it is very doubtfully a syntype (though it could be, if the syntypes contained several species). The *calais* bug is called the holotype in Pelham Catalogue but Miller/Brown wrote that type may be lost (MCZ)? (the Pelham Catalogue is probably correct on this and most of these disagreements). If any of these prove not to have holotypes, as

might be possible in colorado & calais, then they will have Holland lectotypes. So, more research is needed in books and museums to conclusively fix some of these types. An examination of hundreds of original specimens in CM and other museums would be necessary to determine the correct “type” status of some of the names. People who do revisionary work should look at the original descriptions and try to look at the various types/syntypes.

Type localities. Art. 76.2. says that the type locality becomes the locality of the lectotype (76.3. says TL becomes locality of neotype), which means that the type localities of some of the bugs for which Holland designated lectotypes, invalidate the TLs of later-designated lectotypes, so will have to be changed to the locality of Holland’s lectotype. Most of the localities were rather vague, so there will not be many changes here, though there will be a few (the *Oeneis brucei* TL was changed a tiny bit; the platina & flora lectotypes should be examined for the locality label, as that locality overrides the locality of dos Passos & Grey’s and Brown’s later lectotypes; for hanhami, the specimen must be found and its exact locality in Man. or Minn. discovered). The *bremnerii* TL is now Vancouver.

The Butterfly Book first edition was published in 1898 (it was reprinted many times without change through 1929, then the revised 2nd edition appeared in 1931). The 48 plates of the first edition (pl. I-XLVIII) were identical to plates I-XLVIII of the revised 1931 book. But the explanations of the plates were a little different, as misidentifications in the 1898 book were corrected in 1931 and the nomenclature was brought up to date. The 1898 book has comparatively few mentions of the word type or types etc. (28 mentions of the word “type” including two on the explanation of plate XXX, of which eight are valid lectotype designations; 22 mentions of the word “types”; 1 mention of “paratype”; 2 of “paratypes”; 17 of “typical”; 0 mentions of other kinds of types; I carefully read through the book, and used the computer search feature on www.google.com books, to search the book for these words), which I studied to determine whether those are lectotype designations, with the following results. Holland 1898 made only eight valid lectotype designations: platina on p. 117, beani (*Euphydryas*) on p. 140, barnesi on p. 156, ethela on p. 211, duryi on p. 230, itys on p. 243, aemilia on p. 325, carolina (*Amblyscirtes*) on p. 367. Holland 1898 wrote that kodiak is type on p. 207, siva is type on Explanation of pl. XXX, alce is type on Explanation of pl. XXX, and myrtis is type on p. 346 (fig. pl XLVII fig. 11m), but these are not lectotype designations because Brown found that the specimens are not syntypes. Holland 1898 wrote on p. 359 that the fig. of yehl on pl. XLVI fig. 40m is the type, which would be a lectotype designation, except that there is a holotype (in FMNH). Holland 1898 did not make lectotype designations for mardon or simius as the Pelham Catalogue claimed, or for any other name.

The Butterfly Guide, a Pocket Manual for the Ready Identification of the Commoner Species Found in the United States and Canada, was authored in 1915 by Holland, and consisted of photos and brief text for the commoner species found in The Butterfly Book. This is the kind of small popular book we wish we could ignore, but we cannot ignore it, because it included 17 mentions of the word type and one mention of the word paratype, resulting in six valid lectotype designations, for the names atlantis, dymas, flora, macounii, perse, and wittfeldi. The photos in this book are mostly repeated from the Butterfly Book, but reversed left-to-right. P. 78 designates atlantis male lectotype on pl. XIV, which Holland 1931 fig. on pl. X fig. 9m & called a paratype; Brown called it a paratype and it is from Hunter NY thus dos Passos’ TL restriction was unnecessary. P. 86 designates dymas lectotype, a female on pl. XX fig. 4, which Holland 1931 fig. on pl. XVI fig. 18f. P.116 designates flora lectotype, a male on pl. L fig. 1, not fig. by Holland 1931, which is evidently one of the 2m1f syntypes Brown noted in CM. P. 142 designates macounii lectotype, a male on pl. LXXVI, which could be a reversed photo of Holland 1931 pl. XXVII fig. 3m, which Brown evidently treated as syntype, and which may be the male syntype that Brown discussed that has #5 on back of label; it is not the same specimen as Brown’s invalid lectotype. P. 85-86 designates perse lectotype, a male on pl. XX fig. 3, which Holland 1931 fig. on pl. XVI fig. 19m. P. 151 designates wittfeldi lectotype, a female on pl. LXXXI fig. 2, which Holland 1931 fig. on pl. XXIX fig. 19f, which Brown called the only female syntype; Brown designated an invalid male lectotype. Holland called other illustrated specimens type, which are not valid lectotype designations: The rhodope female and paulus male and silenus male are not syntypes according to Brown. The alicia male is not a syntype as it is from Fla. and the holotype and allotype fig. by Edwards were from New Orleans, and Brown found no syntypes in CM. The rubricata male is not a syntype and Brown designated a neotype. The charon male is not a syntype as it is not one of the 2m1f with type labels (Brown designated 1m of these lectotype, and Holland 1931 fig. the other pair on pl. LXIII figs. 24-25 which look too pale to be charon although the whole plate is too pale). The ridingsii male is not a syntype because Edwards named it from 4 females that were returned to James Ridings (Brown designated a neotype). The nais male is not a syntype, as the real male type was not in good condition in 1881 and was lost by 1968. There is no evidence that the ocola male is a syntype, as the specimen is labeled only “P. ocola Edw.” and “Butt. Guide/Pl. 143/Fig. 5”.

Misidentifications of photos in The Butterfly Guide: Pl. XXIV fig. 2 is *Chlosyne lacinia lacinia*. Pl. XXVII is *Polygonia faunus* male. XXXIV is *Vanessa virginiensis* female. XXXVI is *Junonia coenia* female. XLVII is *Limenitis lorquini* female. XLVIII is *Nymphalis californica* female. LXX is *Cercyonis pegala maritima*. LXXII fig. 1 is *Cercyonis meadii* female. LXXVIII fig. 3 is *Calephelis virginiensis* female. LXXXII fig. 2 is *Strymon melinus* male. LXXXIII fig. 2 is *Satyrium edwardsii* female. LXXXVI fig. 3 is *Callophrys polios* female. LXXXVI fig. 2 is *Callophrys henrici* male. LXXXVII fig. 1 is *Callophrys affinis perplexa* female. XCIII fig. 6 is *Brephidium isophthalma exilis* female. XCIV fig. 2 is *Echinargus thomasi* female. CI fig. 3 is *Euchloe hyantis lotta* female. CIX fig. 2 is *Eurema दौरa* male. CXXIX fig. 2 is

Urbanus proteus male. CXXXII fig. 1 is Achalarus lyciades male. CXXXVI fig. 2 is Erynnis horatius female. CXXXVII fig. 2 is E. horatius female. CXXXIX fig. 1 is Oarisma edwardsii male. CXXXIX fig. 3 is Oarisma garita female. CXLIX fig. 3 is Lerodea eufala female. CXLV fig. 2 is Euphyes vestris female. CXLVI fig. 4 is E. vestris female. CXLVII fig. 1 is Poanes aaroni=howardi male. CXLVII fig. 2 is Poanes viator male. CXLVII fig. 4 male & 5 female are Atrytone arogos male.

Correct Identifications and Type Status of Figured Specimens in the 1931/1898 Butterfly Book. In the corrected 1931/1898 plate legends below I note the reasons for the type decisions, and have tried to keep Brown's research visible, for instance by noting all the specimens figured by Holland that Brown discovered to be not syntypes, because Brown was the expert on Holland's types. I was aided in determining lectotype designations by the Miller/Brown checklist and Pelham Catalogue, which indicate whether there is a holotype or just syntypes, etc. Pelham's Catalogue attempted to list all the holotypes & lectotypes figured by Holland, and listed many of the paratypes & syntypes that Holland figured (some of which will no doubt be shown to not be real syntypes) but did not cite all the figs. of paratypes/syntypes.

I have tried to use original spellings of the species and subspecies names (Holland emended some names to paula, oeta, acasta, etc.) in the corrected Holland plate legends below. The Miller/Brown checklist cited the original genus name used in the O.D. for each name, but failed to list original spelling of species/ subspecies/ form names prior to their possible sexual transmogrification, but now we can find the original name in the Pelham Catalogue (at last we find that fuliginosa is the original name!).

Some people may want to make new legends for the plates in their copy of Holland's book, so I list the identification of every specimen on each plate, and put each plate explanation in a separate paragraph.

(Note: the name lupini is so grossly misleading and inappropriate and disgusting—implying that the bug has something to do with Lupinus, which it does not—that I have invoked the *lapsus contrarius* principle of the Biological Catalogue [see Papilio {New Series}#20], which adds the prefix a- or an- [meaning not-] to the front of such misleading names, in order to negate their inappropriateness; thus in this case alupini means “not-lupini”).

Abbreviations: m=male. f=female. CM=Carnegie Museum

Explanation of Plate I (frontispiece). 1. Vanessa cardui cardui m. 2. Vanessa virginiensis f. 3. Polygonia interrogationis m. 4. Colias philodice philodice m. 5. C. philodice philodice f. 6. Nymphalis antiopa f.

Explanation of Plate II (larvae). 1. Colias eurytheme. 2. Phoebis sennae eubule. 3. Eurema lisa. 4. Phoebis sennae eubule. 5. Anthocharis midea midea. 6. Eurema nicippe. 7. Pontia protodice. 8-9. Pieris oleracea. 10. Colias philodice philodice. 11-12. Pieris rapae. 13. Battus philenor philenor. 14. Eurytides marcellus. 15. Papilio glaucus glaucus prepupa. 16. Papilio crespontes. 17. Papilio polyxenes asterias 2nd stage. 18. Papilio troilus. 19. Papilio troilus 3rd stage. 20. Battus philenor philenor. 21. B. p. philenor 3rd stage. 22. Papilio troilus 3rd stage. 23. Achalarus lyciades. 24. Papilio polyxenes asterias 4th stage. 25. Thorybes pylades?. 26. Papilio glaucus glaucus. 27. Papilio polyxenes asterias. 28. Papilio glaucus glaucus. 29. Thorybes pylades. 30-31. Epargyreus clarus (should be green). 32. Thorybes bathyllus. 33. Epargyreus clarus. 34. Urbanus proteus. 35. Epargyreus clarus 3rd stage.

Explanation of Plate III (larvae). 1-2. Oeneis melissa semidea. 3. Megisto cymela. 4. O. melissa semidea. 5. Danaus plexippus. 6. Megisto cymela. 7. Oeneis melissa semidea 1st stage. 8. Neonympha areolata. 9. Lethe eurydice eurydice. 10. Megisto cymela. 11. Oeneis jutta 1st stage. 12. Neonympha areolata. 13-14. Megisto cymela. 15. Oeneis melissa semidea. 16. Lethe portlandia. 17. Limenitis arthemis astyanax. 18. Cercyonis pegala alope. 19. Limenitis archippus archippus. 20. Asterocampa clyton (horns too large). 21. Limenitis arthemis astyanax. 22. Limenitis a. archippus. 23. Polygonia interrogationis. 24. Limenitis a. archippus. 25. L. arth. astyanax. 26. L. arth. arthemis. 27. P. interrogationis? (comma?) 28. Nymphalis antiopa. 29-30. Precis coenia. 31. Polygonia progne. 32. Polygonia faunus. 33. Polygonia satyrus. 34. Vanessa virginiensis. 35. Vanessa atalanta. 36. Aglais milberti. 37. Vanessa cardui cardui. 38. Polygonia comma.

Explanation of Plate IV (pupae). 1-3. Danaus plexippus. 4-5. Oeneis melissa semidea. 6. Lethe portlandia. 7-8. Cercyonis pegala nephele. 9. Lethe eurydice eurydice. 10-11. Neonympha areolata. 12-13. Limenitis arthemis astyanax. 14. Limenitis arthemis arthemis. 15-17. Asterocampa clyton clyton. 18-20. Limenitis archippus archippus. 21-22. Polygonia interrogationis. 23. Limenitis arthemis arthemis. 24-26. Polygonia interrogationis. 27. Polygonia comma. 28. Megisto cymela. 29-30. Polygonia comma. 31. Polygonia faunus faunus. 32. Polygonia progne. 33-35. Polygonia faunus faunus. 36. Roddia l-album j-album. 37-38. Polygonia progne. 39. Polygonia comma. 40. Polygonia interrogationis. 41-42. Polygonia satyrus. 43. Aglais milberti. 44-45. Roddia l-album j-album. 46-48. Polygonia comma. 49-50. Aglais milberti. 51. Nymphalis antiopa. 52-53. Vanessa atalanta. 54. Vanessa virginiensis. 55. Vanessa atalanta. 56-57. Junonia coenia. 58-59. Nymphalis antiopa. 60-62. Vanessa cardui cardui. 63-64. Vanessa virginiensis. 65-66. Junonia coenia. 67. Junonia coenia?

Explanation of Plate V (pupae). 1-3. Argynnis (Speyeria) cybele cybele. 4. Argynnis (Speyeria) idalia. 5. Argynnis (Speyeria) aphrodite aphrodite. 6. Argynnis (Speyeria) atlantis atlantis. 7. Euphydryas phaeton. 8-9. Euptoieta claudia. 10-11. Boloria bellona. 12-14. Boloria selene myrina. 15-16. Euphydryas phaeton. 17-18. Chlosyne harrisii. 19. Chlosyne nycteis nycteis. 20-22. Phyciodes tharos or P. cocyta. 23-24. Libytheana carinenta bachmanii. 25. Satyrium calanus falacer. 26. Callophrys irus irus. 27. Satyrium calanus falacer. 28. Satyrium liparops strigosa. 29. Satyrium

edwardsii. 30-31. *Callophrys gryneus gryneus*. 32-34. *Callophrys irus irus*. 35. *Satyrium acadica*. 36. *Celastrina* sp. 37. *Satyrium titus titus*. 38. *Callophrys niphon*. 39. *Strymon melinus*. 40. *Callophrys niphon*. 41. *Plebejus idas scudderi*. 42. *Cupido comyntas comyntas*. 43-44. *Celastrina neglecta*. 45-46. *Feniseca tarquinius*. 47-48. *Cupido comyntas comyntas*. 49. *Lycaena phlaeas hypophlaeas*. 50. *Lycaena thoe*. 51-52. *Eurema nicippe*. 53. *Colias eurytheme*. 54-55. *Colias philodice philodice*. 56. *Eurema lisa*. 57. *Pieris oleracea*. 58. *Pieris rapae*. 59. *Anthocharis midea midea*. 60-62. *Phoebis sennae eubule*. 63-64. *Pieris oleracea*. 65. *Pieris rapae*. 66-67. *Pontia protodice*.

Explanation of Plate VI (pupae). 1-4. *Papilio glaucus glaucus*. 5-7. *Papilio troilus troilus*. 8-10. *Papilio crespontes*. 11-12. *Eurytides marcellus*. 13. *Papilio polyxenes asterias*. 14-17. *Battus philenor philenor*. 18-19. *Papilio polyxenes asterias*. 20. *Battus philenor philenor*. 21. *Achalarus lyciades*. 22. *Epargyreus clarus*. 23. *Urbanus proteus*. 24. *Thorybes bathyllus*. 25-26. *Epargyreus clarus*. 27. *Erynnis icelus*. 28. *Thorybes pylades*. 29. *Pholisora catullus*. 30-32. *Erynnis lucilius*. 33. *Erynnis juvenalis*. 34. *Erynnis persius*. 35. *Pyrgus communis*. 36. *Pholisora catullus*. 37. *Erynnis martialis*. 38. *Erynnis brizo*. 39. *Hylephila phyleus phyleus*. 40. *Amblyscirtes vialis*. 41. *Pholisora catullus*. 42. *Wallengrenia otho?* (J. Calhoun 2006 J. Lep. Soc. 60:8 says Abbot's otho may not be *Wallengrenia*). 43. *Atalopedes campestris*. 44. *Polites themistocles*. 45. *Amblyscirtes hegon*. 46. *Lerema accius*. 47. *Atalopedes campestris*. 48. *Calpodus ethlius*.

Explanation of Plate VII. 1. *Danaus plexippus plexippus* m. 2. *Danaus gilippus berenice* m. 3. *Danaus gilippus thersippus* m. 4. *Limenitis archippus archippus* m. 5. *Limenitis archippus obsoleta=hulstii* holotype m.

Explanation of Plate VIII. 1. *Dircenna klugi* f. 2. *Mechanitis polymnia* (?) m. 3. *Ithomia anaphissa* (?) m. 4. *Dryas iulia delila* (Jamaica) m. 5. *Heliconius charithonia* ssp. m. 6. *Dryas iulia iulia* (evidently Puerto Rico) m. 7. *Dione vanillae* (perhaps ssp. *nigrior*) m. 8. *Euptoieta hegesia*. 9. *Euptoieta claudia* m.

Explanation of Plate IX. 1-2. *Argynnis* (*Speyeria*) *diana*, m, f. 3-4. *Argynnis* (*Speyeria*) *cybele cybele* (near *krautwurmi*?), m, f. 5-6. *Argynnis* (*Speyeria*) *cybele leto*, neotype m, f.

Explanation of Plate X. 1-2. *Argynnis* (*Speyeria*) *nokomis apacheana*, m, f. 3. *Argynnis* (*Speyeria*) *idalia* (evidently weak ssp. *occidentalis*) f. 4. *Argynnis* (*Speyeria*) *callippe nevadensis* (not a syntype) m. 5. *Argynnis* (*Speyeria*) *mormonia mormonia* m. 6. *Argynnis* (*Speyeria*) *aphrodite alcestis* (not a syntype) m. 7. *Argynnis* (*Speyeria*) *zerene bremnerii* (a lectotype designation by Holland 1931) lectotype m. 8. *Argynnis* (*Speyeria*) *aphrodite ethne* (not a syntype of *electa* which is a ssp. of *S. hesperis*) m. 9. *Argynnis* (*Speyeria*) *atlantis atlantis* lectotype (called paratype on p. 90 but validly designated lectotype by Holland 1915 pl. XIV) m (Hunter, NY).

Explanation of Plate XI. 1-2. *Argynnis* (*Speyeria*) *callippe callippe*, m, f. 3. *Argynnis* (*Speyeria*) *callippe* ssp. (*semivirida?*) m. 4-5. *Argynnis* (*Speyeria*) *edwardsii*, m, f. 6. *Argynnis* (*Speyeria*) *hydaspe rhodope* f (not a syntype). 7. *Argynnis* (*Speyeria*) *mormonia bischoffi* (not a syntype as Brown wrote on p. 316, as 320 says both syntypes were destroyed) m. 8. *Argynnis* (*Speyeria*) *hesperis electa=cornelia* (may have been a syntype of *cornelia*, but now lost) m. 9. *Argynnis* (*Speyeria*) *hesperis nausicaa* (not a syntype) m. 10-11. *Argynnis* (*Speyeria*) *coronis coronis*, m, f.

Explanation of Plate XII. 1-2. *Argynnis* (*Speyeria*) *hesperis hesperis* (not syntypes), m, f. 3-4. *Argynnis* (*Speyeria*) *aphrodite ethne=cypris* (text says both specimens are types, thus neither is Holland's lectotype designation), lectotype m, paralectotype f. 5-6. *Argynnis* (*Speyeria*) *egleis oweni* (text says both specimens are types, thus neither is Holland's lectotype designation), lectotype m, paralectotype m. 7. *Argynnis* (*Speyeria*) *mormonia eurynome* m. 8-9. *Argynnis* (*Speyeria*) *callippe rupestris* 2m. 10. *Argynnis* (*Speyeria*) *zerene hippolyta* lectotype (designated lectotype by Holland 1931 on p. 93) m. 11-12. *Argynnis* (*Speyeria*) *callippe juba=sierra=laura*, m lectotype of *laura*, f. 13. *Argynnis* (*Speyeria*) *mormonia artonis* (not a syntype) m.

Explanation of Plate XIII. 1. *Argynnis* (*Speyeria*) *cybele cybele* m. 2-3. *Argynnis* (*Speyeria*) *coronis semiramis*, m (may be paralectotype), f (not a syntype). 4. *Argynnis* (*Speyeria*) *nokomis nitocris* holotype m. 5-6. *Argynnis* (*Speyeria*) *coronis halcyone* (not syntypes) 2m. 7-8. *Argynnis* (*Speyeria*) *zerene zerene=monticola* 2m. 9. *Argynnis* (*Speyeria*) *callippe macaria* lectotype (text p. 98 says type, so designated lectotype by Holland 1931) m. 10. *Argynnis* (*Speyeria*) *callippe juba=laura=sierra* (unsilvered var.) (not *inornata*, which is a syn. of *rupestris*) (not a syntype of *inornata*) f. 11. *Argynnis* (*Speyeria*) *callippe liliana* (text p. 98 says "type", a Holland 1931 lectotype designation; Pelham Catalogue says that this specimen is possibly a fig. of the dos Passos & Grey lectotype, which is in AMNH, but dos Passos & Grey don't mention that; it is not a lectotype if further research shows that it is not a syntype) m. 12. *Argynnis* (*Speyeria*) *adiaste atossa* lectotype (designated lectotype on p. 100 by Holland 1931; it has no locality but allotype locality is Tehachapi [Mts.] which is TL) m. 13-15. *Argynnis* (*Speyeria*) *egleis egleis*, m, m, f.

Explanation of Plate XIV. 1-2. *Argynnis* (*Speyeria*) *callippe meadii* (not syntypes) 2m. 3. *Argynnis* (*Speyeria*) *aphrodite* lectotype (syntype, called "type" in text p. 88 thus Holland 1931 designated this specimen as lectotype) lectotype m. 4. *Argynnis* (*Speyeria*) *adiaste adiaste* m. 5-7. *Argynnis* (*Speyeria*) *mormonia eurynome=clio* (not *clio* syntypes), m, f, m. 8. *Argynnis* (*Speyeria*) *mormonia opis* lectotype (designated lectotype by Holland 1931 on p. 103) m. 9. *Argynnis* (*Speyeria*) *hydaspe hydaspe* m. 10. *Argynnis* (*Speyeria*) *zerene behrensii* (not a syntype) m. 11. *Argynnis* (*Speyeria*) *aphrodite alcestis* f. 12-13. *Argynnis* (*Speyeria*) *hesperis dennisi=lais*, m (not a syntype), f syntype. 14-15. *Argynnis* (*Speyeria*) *mormonia eurynome*, f, m. 16. *Argynnis* (*Speyeria*) *hesperis wasatchia* (not *chitone*) (not a *chitone* syntype) f. 17. *Argynnis* (*Speyeria*) *zerene zerene=monticola* f.

Explanation of Plate XV. 1-2. *Boloria selene myrina* 2m. 3. *Boloria eunomia caelestis=alticola* m. 4-6. *Boloria titania grandis*, m, m, f. 7-8. *Boloria titania montinus*, m, f. 9-10. *Boloria freija freija* (Alta. phenotype), m, f. 11-12. *Boloria freija freija* 2m. 13-14. *Boloria frigga saga*, m, f. 15. *Boloria alberta* lectotype m. 16. *Boloria bellona* m. 17. *Boloria frigga sagata* (not an epithore syntype) m. 18. *Boloria epithore* (not a syntype) m.

Explanation of Plate XVI. 1. *Euphydryas phaeton phaeton* m. 2. *Euphydryas chalcedona chalcedona* m. 3. *Euphydryas chalcedona macglashanii* m. 4. *Euphydryas editha quino=augusta* (not a syntype) m. 5. *Euphydryas chalcedona colon* lectotype (designated lectotype by Holland 1931 on p. 117; TL is required to be locality of lectotype which is “Columbia River”) m. 6. *Euphydryas anicia capella* Colo. m. 7. *Euphydryas chalcedona chalcedona* (not a baroni syntype) m. 8. *Euphydryas editha editha* f. 9. *Euphydryas anicia capella* Colo. m. 10. *Euphydryas editha rubicunda* (text says “type” so perhaps a syntype and if so would be a Holland 1931 lectotype designation; but male is different from Gunder’s 1929 Pan-Pac. Ent. 6[1]:pl. 10m fig. of “type”; Gunder’s specimen would seem to be the valid lectotype designation) m. 11-12. *Chlosyne sterope acastus* (p. 126 says 3 figs. are types, so none is a Holland lectotype designation), m lectotype, m (probable paralectotype, but now lost). 13-14. *Chlosyne palla palla* 2m. 15. *Chlosyne gabbii* m. 16. *Euphydryas editha taylori* paralectotype m. 17. *Chlosyne leanira fulvia* (not a syntype) m. 18. *Microtia (Dymasia) dymas dymas* lectotype (designated lectotype by Holland 1915 pl. XX fig. 4, later designated lectotype by Holland 1931 on p. 132) f. 19. *Microtia (Texola) elada perse* lectotype (designated lectotype by Holland 1915 pl. XX fig. 3, later designated lectotype by Holland 1931 on p. 132) m. 20. *Chlosyne leanira wrighti* m. 21. *Poladryas minuta nympha* (text says “types” for both this and LVIII fig. 14, so not a lectotype designation by Holland 1931; Holland 1898 made no mention of type in text or plate, so did not designate lectotype either. Brown declared this male lectotype.) lectotype m. 22. *Poladryas minuta minuta aberrant* f.

Explanation of Plate XVII. 1. *Chlosyne leanira alma* m. 2. *Microtia (Texola) elada ulrica* lectotype (designated lectotype by Holland 1931 on p. 132) m. 3-4. *Microtia (Dymasia) dymas chara* (probably paralectotypes but now lost) 2m. 5-6. *Chlosyne harrisii harrisii*, m, f. 7-8. *Chlosyne palla altasierra* 2m. 9-10. *Chlosyne leanira wrighti* (text says both are types so this is not a Holland lectotype designation), lectotype m, allotype (properly paralectotype) f. 11-12. *Poladryas minuta arachne* 2m. 13. *Chlosyne hoffmanni hoffmanni* m. 14. *Chlosyne hoffmanni hoffmanni ab. hollandae* holotype f. 15-16. *Chlosyne theona thekla*, m (Ariz., coll. H. Morrison, may be syntype), m (probable syntype, called “type” in text, now missing, thus if it was a valid syntype it was Holland’s designation of lectotype on p. 130, even though now lost [Art. 74.4], but I will ignore the possibility because maybe it wasn’t a syntype and being lost it cannot be proven to be a syntype). 17-19. *Phyciodes graphica graphica* (not *vesta* syntypes), m, f, spring form f. 20-21. *Phyciodes picta picta*, f (not a syntype), m (probably not syntype, now missing). 22-23. *Phyciodes phaon jalapeno* (Texas), m, f (not a syntype). 24-25. *Chlosyne gorgone* 2m. 26-27. *Phyciodes pulchella montana* 2f. 28-30. *Chlosyne nycteis nycteis*, m, m, f. 31. *Phyciodes orseis orseis* (not a syntype) m. 32-34. *Phyciodes pulchella camillus* (not syntypes), m, f, spring form m. 35-36. *Phyciodes batesii batesii* 2m. 37. *Phyciodes pulchella camillus* m. 38. *Phyciodes mylitta mylitta* spring form. 39. *Phyciodes (Eresia) frisia tulcis=punctata* holotype m. 40-41. *Phyciodes mylitta arizonensis* (not *mylitta* syntypes) 2m. 42. *Phyciodes (Eresia) frisia*, looks like ssp. *dubia* of Venez.-Col. m.

Explanation of Plate XVIII. 1. *Phyciodes cocyta selenis* m. 2. *P. tharos tharos* probably as antenna club is oval (or *Phyciodes cocyta selenis*, as uph has large space) m. 3-4. *Phyciodes tharos tharos* probably as antenna club oval (may be *marcia* syntypes, but specimens now missing) 2m. 5. *Phyciodes pallida barnesi* (p. 138 says pl. XVIII fig. 5m is type, a Holland 1931 lectotype designation; Miller/Brown say holotype is in CM, but Pelham Catalogue says Holland 1931 p. 138 designated this specimen as lectotype, so evidently there was no holotype and this is lectotype; however Holland 1898 p. 156 says this male pl. XVIII fig. 5m is the type, so this is a valid Holland 1898 lectotype designation)(p. 138 says in error “Colorado Springs” where *P. pallida* does not occur) lectotype m. 6. *Argynnis (Speyeria) coronis snyderi* (text says paratype) m. 7. *Argynnis (Speyeria) zerene platina* (p. 97 says paratype, so it must be a syntype, as Pelham Catalogue says dos Passos & Grey designated lectotype in 1947 so there was no holotype; Holland 1898 p. 117 wrote that this pl. XVIII fig. 7m specimen is the type, therefore it is a valid Holland 1898 lectotype designation) lectotype m. 8-9. *Phyciodes (Anthanassa) texana texana* m (paratype?, now missing), holotype m. 10. *Chlosyne lacinia lacinia* (or *adjutrix*) m. 11. *Chlosyne lacinia adjutrix* m. 12. *Phyciodes (Anthanassa)* looks like *ardys subota* from Guat. m (*ianthe* is a syn. of “*Janatella*” *hera*, but fig. isn’t that). 13. *Euphydryas editha beani* (p. 118 says “type”, so this is a Holland 1931 lectotype designation; Miller/Brown say holotype is in CM, but Pelham Catalogue says the lectotype was designated by Gunder 1929 Pan-Pac. Ent. 6(1) pl. 11 in ANSP, which is older, thus Holland’s 1931 designation is invalid; Pelham Catalogue says this fig. 13 specimen is Gunder’s type, but it looks different than Gunder’s so isn’t lectotype and is probably paralectotype; however, Holland 1898 p. 141 says that this pl. XVIII fig. 13m is Dr. Skinner’s original type, which is the valid Holland 1898 lectotype designation and Gunder’s is invalid) lectotype m. 14-15. *Boloria tritonia astarte*, f, m (holotype of *victoria*). 16-17. *Boloria titania helena* (text says both are types, so not a Holland lectotype designation), m (not a syntype), m (possible syntype). 18. *Lethe creola* m. 19. *Lethe portlandia missarkae* (not *creola*) f. 20. *Lethe anthedon* m. 21. *Gyrocheilus patrobas tritonia* m.

Explanation of Plate XIX. 1. *Polygonia interrogationis* spring form *fabricii* m. 2. *Polygonia interrogationis* summer form *umbrosa* f. 3. *Polygonia comma* summer form *dryas* m. 4. *Polygonia comma* spring form *harrisii* m. 5-6. *Polygonia*

oreas silenus, m (not a syntype), f. 7. *Polygonia faunus hylas* (S Colo., not a syntype) m. 8. *Polygonia faunus hylas* (possible hylas syntype) form *silvius=orpheus* Colo. f. 9. *Roddia l-album j-album* f. 10-11. *Polygonia gracilis gracilis* 2m. 12. *Polygonia faunus faunus* (may be topotype Hunter, Green Co. NY) m. 13. *Polygonia faunus smythi* W. Va. m. 14-15. *Polygonia satyrus satyrus* (not the European marsyas) 2m (both now missing).

Explanation of Plate XX. 1-2. *Polygonia satyrus satyrus* 2m. 3-4. *Polygonia progne*, winter form, m, m. 5-6. *Polygonia gracilis zephyrus*, m, syntype m. 7. *Junonia coenia* f. 8. *Junonia genoveva hubneri* form *pallescens* m. 9. *Junonia evarete zonalis* m. 10. *Aglais milberti* m. 11. *Nymphalis californica* m. 12. *Vanessa carye annabella* f. 13. *Anartia jatrophae guantanamo* m.

Explanation of Plate XXI. 1-2. *Marpesia zerynthia=coresia* 2m. 3. *Marpesia petreus* m. 4. *Marpesia chiron* m. 5-6. *Diaethria clymena* 2m. 7-8. *Eunica monima*, m, f. 9-10. *Hypolimnias misippus*, m, f.

Explanation of Plate XXII. 1. *Limenitis arthemis astyanax* m. 2. *Adelpha bredowii californica* f. 3. *Limenitis lorquini lorquini* m. 4. *Limenitis arthemis arthemis* m. 5. *Limenitis arthemis arthemis X astyanax* (f. *proserpina*) m. 6. *Limenitis weidemeyerii latifascia* m.

Explanation of Plate XXIII. 1-2. *Asterocampa clyton flora*, lectotype m, f. 3-4. *Asterocampa celtis celtis*, m, f. 5. *Asterocampa clyton clyton* form *proserpina* (Scott's form *geneumbrosa*) m. 6. *Asterocampa clyton clyton* f. 7-8. *Asterocampa celtis celtis X montis* Colo., m, f. 9-10. *Asterocampa celtis reinthali=alicia*, m (Florida; not a syntype; too tawny, the same male in *The Butterfly Guide* is browner like *celtis*), f. 11. *Asterocampa celtis montis* lectotype (tawny, but atypical ocelli resembling *antonia*) m. 12. *Asterocampa celtis antonia* lectotype m. 13. *Asterocampa celtis celtis* m.

Explanation of Plate XXIV. 1. *Anaea andria* f. 2. *Anaea aidea morrisonii* syntype f. 3. *Anaea troglodyta floridensis* m. 4. *Hamadryas feronia* m. 5. *Hamadryas fornax* m. 6. *Siproeta stelenes biplagiata* m. 7. *Mestra dorcas amymone* m. 8-9. *Chlosyne lacinia crocale*, f, m. 10. *Hypanartia lethe* m.

Explanation of Plate XXV. 1. *Lethe eurydice eurydice* m. 2. *Cyllopsis gemma* m. 3. *Megisto rubricata cheneyorum* (not a *rubricata* syntype) m. 4. *Megisto cymela* (or *Megisto eurytris* Maynard 1891?) f. 5. *Hermeuptychia hermes sosybia* m. 6. *Neonympha mitchellii* m. 7. *Neonympha areolata areolata=phocion* m. 8. *Cyllopsis pertepida dorothea=* form *edwardsii* (Trans. Amer. Ent. Soc. 90:331) f. 9. *Coenonympha tullia californica=galactinus* m. 10. *Coenonympha tullia californica=eryngii* m. 11. *Coenonympha tullia ochracea* m. 12. *Coenonympha tullia* (looks like ssp. *pseudobrenda*) m. 13. *Coenonympha tullia inornata* m. 14. *Coenonympha tullia californica* f. 15. *Neominois ridingsii ridingsii* m (not a syntype). 16. *Neominois ridingsii stretchii=dionysus* m. 17. *Erebia magdalena magdalena* m. 18. *Erebia stubbendorffii ethela* (p. 205 says two specimens are type, so this is not a Holland 1931 lectotype designation; however, Holland 1898 p. 211 says "The figure in the plate [XXV fig. 18f] is that of the female type of Edwards' ethela, ethela being a synonym for sofia.", and Brown stated that this female is a syntype, therefore this female is a Holland 1898 lectotype designation) lectotype f. 19. *Erebia discoidalis* m. 20. *Erebia callias syntype* m (Brown stated he could not find original specimen matching this, so it may be missing). 21. *Coenonympha tullia insulanus* Vancouver Is., m. 22. *Coenonympha tullia kodiak* (not a syntype) f. 23. *Erebia mancinus* m. 24. *Coenonympha haydenii* m. 25-26. *Coenonympha tullia elko* syntypes (both called types on p. 184, thus not a Holland lectotype designation), 2m. 27. *Coenonympha tullia pseudobrenda* f. 28. *Erebia episodea=rhodia* m. 29. *Coenonympha tullia inornata* (called type in text, and missing so stated to be not a syntype by Brown; thus it might have been a syntype and thus might be considered Holland's lectotype, but the odds of it being not a syntype were high so it should not be considered to be a valid lectotype) m. 30. *Coenonympha tullia insulanus* Vanc. Is., m. 31. *Coenonympha tullia pseudobrenda* m.

Explanation of Plate XXVI. 1-2. *Cercyonis pegala maritima* (replaces *alope*, which is an intergrade of *pegala*X*carolina* says R. Gatrell), m, f. 3. *Cercyonis pegala nephele* m. 4. *Cercyonis pegala maritima* f. 5-6. *Cercyonis pegala boopis* (text says near San Francisco), m, f. 7. *Cercyonis oetus oetus* m. 8. *Cercyonis sthenele behrii* m. 9-10. *Cercyonis pegala nephele* (may be *olympus* syntypes but both are now lost, male is not *olympus* lectotype), m, f. 11-12. *Cercyonis oetus charon* syntypes (properly *paralectotypes*), m, f. 13-14. *Cercyonis meadii meadii*, f, m. 15-16. *Cercyonis pegala boopis=baroni* (not syntypes) 2m. 17. *Cercyonis pegala gabbii=utahensis* (not a *gabbii* syntype) f. 18. *Cercyonis pegala pegala* f. 19. *Cercyonis sthenele paulus* m (not a syntype). 20. *Cercyonis sthenele sthenele* m.

Explanation of Plate XXVII. 1-2. *Oeneis nevadensis gigas*, m, f. 3. *Oeneis* (nevadensis?) *macounii* (designated lectotype by Holland 1915 pl. LXXVI [bad printing and antenna fakery have made the 1915 & 1931 photos a little different], evidently a syntype and may be the syntype Brown found to have #5 on uns of label) m. 4. *Oeneis nevadensis iduna* (not a syntype; two figs. claimed to be "types" on p. 194 but only pl. LXII fig. 6m called type on plate, thus pl. LXII fig. 6 is Holland's designation of lectotype) m. 5. *Oeneis jutta* (ssp. *terraenovae* or *ridingiana* or *reducta*) f. 6. *Oeneis bore taygete* (Nain, Labrador) m. 7. *Oeneis polixenes brucei* (stated to be the "type" on p. 198, and Brown wrote that it is a syntype, therefore it is Holland's 1931 designation of lectotype; Brown says this fig. 7 male was coll. at the TL Bullion Mtn. July 3 and is a syntype, but did not consider it for Brown's lectotype designation because the O.D. said Aug.; however, Brown's type does not list Aug. either, and has just the locality "Colo." which does not match the TL, and none of the syntypes say Aug.; the bottom line is that this Aug. discrepancy does not matter, because fig. 7 is a syntype and Holland validly designated it as the lectotype; the valid TL is required to be locality of valid lectotype [this specimen] therefore is Bullion Mtn. [above Hall Valley, Park Co. Colo.]. Holland 1898 made no mention of type therefore did not designate a

lectotype.) lectotype m. 8. *Oeneis uhleri varuna* (probably not a syntype) m. 9. *Oeneis calais ivallda* (p. 194 says both book specimens are type, plate does not call this male type, so this male is paralectotype, and pl. LXII fig. 4 calls that female type so that female is lectotype designated by Holland 1931) m. 10. *Oeneis chryxus* m. 11. *Oeneis melissa semidea* m. 12. *Oeneis uhleri uhleri* atypical m.

Explanation of Plate XXVIII. 1-2. *Libytheana carinenta bachmanni* m. 3. *Libytheana carinenta larvata* f. 4-5. *Apodemia virgulti mejicanus* (not syntypes) f, Ariz. m. 6. *Apodemia virgulti davenporti=pratti* m. 7. *Apodemia mormo mormo* m. 8-9. *Apodemia nais nais*, m (coll. Morrison; not a syntype), lectotype f. 10. *Apodemia virgulti duryi* (designated lectotype on p. 213 by Holland 1931; however Holland 1898 p. 230 stated “The only specimen as yet known is the type figured in our plate [pl. XXVIII fig. 10f].”, therefore this is actually a Holland 1898 lectotype designation and not a 1931 designation) lectotype f. 11. *Apodemia palmeri palmeri* (not a syntype) m. 12-13. *Calephelis borealis* 2m. 14. *Calephelis rawsoni arizonensis* (W. McAlpine J. Res. Lep. 10:29)(not a syntype of *australis*) m. 15. *Calephelis nemesis nemesis* (not a syntype of *nemesis*, as F. Brown states it is *australis*) m. 16. *Calephelis virginienensis* m. 17-18. *Emesis ares* (not syntypes), m, f. 19-20. *Emesis zela cleis* (p. 216 says both specimens are types thus neither is Holland’s designation of lectotype), m lectotype (Brown claimed that this specimen was both fig. 19 & Brown’s lectotype, but Brown’s fig. of lectotype has fw tip a bit more falcate and the spots are a bit different, so even after sagging from moisture there is a little doubt that fig. 19 is Brown’s lectotype; if it isn’t it would be paralectotype), f paralectotype. 21. *Feniseca tarquinius* spring form m. 22. *Eumaeus atala* m. 23-24. *Lycaena arota arota*, m (could be ssp. *virginiensis*), f (female looks more like ssp. *arota* and not *virginiensis*). 25. *Lycaena phlaeas hypophlaeas* f. 26-27. *Lycaena xanthoides editha* (text p. 247 says both are types, thus neither is Holland’s designation of lectotype; Pelham Catalogue says these are syntypes currently in FMNH), m, f. 28. *Lycaena epixanthe phaedra=amicetus* (Boisduval & Scudder did not validly publish *amicetus* according to Miller/Brown note 348, but Holland validly published *amicetus* in this 1931 book, thus his 3 figs. on pl. XXVIII & LXIV are syntypes, and Clench 1968 later designated this male lectotype according to Pelham Catalogue) lectotype m. 29-30. *Lycaena xanthoides xanthoides*, m, f. 31-32. *Lycaena thoe*, m, f. 33-34. *Lycaena helloides*, m, f. 35-36. *Lycaena gorgon*. 37-38. *Lycaena mariposa mariposa*, m, f.

Explanation of Plate XXIX. 1-2. *Lycaena arota arota*, m, f. 3-4. *Lycaena rubidus sirius* (p. 252 says both are types thus neither is a lectotype designation by Holland), m lectotype, f (not a syntype). 5-6. *Lycaena rubidus rubidus*, m, f. 7-8. *Lycaena cupreus snowi* (not syntypes), m, f. 9. *Atlides haesus haesus* m. 10. *Parrhasius m-album* m. 11. *Hypaurotis crysalus crysalus* f. 12. *Habrodais grunus grunus* m. 13. *Satyrium favonius autolytus* f. 14. *Phaeostrymon alcestis alcestis* holotype f. 15-16. *Satyrium acadica*, m (now missing), f (the syn. *montanensis* according to Brown). 17. *Satyrium sylvinus putnami=itys* lectotype (designated *itys* lectotype by Holland 1931 on p. 238; however Holland 1898 p. 243 already designated this female the valid lectotype) lectotype f. 18. *Calycopis cecrops* f. 19-20. *Satyrium calanus calanus=wittfeldi*, f lectotype (designated lectotype by Holland 1915 pl. LXXI fig. 2), m paralectotype (p. 236 says both specimens are types, thus neither is a lectotype designation by Holland 1931). 21. *Callophrys spinetorum* f. 22. *Satyrium favonius favonius* m. 23-24. *Erora laeta laeta*, m, f (f now missing). 25. *Satyrium tetra=adenostomatis* paratype m. 26. *Satyrium calanus falacer* m. 27. *Satyrium edwardsii* f. 28. *Satyrium liparops liparops* f. 29. *Callophrys gryneus castalis=discoidalis* Tex. (now missing) m. 30. *Satyrium auretteorum auretteorum=tacita* m. 31. *Strymon melinus* f. 32. *Callophrys gryneus gryneus=smilacis* (not *castalis*) Md. m. 33-34. *Satyrium saepium saepium*, m, f. 35. *Ministrymon leda form ines* lectotype (p. 240 says this is type, and Brown says this is “paratype” syntype, so this is Holland’s 1931 designation of lectotype, which may now be in USNM or AMNH) f. 36-37. *Satyrium saepium saepium=chalcis*, m, f. 38. *Strymon acis bartrami* f. 39. *Chlorostrymon simaethis tacita* f.

Explanation of Plate XXX. 1. *Callophrys perplexa perplexa* m. 2. *Callophrys* probably *perplexa perplexa* (Cal.)(identified as *viridis* by Brown, but has orange unf and grass-green unh so is probably *perplexa* despite a few white dots), m. 3. *Callophrys perplexa perplexa* Cal. (not a syntype of *affinis*) m. 4-5. *Satyrium behrii behrii* (not syntypes) 2m. 6. *Ministrymon clytie* holotype f. 7. *Calycopis cecrops* m. 8. *Callophrys gryneus nelsoni* f. 9. *Callophrys gryneus siva* (not a syntype) m. 10. *Satyrium titus titus* m. 11. *Callophrys nippon nippon* f. 12. *Callophrys irus irus* m. 13. *Callophrys gryneus nelsoni* f. 14. *Satyrium titus titus* (neon band is bad printer alteration of photo) m. 15. *Callophrys augustinus* f. 16. *Satyrium fuliginosa fuliginosa* (not a syntype) m. 17. *Callophrys eryphon eryphon* f. 18. *Strymon martialis* m. 19. *Celastrina lucia Auctorum?*, form *marginata* (Winnipeg) m. 20. *Celastrina lucia* form *lucimargina* m. 21. *Callophrys henrici henrici* f. 22. *Callophrys nippon nippon* f. 23. *Glaucopsyche lygdamus* (not the silvery couperi, looks like ssp. *oro*) m. 24-25. Possibly *P. icarioides fulla*, but they look most like *Glaucopsyche lygdamus columbia*, m (Brown found it missing among *icarioides*, perhaps because it is in *lygdamus* drawer), f (“Mt. Hood Ore.”). 26. *Lycaena heteronea clara* (Holland 1898 p. 259 and Holland 1931 p. 253 say “The figure in the plate is that of the type of the female”, which sounds a little like he might be just describing the bluer female form; however Holland 1931 p. 253 also lists “Pl. XXX, fig. 26, female, type”, which is a definite Holland 1931 lectotype designation; it is a syntype according to Pelham Catalogue [syntypes in MGCL & AMNH]) lectotype f. 27. *Leptotes marina* f (m?). 28. *Plebejus saepiolus hilda* (see p. 261)(*daedalus* belongs to *icarioides*) f. 29. *Plebejus icarioides icarioides* m. 30. *Euphilotes enoptes enoptes* f. 31. *Euphilotes* (most likely *enoptes enoptes* or *e. tildeni* or *intermedia*) f. 32. *Celastrina neglectamajor* m. 33. *Echinargus isola* (not a syntype of *alce*) m. 34. *Glaucopsyche lygdamus couperi* m. 35. *Glaucopsyche lygdamus* (35-36 are not ssp. *xerces* f. *antiacis*)(looks like

ssp. columbia) m. 36. *Glaucopsyche lygdamus* (looks like ssp. couperi) m. 37. *Plebejus icarioides pheres* m. 38. *Echinargus isola* f. 39. *Euphilotes* perhaps *glaucan* (Not a syntype. Not *anna* as Brown claimed, the figure is obviously *Euphilotes* and could be *glaucan*. P. 266 calls it type, but the label on this *anna* specimen in CM reads “*Glaucan* male/Nev. Morr.”, the collector Herbert Morrison, so the original lost *Euphilotes* specimen could not have been a syntype because Brown wrote that the source of *glaucan* types was “2m, 1f, taken by Mr. Henry Edwards) m. 40. *Plebejus scudderi* *scudderi*=aster paralectotype m. 41. *Glaucopsyche lygdamus* (ssp. *xerces* form *antiacis*?, or could be ssp. *incognitus*, text calls figs. 35-36 & 41 “*behri*”) f. 42. *Plebejus icarioides pheres* f. 43. *Glaucopsyche lygdamus xerces* m. 44. *Glaucopsyche piasus piasus* f. 45. *Echinargus thomasi*, perhaps ssp. *bahamensis* (not *catalina*, a ssp. of *Leptotes cassius* from Virgin Is.) f. 46-47. *Plebejus scudderi scudderi*=aster paralectotypes, f, m. 48-49. *Plebejus samuelis* (according to Brown, & J. Masters *J. Lepid. Soc.* 26:153) m (Albany, N.Y.), f (N.Y.). 50. *Glaucopsyche lygdamus* probably ssp. *oro*, f. 51. *Euphilotes* *noptes* (*battoides* has similar ups) m.

Explanation of Plate XXXI (too whitish). 1-3. *Celastrina lucia* Auct. or *C. ladon* (#1 form *lucia* m, now missing; #2 form *marginata* m, missing; #3 form *lucimargina* f, ?Long Is.). 4. *Celastrina nigra* Coalburgh W. Va. m. 5. *Celastrina ladon* f. *violacea* W. Va. Apr. 13, m. 6-9. *Celastrina neglecta* (#8 missing, #9 reared)(Brown det. #6 & #9 as *neglecta*), m, f, m, f. 10. *Celastrina* (*neglecta*) *echo* S. Cal. W. Wright m. 11-12. *Plebejus saepiolus hilda* (*daedalus* belongs to *icarioides*), m, f. 13-14. *Lycaena heteronea* (syn. *klotsi*?), m, f. 15-16. *Plebejus saepiolus* (looks like N Cal. near-*rufescens*), m, f. 17-18. *Glaucopsyche lygdamus couperi* (silvery ups, however the whole plate is too whitish), m, f. 19-20. *Glaucopsyche piasus piasus*, m, f. 21-22. *Philotes sonorensis*, m, f. 23-24. *Plebejus shasta shasta*, m, f. 25. *Plebejus melissa melissa* Colo. T. L. Mead paralectotype (now broken) m. 26. *Plebejus anna anna* (not a syntype of *melissa*)(probably from “Summit” area W of Donner Pass Cal.) f. 27-28. *Plebejus acmon*, m, spring form f. 29-30. *Cupido comyntas comyntas*, m, f. 31. *Echinargus thomasi* (ssp. *bethunebakeri* or *bahamensis*) f. 32. *Leptotes marina* f.

Explanation of Plate XXXII. 1-2. *Philotiella speciosa speciosa*, m (Pelham Catalogue says this might possibly be holotype, but that is dubious as text only mentions fig. 2 is paratype), f (text says paratype). 3. *Hemiargus hanno gyas* m. 4. *Brephidium* (*exilis*) *pseudofea* m. 5. *Brephidium exilis exilis* m. 6. *Leptotes cassius theonus* f. 7-8. *Cupido amyntula amyntula*, m, f. 9-10. *Plebejus glandon franklinii* 2m. 11. *Euphilotes*, maybe *battoides* ssp. but looks most like *centralis* m. 12. *Cupido comyntas comyntas* m. 13. *Plebejus atrapraetextus annetta* (det. by F. M. Brown, now missing and ups resembles *P. melissa* so may really be *melissa*; not a syntype of *annetta*) m. 14. *Plebejus atrapraetextus annetta* (p. 264 says 3 specimens are types so neither is Holland’s designation of lectotype) paralectotype f. 15-16. *Plebejus podarce podarce* 2m. 17. *Plebejus glandon rustica* (now missing) m. 18. *Plebejus icarioides lycea* (not a syntype) S Colo. m. 19. *Lycaena heteronea heteronea*=*klotsi* f. 20. *Strymon melinus* ssp. m. 21-22. *Nathalis iole*, m, f. 23. *Euchloe hyantis hyantis* m. 24-25. *Euchloe ausonides ausonides*, m, f. 26-27. *Anthocharis cethura deserti*, m (aberrant fw apex), f. 28-29. *Anthocharis*, looks like *A. julia flora*, m, f. 30. *Anthocharis lanceolata lanceolata* m. 31-32. *Anthocharis sara sara* (not syntypes of *reakirtii*), m, f. 33. *Anthocharis cethura pima* lectotype (designated lectotype by Holland 1931) m. 34. *Anthocharis sara* (looks more like ssp. *sara*)(not syntype of *julia*) m. 35-36. *Anthocharis julia stella* (not syntypes of *stella*), m (aberrant, ssp. unknown), f. 37-38. *Anthocharis midea annickae*, m, f. 39. *Euchloe olympia=rosa* lectotype (designated *rosa* lectotype here by Holland 1931; Pelham Catalogue wrongly says Brown’s later invalid lectotype was this male but the two clearly differ in maculation) m.

Explanation of Plate XXXIII. 1. *Phoebis agarithe* m. 2-3. *Phoebis sennae sennae* wet season 2m. 4. *Phoebis philea* m. 5. *Colias eurytheme* m. 6. *Vanessa virginiensis* f.

Explanation of Plate XXXIV. 1. *Anthocharis cethura morrisoni* m. 2. *Euchloe* (*hyantis*) *lotta* f. 3. *Euchloe ausonides ausonides* m. 4-5. *Anthocharis julia flora*, m, f. 6. *Anthocharis julia julia* f. 7. *Neophasia menapia menapia* m. 8-9. *Pontia beckerii* (not syntypes), m, f. 10-11. *Pontia protodice*, m, f. 12. *Pontia sisymbrii sisymbrii* m. 13. *Pontia callidice occidentalis* m. 14. *Pieris virginiensis* neotype m. 15. *Pieris*, perhaps *marginalis marginalis* form *pallida* (a hybrid with *P. rapae*?, odd-looking with black upf bar) f. 16. *Pieris oleracea* m. 17. *Pieris hulda pseudobryoniae* f. 18. *Pontia protodice* spring form *vernalis* (not a syntype) m. 19. *Pieris oleracea=frigida=acadica* (paralectotype of *acadica*) f. 20-21. *Kricogonia lyside*, m, f.

Explanation of Plate XXXV. 1-2. *Ascia monuste phileta*, m, f. 3. *Pieris rapae* f. 4-5. *Appias drusilla neumogeni*, m, f. 6-7. *Colias alexandra alexandra* (not syntypes), m, f. 8-9. *Colias scudderii scudderii*, m, f. 10-11. *Colias interior*, m, f. 12-13. *Colias occidentalis chrysomelas*, m, f. 14. *Colias pelidne pelidne* (borders unusually narrow) m. 15. *Colias philodice philodice* m.

Explanation of Plate XXXVI. 1-2. *Zerene eurydice*, m, f. 3-4. *Zerene cesonia cesonia*, m, f. 5-6. *Colias meadii meadii* (p. 297 says both figs are types, thus neither is Holland’s designation of lectotype) lectotype m, paralectotype f. 7-8. *Colias eurytheme* form *ariadne*, m, f. 9-10. *Colias eurytheme* summer form=“*keewaydin*”, m (lectotype of *keewaydin*), f. 11-12. *Colias nastes* (looks like ssp. *gueneei*), m, f. 13-14. *Colias meadii elis* (p. 298 says both are types so neither is Holland’s designation of lectotype; Pelham Catalogue says one of the two is possibly holotype & states that current holotype is in Field Museum, but actually there is no holotype as O.D. failed to designate one), m, f. 15. *Colias pelidne pelidne* m. 16. *Colias pelidne pelidne* (but looks more like *scudderii gigantea*) f. 17. *Colias behrii* lectotype (designated lectotype by Holland 1931 on p. 299) m.

Explanation of Plate XXXVII. 1. *Eurema proterpia* f. *gundlachia*=*longicauda* m. 2. *Eurema proterpia* m. 3-6. *Eurema nicippe*, m, pale aberration f, yellow aberration *flava* m, f. 7-8. *Eurema mexicana* 2m. 9-10. *Eurema salome* (not *damaris*, which is a syn. of *mexicana*), m, f. 11. *Eurema dina westwoodi* m. 12. *Eurema दौरα lydia* f. *lydia* (summer form) m. 13. *Eurema lisa* m. 14. *Eurema दौरα दौरα*=*delia* winter f. *daura* m. 15-16. *Eurema दौरα दौरα* summer f. *jucunda* 2m. 17-18. *Enantia melite*, m, f.

Explanation of Plate XXXVIII. 1. *Papilio zelicaon* m. 2. *Papilio multicaudata multicaudata* m. 3. *Papilio pilumnus* m.

Explanation of Plate XXXIX. 1-2. *Parnassius phoebus behrii*, m, f. 3-4. *Parnassius phoebus smintheus* variants, m, f. 5-6. *Parnassius phoebus maximus*, m, f. 7-8. *Parnassius clodius "baldur"*, m, f. 9-10. *Parnassius clodius clodius*, m, f.

Explanation of Plate XL. 1. *Papilio polyxenes asterias* m. 2. *Papilio machaon bairdii* (not a syntype) m. 3. *Papilio machaon bairdii* form *hollandii* (designated lectotype by Holland 1931) lectotype m. 4. *Papilio machaon brucei* (designated lectotype by Holland 1931) lectotype f. 5. *Papilio brevicauda brevicauda* f.

Explanation of Plate XLI. 1. *Papilio machaon hudsonianus* (Rupert's House, Hudson Bay)(p. 314 says "type" of *aliaska*, but not a syntype; Pelham Catalogue claims this was Holland's 1931 designation of *aliaska* lectotype, which if true would make *hudsonianus* a syn. of *aliaska*. However, p. 314 also says Holland has another type from Scudder's original material labeled *Alaska*; thus Holland called two specimens types, therefore neither is a lectotype designation by Holland, and *hudsonianus* is a valid ssp. And p. 314 says this specimen is from Rupert's House, and Scudder's *aliaska* types were all from *Alaska*, thus this male is not a syntype of *aliaska*) m. 2. *Papilio zelicaon* form *nitra* lectotype (designated lectotype by Holland 1931) m. 3. *Papilio indra indra* f. 4. *Battus polydamas polydamas* m. 5. *Papilio troilus troilus* m. 6. *Papilio polyxenes asterias* aberration *calverleyi* (evidently holotype; text says type; Miller/Brown say HT may be in AMNH or Albany; Pelham Catalogue says this is holotype probably in CM) f.

Explanation of Plate XLII. 1. *Papilio palamedes palamedes* f. 2. *Battus philenor philenor* m. 3. *Papilio crespontes* m. 4. *Papilio thoas cinyras* (*Amazons* in Rothschild & Jordan, *Brazil* in text) m.

Explanation of Plate XLIII. 1. *Papilio glaucus glaucus* (yellow form "turnus") m. 2. *Papilio glaucus glaucus*, f black form. 3. *Colias philodice=hageni* f. 4. *Vanessa atalanta rubria* f. 5. *Epargyreus clarus* m.

Explanation of Plate XLIV. 1. *Eurytides marcellus*, spring form m. 2. *Eurytides marcellus*, spring form m (Holland 1898 named *floridensis* & designated holotype by monotypy on p. 307 "Another winter form, for which I propose the name *floridensis*, is represented in Plate XLIV, Fig. 2, by a male specimen.") spring form m. 3-4. *Eurytides marcellus*, summer form m, spring form (holotype *walshii*) m. 5. *Papilio eurymedon* m.

Explanation of Plate XLV. 1. *Papilio glaucus rutulus* m. 2. *Hesperopsis alpheus* (not a syntype) Ariz. m. 3. *Calpodus ethlius* f. 4. *Pholisora catullus* m. 5. *Erynnis afranius* m. 6. *Urbanus proteus* m. 7. *Erynnis brizo* f. 8. *Erynnis juvenalis clitus* (evidently not a syntype) m. 9. *Apyrrothrix araxes arizonae* m. 10. *Achalarus lyciades* f. 11. *Zestusa dorus* lectotype (designated lectotype here by Holland 1931) m. 12. *Autochton cellus* m.

Explanation of Plate XLVI. 1. *Systasea pulverulenta* (not a syntype of *zampa* as stated in text; the real type was lost at sea) m. 2-3. *Hesperia comma manitoba*, m, f. 4-5. *Atalopedes campestris*, m, f. 6. *Atrytone arogos* m. 7-8. *Poanes melane melane* (not syntypes as text claims)(from San Bernardino, W. Wright), m, f. 9-10. *Atrytonopsis hianna hianna*, m, f. 11-12. *Hesperia ottoe* (not syntypes as text claims) m (Neb.) f. 13. *Hesperia sassacus sassacus* m. 14. *Poanes aaroni=howardi* (not a type on p. 391) m. 15. *Poanes viator zizaniae* (not a syntype of *viator* as text claims) Tex. f. 16-17. *Euphyes conspicua*, m (now missing), f (Long Island). 18-19. *Hylephila phyleus phyleus*, m, f. 20. *Problema byssus byssus* (text p. 386 says both figs. on 2 pl. are types, but this is not called type on plate whereas male on pl. LIII fig. 38 is called type on plate, thus LIII fig. 38 is Holland 1931 lectotype designation) paralectotype f. 21. *Euphyes pilatka* m. 22-23. *Polites mystic mystic*, neotype m, f. 24-25. *Anatrytone logan logan* (Kanawha R., W. Va.), neotype of *delaware* m, neotype of *logan* f. 26-27. *Stinga morrisoni* (So. Colo., 7500')(p. 372 says 3 specimens are types, but on plates only pl. LII fig. 15 male is called type, thus the latter is Holland 1931 designation of lectotype, not this specimen), paralectotype m, paralectotype f. 28-29. *Wallengrenia otho otho*, m, f. 30. *Polites themistocles=taumas* f. 31. *Euphyes vestris metacomet* (not *ruricola*) m (now missing). 32. *Pompeius verna* m. 33. *Lerodea eufala* f. 34. *Panoquina ocola* m (now missing). 35. *Oligoria maculata* m. 36. *Amblycirtes carolina* (Holland 1931 p. 362 designates this male lectotype; Miller/Brown say HT in CM suggesting this may be holotype, but Pelham Catalogue says this is syntype [syntypes in ANSP]; actually Holland 1898 p. 367 stated that this specimen is the type, so it is the valid lectotype designation is by Holland 1898) lectotype m. 37. *Poanes aaroni* paratype (now lost) m. 38. *Poanes aaroni* syn. *howardi* (designated lectotype by Holland 1931 on p. 391 [the fig. on pl. XLVI fig. 14 is not type and fig. on LIV fig. 1 is paratype according to p. 391 & plates]; Miller/Brown say HT in CM, but Gatrell Tax. Report of Int. Lep. Survey 2[2]:6-7 wrote that no holotype was designated and fig. 38 was a syntype; the bug in fig. 38 is now lost [in Gatrell's paper, pl. XLVI fig. 38m *Poanes howardi* in 1931 book is the same bug as *Phycanassa howardi* in 1898 book {which Gatrell wrote was on pl. 46 then wrongly wrote LXVI, actually XLVI}); Gatrell designated a 2nd lectotype in CMNH with "type 7092" label, which is invalid; Holland's earlier 1931 lectotype is valid even though lost. Holland 1898 did not designate a lectotype.) lectotype m. 39. *Thorybes mexicana=aemilia* (Fort Klamath Ore.)(designated *aemilia* lectotype by Holland 1898 on p. 325; Miller/Brown say HT should be in CM, but there must not be a holotype as Pelham Catalogue says Holland 1931 designated this lectotype, which is true but Holland 1898

already designated it) lectotype f. 40. *Poanes yehl* (text says type; this would be a Holland 1898 lectotype designation, but there is a holotype, in FMNH) m.

Explanation of Plate XLVII. 1. *Carterocephalus* "palaemon" "magnus" Mendocino Cal. m. 2. *Ancyloxypha numitor* m. 3. *Oarisma edwardsii* m. 4. *Oarisma poweshiek* m. 5. *Amblyscirtes vialis* (Hunter N.Y.) m. 6. *Amblyscirtes hegon*=samoset (now missing) m. 7. *Amblyscirtes aenus aenus* f (now missing). 8. *Notamblyscirtes simius* (designated lectotype on p. 369 in Holland 1931 Butt. Book—Pelham Catalogue says Holland 1898 designated lectotype but he surely meant 1931 as *simius* is on p. 369 in 1931 book and on p. 341 in 1898 book which was thoroughly revised in 1931. Also, Holland 1898 did not say any specimen was type thus made no lectotype designation) lectotype m. 9. *Copaeodes aurantiaca*=waco (Ariz., Morrison coll.) m. 10. *Pseudocopaeodes eunus eunus*="wrighti" (p. 369 says this is type, and Brown wrote it may have been a syntype but Brown found no Edwards label on it thus there is no proof it was syntype, so this is not a valid Holland lectotype designation; Brown designated the valid lectotype) paralectotype m. 11. *Ancyloxypha arene* (not a *myrtis* syntype) m. 12. *Pyrgus scriptura* f. 13. *Pyrgus centaureae* (ssp. *wyandot* evidently, maybe *loki*) m. 14. *Pyrgus ruralis ruralis* f. 15. *Pyrgus xanthus* So. Col. paralectotype f. 16. *Amblyscirtes aesculapius*=wakulla lectotype m. 17. *Celotes nessus* Bastrop Co. Tex. m. 18. *Pyrgus communis* (not *P. philetas*=occidentalis which has darker ups) m. 19. *Heliopetes domicella* m. 20. *Polites themistocles*=taumas m. 21-22. *Poanes massasoit massasoit*, m, f. 23. *Hesperia attalus attalus* m (now missing). 24-25. *Polites peckius peckius*, m, f. 26. *Polites mardon mardon* (p. 381 says both figs. pl. XLVII fig. 26 male and pl. LIII fig. 12 female are types, but plates say only pl. LIII fig. 12 female is type, so the latter female is a Holland 1931 lectotype designation, even though pl. XLVII fig. 26m is a syntype; Pelham Catalogue states that Holland's 1898 Butt. Book designated the pl. 47 fig. 26 male as lectotype, but Holland 1898 p. 354 & plate explanation does not say that pl. XLVII fig. 26m is a type, so this is NOT a Holland 1898 lectotype designation) paralectotype m. 27-28. *Hesperia uncas uncas*, m neotype N Col., f. 29-30. *Paratrytone snowi* Ariz., m, f. 31-32. *Poanes taxiles* (p. 390 says both specimens are types, so neither is a Holland 1931 lectotype designation), lectotype m, f (evidently paralectotype, now missing). 33-34. *Hesperia metea metea*, m, f. 35-36. *Hesperia leonardus leonardus*, m, f. 37-38. *Poanes hobomok hobomok*, m, f. 39. *Poanes zabulon* f. 40. *Hylephila phyleus phyleus* m. 41. *Polites vibex vibex*=brettus f. 42-43. *Polites sabuleti sabuleti*, m, f. 44. *Hesperia comma* (looks like ssp. *ochracea*) m.

Explanation of Plate XLVIII. 1. *Erynnis persius* m. 2. *Erynnis brizo brizo*=somnus m. 3. *Erynnis zarucco terentius* f. 4. *Erynnis martialis* m. 5. *Thorybes bathyllus* f. 6. *Thorybes pylades pylades* f. 7. *Erynnis horatius* f. 8. *Lerema accius* m. 9. *Erynnis pacuvius pacuvius* f. 10. *Erynnis lucilius* (?) m. 11. *Erynnis juvenalis juvenalis* f. 12. *Erynnis funeralis* m. 13. *Achalarus casica*=epigena m. 14. *Hesperopsis libya libya* m. 15. *Erynnis horatius* f. 16. *Staphylus hayhurstii* (Indian River Fla.) f. 17. *Erynnis icelus* m. 18. *Colias eurytheme albino* f.

Explanation of Plate XLIX. 1. *Phocides pigmalion okeechobee* m. 2. *Phocides polybius lilea* m. 3. *Phocides urania* m. 4. *Dyscophellus euribates* m. 5. *Polygonus leo*=amyntas m. 6-7. *Proteides mercurius mercurius*=idas, m, f. 8-9. *Epargyreus exadeus*, m, f. 10. *Epargyreus zestos* m. 11-12. *Urbanus dorantes dorantes*, m, syn. *rauterbergi* paratype m. 13. *Urbanus simplicius* m. 14. *Urbanus teleus*=eurycles m. 15. *Codatractus arizonensis* paratype m. 16. *Codatractus carlos* f. 17. *Chioides catillus albofasciatus* f. 18. *Chioides zilpa* m. 19. *Aethilla memmius*=hahneli paratype m.

Explanation of Plate L. 1. *Thorybes confusus* paratype m. 2. *Thorybes confusus* (spots wider than normal but spot in cell M₃ far from spot in cell CuA₁) paratype f. 3. *Thorybes mexicana mexicana*=nevada (? , atypical) paratype (properly syntype, as there is no holotype) m. 4. *Thorybes drusius* (Holland 1931 designation of lectotype) lectotype m. 5-6. *Thorybes mexicana mexicana* 2m. 7-8. *Cogia hippalus*, f (not a syntype), m (may be a syntype). 9. *Autochton pseudocellus* m. 10. *Cabares potrillo* m. 11. *Cogia outis* paratype m. 12. *Cogia calchas* m. 13-14. *Cogia caicus moschus*, lectotype (designated lectotype by Holland 1931) m, paralectotype m. 15. *Pyrgus oileus* (doubtfully *orcas*, which is the Costa Rica-S. Amer. ssp.) m. 16-17. *Pyrgus oileus oileus*=syrictus, f, m. 18. *Pyrgus oileus oileus*=montivaga m. 19-22. *Pyrgus communis*, m, f, m, f. 23. *Pyrgus communis* (not *occidentalis*, which is a syn. of *P. philetas*) m. 24. *Pyrgus philetas* holotype f. 25-26. *Heliopetes laviana*, m, f. 27-28. *Heliopetes ericetorum*, m, f. 29. *Heliopetes macaira macaira* pale form (not *nivella*)(according to Evans' Amer. Hesperidae), m. 30-31. *Heliopetes macaira macaira* 2m. 32. *Pholisora mejicanus* m. 33-34. *Staphylus mazans* (received from Staudinger, so evidently ssp. *ascalaphus* paratypes from S Mex.-S Amer.), m, f. 35. *Staphylus ceos* (not a syntype) Ariz. m. 36. *Hesperopsis libya lena* lectotype (designated lectotype by Holland 1931) m (Montana). 37. *Chiomara asychis* m. 38. *Xenophanes trixus* m. 39. *Achlyodes thraso thraso* (probably Chiriqui, Panama) m. 40-41. *Ephyriades brunnea*, m, f.

Explanation of Plate LI. 1-2. *Gesta gesta invisus*, m, f. 3-4. *Erynnis brizo brizo*=somnus (multiple specimens are called type on p. 350 & on plate, thus none is a Holland lectotype designation) lectotype (designated by J. Calhoun) m, paralectotype f. 5. *Ephyriades brunnea* f. 6. *Ephyriades zephodes* f. 7. *Erynnis brizo burgessi* paratype m. 8. *Erynnis brizo lacustra* m. 9. *Erynnis "scudderi paratype"* (looks like *pacuvius* to me; some of these *Erynnis* would require genitalic dissection for positive identification) m. 10. *Erynnis tristis tristis* m. 11. *Erynnis horatius* m. 12. *Erynnis propertius propertius* m. 13. *Grais stigmaticus* m. 14. *Timochares ruptifasciatus* m. 15. *Piruna pirus* (Water Can., Socorro Co. NM, not a syntype) m. 16-17. *Oarisma garita*, 2m. 18. *Carterocephalus palaemon skada* (Rampart, Alaska, not a syntype) m. 19-20. *Piruna polingii* (plate says 19 is paratype, text says 20 is paratype), 2m. 21. *Copaeodes aurantiaca* m. 22-23. *Copaeodes minima* (not syntypes), m Comfort Tex., f Tex. 24-25. *Amblyscirtes nysa* (both figs. are called types on p. 364

& on plate, thus neither is a Holland designation of lectotype), m (not a syntype), lectotype f. 26-27. *Amblyscirtes eos=comus* (W Tex.), f (text says type of *comus*, but not a syntype), m (neotype of *comus*). 28-29. *Erynnis persius=avinoffi* syntypes (male is holotype, female allotype, according to O.D.), m, f. 30. *Amblyscirtes celia* paratype m. 31. *Amblyscirtes cassus* lectotype (designated lectotype by Holland 1931) m. 32. *Amblyscirtes phylace* S Colo. holotype f. 33-34. *Amblyscirtes alternata*, f, m. 35-36. *Amblyscirtes exotera=nanno* (Ariz., H. Morrison, not syntypes), m, f. 37. *Thymelicus lineola* m. 38. *Amblyscirtes fimbriata=bellus* (Huachuca Mts., Ariz., not a syntype) m. 39. *Amblyscirtes oslari* paratype m. 40. *Copaeodes aurantiaca=waco=candida* (Ariz., H. Morrison) f. 41. *Choranthus haitensis* paratype m. 42. *Amblyscirtes nereus* (Ariz., H. Morrison, not a syntype) m. 43-44. *Polites* (*Yvretta*) *carus* (both figs are called types on p. 369 & on plate, thus neither is a Holland lectotype designation), m (Ariz., not a syntype), lectotype f. 45-46. *Polites* (*Yvretta*) *rhesus* (both figs are called types on p. 369 & on plate, thus neither is a Holland lectotype designation), lectotype m, paralectotype f. 47. *Choranthus radians* m. 48-49. *Hesperia comma laurentina*, m, f.

Explanation of Plate LII. 1. *Hesperia nevada* m (so doubtfully a syntype of *colorado* unless it was misidentified in type series—and I greatly doubt that this is the specimen declared *colorado* lectotype by Barnes & McDunnough [1916] as some people might interpret Pelham Catalogue—because this is the wrong species and fig. 2 is not called type on plate; p. 371 says both figs. are types but plate says only fig. 1 is type, so fig. 1 would be a lectotype designation, but I do not consider it to be a valid Holland 1931 designation of *colorado* lectotype because it is very doubtfully a syntype of *colorado*). 2. *Hesperia comma colorado* (p. 371 says figs. 1-2 are both types, but this is not called type on plate, so doubtfully a syntype) f. 3. *Hesperia comma idaho* (designated lectotype by Holland 1931; Brown & Miller wrongly claimed that Barnes & McDunnough 1916 [Contrib. Nat. Hist. Lepid. 3 #2:127] designated it earlier as lectotype. Actually Barnes & McDunnough wrote “*Idaho* Edw., described from Oregon, Washington, and California is a form with paler, yellowish green underside of secondaries, on which the white markings stand out very distinctly; we would restrict the type to the male from *East Calif.* in the Edwards’ Collection.” They would have—but did not. They would have designated a lectotype if they had said that “we restrict the lectotype”; instead Holland 1931 validly designated this lectotype.) lectotype m. 4. *Hesperia comma assiniboia* paratype m. 5-6. *Hesperia comma hulbirti* X *harpalus* “*oregonia*” (worthless intergrade name)(p. 372 & plate say that both specimens are types, therefore neither is a Holland lectotype designation) (Cal., O. Baron), m lectotype, f paralectotype. 7-8. *Hesperia juba*, m, f. 9. *Hesperia juba=ogdenensis* holotype f. 10. *Hesperia viridis* m. 11. *Hesperia woodgatei* paratype m. 12-14. *Hesperia nevada* (p. 372 & plates say that all 3 specimens are types, thus none is Holland lectotype designation) paralectotypes (lectotype is in MCZ, fig. 12 sent from Scudder, 13-14 in Mead coll.), m (not holotype), f, f. 15. *Stinga morrisoni* (evidently syntype given by Edwards to his brother-in-law Theodore Mead [as in *Problema byssus*]; p. 372 calls this plus the two figs. on pl. XLVI types, but on plates only this fig. 15 is called type, thus this pl. LII fig. 15 male is lectotype designation by Holland 1931) lectotype m. 16. *Hesperia comma idaho* (or *colorado*) m. 17-18. *Hesperia comma harpalus=cabelus* (not “*Nev.*”)(H. Morrison)(p. 373 says both specimens are types, but only fig. 17 is called type on plate, which is thus Holland 1931 designation of lectotype), lectotype m, paralectotype m. 19. *Hesperia comma harpalus* (not “*Nev.*”)(H. Morrison)(p. 373 & plate says that both specimens are types, thus neither is Holland 1931 designation of lectotype) lectotype m. 20. *Polites sabuleti* (doubtfully a *harpalus* syntype but perhaps a *harpalus* paralectotype f that was misidentified) f. 21. *Hesperia sassacus=manitoboides* f. 22. *Hesperia dacotae* paratype m. 23. *Hesperia sassacus* f. 24. *Hesperia attalus attalus* (Tex.)(designated lectotype by Holland 1931) lectotype f. 25-27. *Hesperia attalus slossonae=seminole*, m, f, m. 28. *Hesperia metea licinus* (Tex.) holotype m. 29. *Hesperia leonardus montana* paratype m. 30-31. *Hesperia leonardus pawnee*, m paratype (or just Neb. topotype; Pelham Catalogue says syntype as there is evidently no holotype/lectotype, Miller/Brown say type location unknown), f (Mont., not a syntype according to text). 32-34. *Hesperia meskei meskei*, paralectotype of *straton* (Indian River Fla.; Brown wrote that this is *straton*) m, holotype of *meskei* (Bastrop Tex.) f, lectotype of *straton* (designated lectotype of *straton* by Holland 1931) (Indian River Fla.) m. 35-36. *Ochlodes sylvanoides sylvanoides*, m, f. 37-38. *Ochlodes sylvanoides sylvanoides=napa* (not syntypes), m Colo. D. Bruce, f Idaho Springs Colo. T. Mead. 39-40. *Ochlodes agricola agricola=nemorum*, m, f. 41. *Ochlodes sylvanoides sylvanoides=pratincola* m. 42-43. *Ochlodes agricola verus* (p. 378 & plate both say both specimens are types, thus neither is a Holland lectotype designation) Havilah, Kern Co. Calif., lectotype m, paralectotype f. 44. *Ochlodes agricola agricola=milo* holotype m. 45-46. *Ochlodes agricola agricola*, m, f.

Explanation of Plate LIII. 1. *Ochlodes yuma yuma* m (Jensen P.O., Uintah Co. Utah). 2. *Ochlodes yuma yuma=scudderi*, paratype m. 3. *Polites origenes origenes* Coalburgh W.Va. *melanic* m. 4. *Polites origenes rhenae* (not a syntype) m. 5. *Polites mystic* m (*alcina* is a syn. of *rhenae*)(perhaps a paratype of *alcina* and misidentified). 6. *Polites themistocles=taumas* f. 7. *Polites origenes origenes* f. 8. *Euphyes vestris* f (now missing). 9. *Polites sonora siris* (Mt. Hood, Ore.) (designated lectotype by Holland 1931) lectotype m. 10-11. *Polites sonora sonora*, m (p. 381 says paratype), f. 12. *Polites mardon mardon* (p. 381 says both figs. on pl. XLVII fig. 26m & pl. LIII fig. 12f are types, but in plates only LIII fig. 12f is called type, thus the latter female is a Holland 1931 lectotype designation; Pelham Catalogue incorrectly stated that Holland 1898 designated pl. XLVII fig. 26 as type, which Holland did not do) lectotype f. 13. *Polites sonora utahensis* paratype m. 14. *Polites peckius peckius* f. 15-16. *Polites draco* (Colo.)(p. 382 & plate both say that both specimens are types, thus neither is Holland lectotype designation) lectotype m, paralectotype f. 17. *Polites sabuleti tecumseh* m. 18-19. *Polites sabuleti chusca* holotype m, f. 20. *Polites mystic mystic* f. 21-22. *Polites baracoa*, m, f. 23. *Polites mystic dacotah*

(Colo.) holotype m. 24-25. *Polites vibex brettus* 2m. 26. *Polites vibex praeceps=brettoides* (Tex.) (designated *brettoides* lectotype by Holland 1931) lectotype m. 27. *Polites vibex vibex* m. 28. *Polites vibex praeceps=stigma* paratype m. 29. *Wallengrenia otho clavus=curassavica* f. 30. *Anatrytone logan lagus* (E Mont., not a syntype) m. 31-32. *Polites origenes origenes* Coalburgh W.Va., m, f. 33-34. *Euphyes arpa*, m, f. 35. *Euphyes pilatka* (Indian River Fla.) f. 36-37. *Euphyes dion* (Whiting, Ind.), (designated lectotype by Holland 1931) lectotype m, paralectotype f. 38. *Problema byssus byssus* (text p. 386 says both specimens are type, but plates says only pl. LIII fig. 38 is type, thus this fig. 38 is Holland 1931 designation of lectotype) lectotype m. 39. *Problema bulenta* (text p. 386 says this is *kumskaka* paratype, but plate says only det. Scudder [in error], and specimen is not *P. byssus kumskaka*; it cannot be a *kumskaka* paratype) m. 40-41. *Euphyes vestris vestris=osceola* paratypes, m, f. 42-45. *Poanes hobomok hobomok*, m, female form *pocahontas* f, *pocahontas=quadaquina* paratype f, *pocahontas=alfaratta* holotype f. 46-47. *Poanes zabulon* 2m.

Explanation of Plate LIV. 1. *Poanes aaroni=howardi* paratype f. 2. *Poanes viator viator* (Ont., not a syntype) m. 3. *Atrytonopsis vierecki* paratype m. 4-5. *Atrytonopsis python python*, holotype m, paratype f (Brown mistakenly illustrated this female and called it the holotype). 6-7. *Atrytonopsis hianna loammi*, m Lutz Fla., f (now missing). 8. *Hesperia attalus slossonae* (Fla., H. Morrison) f. 9-10. *Thespieus macareus* 2m. 11. *Atrytonopsis pittacus* (designated lectotype by Holland 1931) lectotype m. 12. *Atrytonopsis python margarita* (evidently paratype) m. 13-14. *Atrytonopsis hianna deva* (Ariz., H. Morrison, not syntypes), m, f. 15-16. *Atrytonopsis lunus* (p. 393 & plates both say both specimens are types, thus neither is Holland lectotype designation), lectotype m (Ariz., H. Morrison), f (probable paralectotype). 17. *Nastra neamathla neamathla* paratype m. 18-19. *Nastra lherminieri*, m, f (odd hw shape but evidently not *Oarisma*). 20-21. *Panoquina panoquin*, m (Fla.), f. 22. *Lerema accius* (not a syntype of *ocola*; Brown said the figured bug was missing but he was looking in the *ocola* drawer and it is surely in the *accius* drawer) f. 23. *Panoquina panoquinoides panoquinoides* paratype m. 24. *Panoquina panoquinoides errans* paratype m. 25. *Panoquina lucas=sylvicola* m. 26. *Euphyes dukesi dukesi* (text says type, plate says paratype; called holotype by Pelham Catalogue) paratype or holotype m. 27-28. *Megathymus yuccae coloradensis* 2m. 29. *Agathymus aryxna aryxna* m. 30. *Megathymus streckeri streckeri* m. 31-32. *Agathymus polingi polingi* paratypes, m, f. 33. *Agathymus alliae alliae* m. 34-35. *Megathymus cofaqui cofaqui*, f, m.

Explanation of Plate LV. 1. *Lycorea halia atergatis* m. 2-4. *Argynnis* (Speyeria) *nokomis coeruleascens* (p. 85 & plate says two specimens are types, thus neither is a Holland lectotype designation), f paralectotype, m lectotype, f paralectotype. 5-6. *Argynnis* (Speyeria) *cybele carpenteri* (p. 87 & plate says both are types, thus neither is Holland lectotype designation), lectotype m, allotype (properly paralectotype) f. 7. *Argynnis* (Speyeria) *hydaspae purpurascens* paralectotype m. 8. *Boloria selene nebraskensis* holotype m. 9-10. *Boloria frigga gibsoni=lehmanni*, holotype m, allotype f. 11. *Boloria eunomia tricularis* m (the black wing bases esp. on right are evidently smut). 12. *Boloria polaris* m. 13. *Boloria selene terraenovae* holotype m (too melanic, more smut?). 14-15. *Boloria bellona toddi*, holotype m, allotype f. 16-18. *Boloria titania* (*chariclea*) *butleri*, m & f (not syntypes), lectotype (designated lectotype by Holland 1931) m. 19-21. *Boloria selene atrocotalis=albequina* (p. 107 & plate says two specimens are types, thus none is a Holland lectotype designation) syntypes, m, f, m. 22-23. *Boloria titania* (*chariclea*) *arctica*, m, f. 24-25. *Boloria alaskensis alaskensis*, m holotype, f. 26. *Boloria bellona bellona=pardopsis* holotype f. 27-28. *Boloria improba improba*, m, aberration *youngi* holotype m.

Explanation of Plate LVI. 1. *Argynnis* (Speyeria) *cybele charlottii* f (evidently syntype; Pelham Catalogue calls this holotype, which is in USNM). 2. *Argynnis* (Speyeria) *cybele krautwurmi* holotype f. 3. *Argynnis* (Speyeria) *egleis utahensis* paratype m. 4. *Argynnis* (Speyeria) *hydaspae rhodope=sakuntala* paralectotype f. 5. *Argynnis* (Speyeria) *coronis coronis=californica* paratype m. 6. *Argynnis* (Speyeria) *mormonia luski* paralectotype f. 7. *Argynnis* (Speyeria) *mormonia erinna* (designated lectotype by Holland 1931) lectotype m. 8. *Argynnis* (Speyeria) *mormonia washingtonia* (text says paralectotype) m. 9. *Boloria selene atrocotalis=jenningsae* holotype m. 10. *Euphydryas anicia maria* paratype (properly syntype as there is no holotype) m. 11-12. *Boloria eunomia dawsoni* paratypes, m, f. 13. *Chlosyne whitneyi damoetas* paratype m. 14-15. *Chlosyne definita definita* paratype (properly paralectotype, as there is no holotype) 2m. 16. *Chlosyne sterope* (*acastus*) *neumoegeni* paratype m. 17. *Boloria kriemhild=laurenti* paratype (properly syntype as there is no holotype or lectotype) m. 18-19. *Euphydryas chalcedona chalcedona=cooperi* (not syntypes), m, f. 20-21. *Euphydryas anicia carmentis* paratypes, m, f. 22. *Euphydryas gillettii* paratype (properly paralectotype, as there is no holotype) m. 23-24. *Euphydryas anicia magdalena* paratypes, m, f. 25. *Chlosyne palla flavula* paratype (properly syntype, as there is no holotype or lectotype) m.

Explanation of Plate LVII. 1. *Euphydryas chalcedona dwinellei* m. 2-3. *Euphydryas chalcedona paradoxa=perdiccas* (p. 117 & plate say both specimens are types, thus none is Holland lectotype designation), lectotype m, allotype (properly paralectotype) f. 4-5. *Euphydryas chalcedona olancha*, m, f. 6-7. *Euphydryas chalcedona sierra*, m, f. 8-9. *Euphydryas chalcedona hennei="quino"*, m, f. 10-11. *Euphydryas anicia anicia*, m, f. 12-13. *Euphydryas anicia helvia*, m, f. 14. *Euphydryas bernadetta bernadetta* paratype (properly paralectotype, as there is no holotype) m. 15-16. *Euphydryas anicia eurytion* syntypes, m, f. 17. *Euphydryas chalcedona macglashanii=truckensis* f. 18-19. *Euphydryas anicia capella* paratype (properly paralectotype, as there is no holotype), m, f. 20-21. *Euphydryas anicia alena* paratypes, m, f. 22-23. *Euphydryas anicia morandi* holotype m, allotype f. 24-25. *Euphydryas chalcedona irelandi* holotype m, allotype f. 26-27. *Euphydryas editha edithana=monoensis* paratypes, m (p. 123 & Pelham Catalogue calls this *monoensis* holotype but plate says it is paratype, and the fig. look a little different from Gunder's fig. of holotype), f. 28-29. *Euphydryas editha aurilacus*

(or a syn. of *rubicunda*) paratypes (text says holotype & allotype)(Pelham Catalogue calls fig. 28 holotype but the fig. looks a little different from Gunder's fig. of "type", so probably Gunder's is holotype and Holland's is paratype), m, f.

Explanation of Plate LVIII. 1. *Euphydryas anicia wheeleri* paratype (G. Austin, Syst. W N.A. Butt. chap. 50:635) m. 2. *Euphydryas editha colonia* m. 3. *Euphydryas anicia* (ssp. *wheeleri*?) m. 4-6. *Euphydryas editha editha*, f (not a *rubicunda* syntype as claimed in text), m, f. 7. *Euphydryas editha quino=augusta* (not a syntype) f. 8. *Euphydryas editha baroni* (not a syntype) f. 9. *Poladryas minuta arachne* (label on specimen "minuta Colo."; not a syntype of *arachne*) f. 10. *Poladryas minuta arachne=ab. gunderiae* (Beulah, Colo., H. Nash)(actually named in 1930, not on p. 128)(called type on p. 128 & on plate, thus a Holland 1931 lectotype designation, as Pelham Catalogue is vague as to whether O.D. named a holotype; however this is certainly a unique specimen, so it is the holotype by monotypy, as the original description clearly states "The type is a female.") holotype f. 11. *Poladryas minuta monache* m. 12-13. *Poladryas minuta minuta* (not syntypes), neotype m, f. 14. *Poladryas minuta arachne* (not nymph; not a syntype) f. 15. *Poladryas minuta nympha* (Chih., Mex.) f. 16-17. *Chlosyne gabbii*, m, f. 18. *Poladryas minuta arachne=gilensis* holotype f. 19-20. *Chlosyne palla calydon* (p. 125 & plate say both are types, so neither is Holland designation of lectotype) syntypes, m, f. 21-22. *Chlosyne palla palla*, m, f. 23. *Chlosyne sterope sterope* (designated lectotype by Holland 1931) lectotype f. 24. *Chlosyne palla palla=eremita* (or weak ssp.) f. 25-26. *Chlosyne whitneyi whitneyi=malcolmi*, m, f. 27. *Chlosyne sterope acastus* (not a syntype as text states) f. 28-29. *Chlosyne hoffmanni segregata* paratypes, m, f. 30-31. *Chlosyne cyneas*, m, f.

Explanation of Plate LIX. 1. *Euphydryas chalcedona chalcedona* f. 2. *Euphydryas anicia capella* (not *nubigena*, not a type, not Calif., prob. Colo.) m. 3-4. *Euphydryas anicia brucei* (p. 118 & plate say both are types, thus neither is Holland designation of lectotype; Brown designated this lectotype), lectotype m, paralectotype f. 5. *Chlosyne theona theona* m. 6. *Chlosyne leanira fulvia* (Pelham Catalogue wrongly says this is lectotype designated by Holland 1931; actually p. 131 says pl. XVI fig. 17 is type but plate says nothing, while plate says pl. LIX fig. 6 is type but p. 131 says nothing, thus two bugs are called type in book, therefore neither is a Holland lectotype designation) lectotype (designated by Brown) m. 7-8. *Chlosyne theona thekla* (not syntypes of *bolli*), m, f. 9-10. *Chlosyne leanira leanira*, m, f. 11. *Chlosyne harrisii harrisii=albimontana* holotype m. 12-13. *Chlosyne harrisii liggetti* (p. 126 & plate says both are types, thus neither is Holland lectotype designation) syntypes, m (Pelham Catalogue calls this holotype), f. 14. *Phyciodes (Anthanassa) texana seminole* f. 15-16. *Microtia elva elva*, m, f. 17. *Chlosyne erodyle* m. 18. *Chlosyne lacinia crocale=rufescens* f. 19-20. *Chlosyne californica*, m, f. 21. *Microtia (Texola) elada callina* ("Sonora" in error, evidently S Mex.)(on p. 131-2 Holland clearly designates this f the lectotype in this book [F. Brown 1965 J. Lep. Soc. 19:209 discusses this situation and figures both female and male syntypes], because Boisduval's O.D. indicated the TL is Sonora and the X on this female's label is his indication that "the specimen was to be returned because he needed it to hold the name in his own collection.", thus Barnes & McDunnough's [1916] designation of the Mexican syntype male as lectotype is invalid, and Holland designated the female as lectotype/holotype; however there were two syntypes so Brown thought there was no holotype and a lectotype was needed, but because Bdv. indicated Sonora was the TL and Bdv. placed the X on the female's label and not the male's label indicating the Sonora female was his choice of type, the female should be considered the holotype as Higgins [1960] declared it) holotype f. 22. *Phyciodes mylitta arida* (plate says type, but p. 137 says this is just "a typical specimen" thus is not the holotype and is not a lectotype designation by Holland; it is probably a syntype; J. Rawlins selected another male in CM with "type 7020" label that he thinks is the true holotype, see *Papilio* [N.S.] #10; a female in CM also is labeled type) syntype m. 23. *Boloria improba improba* f. 24. *Polygonia oreas oreas* (not a syntype) f (uns should be checked to be positive it is *oreas*, though ups looks like *oreas*). 25. *Argynnis (Speyeria) zerene gunderi* m. 26. *Polygonia faunus rusticus* (Calif.; not a syntype) m. 27. *Polygonia oreas oreas neotype* m (should check uncus, as this photo looks like possibly *zephyrus*, Brown's photo of remounted ups and uns is too dark on uns though ups looks like *oreas*). 28. *Anartia amathea fatima* m. 29. *Eunica tatila tatila* m. 30. *Limenitis archippus archippus=ab. pseudodorippus* (Pelham Catalogue says there are only "syntypes", making this a Holland 1931 lectotype designation, however it is an extreme aberration thus is surely a unique specimen that is holotype by monotypy, as Holland 1898 wrote that he has the original Strecker type) holotype f.

Explanation of Plate LX. 1. *Historis odius* m. 2. *Historis acheronta* m. 3-4. *Myscelia ethusa*, m, f. 5. *Dynamine serina dyonis* m. 6. *Asterocampa clyton texana* paratype f. 7-8. *Boloria titania grandis* paratype (properly syntype at most, as there is no holotype or lectotype), 2f. 9-10. *Boloria titania ranieri* paratypes (properly syntypes, as there is no holotype or lectotype), m, f. 11. *Paramacera xicaque allyni* m. 12. *Boloria improba improba* m. 13. *Boloria tarquinius tarquinius* (Baffin Is.) m. 14-15. *Boloria eunomia nichollae* holotype m, allotype f. 16. *Coenonympha tullia ampelos* (Ore.) m. 17-18. *Coenonympha tullia ampelos=columbiana* paratypes, 2m. 19-20. *Coenonympha tullia inornata=benjamini* (weak ssp.) paratypes, m, f. 21-22. *Coenonympha tullia kodiak=yukonensis*, (p. 182 says both specimens are types, but pl. LX says fig. 21 is type & 22 is paratype, thus this is Holland 1931 designation of lectotype; Miller/Brown say there is no holotype; Pelham Catalogue wrongly says this is holotype, but O.D. has no designation of holotype; thus fig. 21 is lectotype) lectotype m, paratype (properly paralectotype) f. 23-24. *Coenonympha tullia insulanus* paratypes, f, m. 25-26. *Coenonympha tullia furcae*, m, f.

Explanation of Plate LXI. 1-4. *Erebia disa steckeri* (p. 203 & plate say all four are types, thus none are designated lectotype) syntypes, m, m, f, f. 5-6. *Erebia fasciata* paratypes (properly paralectotypes, as Miller/Brown says there is no holotype), m, f. 7-9. *Erebia fasciata form avinoffi*, holotype f, allotype f, paratype f. 10-11. *Erebia stubbendorffii ethela* (p.

205 & plate says two specimens are type, so neither is a Holland 1931 lectotype designation; Holland 1898 designated lectotype of pl. XXV fig. 18 female), paralectotype (wrongly designated lectotype by Brown) m, paralectotype f. 12-15. *Erebia stubbendorffii pawlowskii=alaskensis*, f, lectotype (p. 206 & plate call this type, a Holland 1931 lectotype designation) m, paratype (properly paralectotype, as there is no holotype) m, paratype (properly paralectotype, as there is no holotype) m. 16. *Erebia discoidalis discoidalis* m. 17. *Erebia mancinus* m. 18-19. *Erebia vidleri*, m, f. 20. *Erebia epipsodea form brucei* f. 21-23. *Erebia rossii kuskokwima* (weak ssp.)(two specimens are called type on plate [none in text], thus neither is Holland designation of lectotype), m syntype, f syntype, m probable syntype. 24-27. *Erebia epipsodea*, f (allotype of syn. *remingtoni*), m (syn. *rhodia*), f (syn. *rhodia*), m (holotype of *remingtoni*). 28-30. *Erebia dabanensis youngi*, lectotype (called type on p. 205 & plate, which is Holland's 1931 designation of lectotype, as there was no holotype--dos Passos' later designation of this male as lectotype was therefore invalid) m, allotype (called type on p. 205 but allotype on plate; properly paralectotype) f, paratype (called paratype on p. 205 & plate; properly paralectotype) f.

Explanation of Plate LXII. 1. *Oeneis nevadensis* (pale, so ssp. *iduna*?) m. 2. *Oeneis nevadensis nevadensis* m. 3. *Oeneis nevadensis gigas* m. 4. *Oeneis calais ivallda* (syntype; p. 194 says two specimens are types, but only this one called type on plate, thus this female is Holland's 1931 designation of lectotype) lectotype f. 5. *Oeneis* (*nevadensis*?) *macounii* (not a syntype) m. 6. *Oeneis nevadensis iduna* (p. 194 says both specimens are type, but plates say only this one is type, thus a Holland 1931 designation of lectotype; TL is required to be locality of lectotype which is "Mendocino") lectotype m. 7. *Oeneis calais calais* (Rupert House, James Bay, Que.)(which would be a valid Holland 1931 lectotype designation, except there was a holotype, which Miller/Brown say is lost, but Pelham Catalogue evidently says this is holotype) holotype f. 8-9. *Oeneis chryxus*, f, m. 10-11. *Oeneis jutta alaskensis* (both specimens are called types on p. 195 & plate, thus neither is a Holland lectotype designation) evidently paralectotypes, m (may be lectotype, as Masters & Sorenson later designated lectotype male in CM), f. 12. *Oeneis uhleri uhleri*, probably male. 13-14. *Oeneis uhleri uhleri=varuna* (N.D.)(several specimens are called types on p. 196 and on plate, thus none is Holland lectotype designation), lectotype m, paralectotype f. 15-16. *Oeneis alberta alberta*, m, f. 17. *Oeneis jutta jutta?* (maybe *O. norna?*, but does not look like *norna* photos)(Finland) m. 18. *Oeneis jutta alaskensis* (Alaska) f. 19-20. *Oeneis polixenes katahdin* paratypes (properly paralectotypes, as there is no holotype), m, f. 21. *Oeneis polixenes subhyalina=peartiae* holotype m. 22. *Oeneis jutta jutta* (Finland) m. 23-24. *Oeneis melissa assimilis* 2m. 25-26. *Oeneis melissa* (looks like ssp. *gibsoni*) "oeno", f, m.

Explanation of Plate LXIII (too pale). 1-3. *Oeneis melissa assimilis=semplei*, lectotype (p. 200 says all 3 figs. are types, but on plate only fig. 1 is called type, and the other two are called allotype and paratype, so this is a Holland 1931 *semplei* lectotype designation) m, allotype (properly paralectotype) f, paratype (properly paralectotype) m. 4-5. *Oeneis melissa beani*, m, f. 6-7. *Oeneis polixenes polixenes*, m, f. 8. *Oeneis melissa semidea* m. 9-10. *Oeneis melissa gibsoni* (p. 200 says fig. 9 is allotype & fig. 10 is holotype, and the "sp. nov." implies that this is the O.D., and Miller/Brown say HT is in CM; however Pelham Catalogue says O.D. was really in 1930 in Ann. Carn. Mus. 20:51, which evidently designated no holotype; so, fig. 10 is a Holland 1931 lectotype designation as Pelham states), paralectotype f, lectotype m. 11. *Lethe eurydice fumosus* paratype m. 12. *Cercyonis sthenele silvestris* (not a syntype) f. 13. *Cercyonis oetus oetus* (not a syntype of *silvestris*) f. 14. *Cercyonis sthenele silvestris* (not a syntype) m. 15. *Cercyonis pegala maritima* m. 16. *Cercyonis pegala wheeleri* lectotype (designated lectotype by Holland 1931) f. 17. *Cercyonis pegala boopis* f. 18. *Cercyonis pegala nephele* (not a syntype of *boopis*) (lectotype of *incana*, which is a syn. of *nephele*) m. 19. *Cercyonis sthenele paulus* (not a syntype) f. 20. *Cercyonis pegala* (Ore., *nephele* X *gabbii*) f. 21. *Cercyonis pegala maritima= weak ssp. texana* (designated *texana* lectotype by Holland 1931 here) m. 22. *Cercyonis oetus oetus* (pale like ssp. *pallescens*) lectotype m. 23. *Cercyonis sthenele paulus* f. 24-25. *Cercyonis oetus charon* (p. 192 & plate says both specimens are type, thus neither is Holland lectotype designation) syntypes (properly paralectotypes, as lectotype is in CM), m (abnormal), f (both are too pale as whole plate is too pale). 26-27. *Oeneis polixenes polixenes* (N Labrador)(ssp. *polixenes* because Pelham Catalogue defined TL as Chateau Bay, Labrador coast), f, m.

Explanation of Plate LXIV (too white). 1. *Apodemia palmeri marginalis* paratype f. 2-3. *Lasaia narses* (not in U.S.) paratypes, m, f. 4-5. *Eumaeus toxea* (=minijas="minyas"), f, m. 6-7. *Callophrys augustinus iroides*, m, f. 8. *Callophrys irus irus* f. 9. *Callophrys polios* paratype m. 10. *Callophrys mossii mossii* (not a syntype according to text) m. 11. *Callophrys fotis fotis* (Ariz.) f. 12-13. *Arawacus jada* 2m. 14-15. *Cyanophrys longula* (not pastor), m, f. 16. *Callophrys [viridis—I have petitioned to protect sheridanii from the name viridis] sheridanii* (not a syntype) m. 17. *Callophrys affinis apama* (p. 229 & plate say this is type, a Holland 1931 lectotype designation, preceding Brown's later invalid lectotype) lectotype f. 18. *Callophrys gryneus loki* paratype f. 19. *Satyrrium auretteorum auretteorum=tacita* holotype m. 20. *Callophrys xami xami* m. 21. *Satyrrium calanus falacer* m. 22-23. *Chlorostyrymon maesites telea* 2m. 24-25. *Satyrrium californica californica* (not syntypes) 2f. 26. *Strymon melinus melinus=pudica* m. 27-28. *Satyrrium sylvinus dryope*, m (not a syntype), m. 29-30. *Ministrymon azia* 2m. 31. *Satyrrium sylvinus putnami=itys* m. 32. *Electrostyrymon angelia boyeri* (text says from Haiti) m. 33-34. *Satyrrium sylvinus putnami*, m, f. 35-36. *Strymon istapa modesta*, m, f. 37-38. *Ministrymon leda* (not syntypes), m, f. 39-40. *Lycaena cupreus cupreus* (p. 252 & plate say both are types, thus neither is Holland lectotype designation), male (not a syntype), paralectotype female. 41-42. *Lycaena epixanthe phaedra=amicetus* Holland 1931, paralectotypes of *amicetus*, m, f. 43. *Lycaena helloides* (Winnipeg) m. 44. *Lycaena florus "castro" florus* (not a syntype) m. 45. *Lycaena phlaeas hypophlaeas=ab. fasciata* paratype m. 46. *Callophrys irus hadros* paratype m. 47-49.

Lycaena mariposa charlottensis, holotype m, paratype m, allotype f. 50. *Lycaena hermes* (broken tails fixed on printing plate)(not a syntype) neotype m.

Explanation of Plate LXV (too white). 1-3. *Lycaena nivalis nivalis* (not syntypes of syn. *ianthe* despite text), m, f, f. 4-5. *Lycaena dione*, m, f. 6-8. *Satyrrium auretteum*=*spadix*, m, f, m. 9. *Strymon avalona* (p. 241 says paratype [properly paralectotype]) m. 10. *Strymon melinus atrofasciata* m. 11. *Satyrrium favonius autolyceus* m. 12. *Phaeostrymon alcestis alcestis* m. 13. *Phaeostrymon alcestis oslari* syntype (p. 237 & plate says type, so would be a Holland lectotype designation except Pelham Catalogue & Miller/Brown says it is holotype, which is in USNM) holotype m. 14. *Satyrrium auretteum* (not tetra, not a tetra syntype), m. 15. *Satyrrium favonius ilavia* pale aberration, paratype (evidently syntype, as there is no holotype or lectotype) f. 16. *Satyrrium saepium saepium*=*provo* paratype m. 17. *Satyrrium saepium saepium*=*chlorophora* paratype m. 18. *Plebejus icarioides icarioides* m. 19. *Plebejus icarioides icarioides*=*mintha* (not a *mintha* syntype according to Brown, so cannot be Holland lectotype designation) m. 20-21. *Plebejus icarioides lycea*, m (not a syntype, and now missing), f (S Colo.). 22. *Plebejus icarioides fulla* (not a syntype)(misabeled "Mt. Hood" according to Brown) m. 23. *Plebejus icarioides fulla*=*ardea* (not a syntype, despite text p. 259)(Cal.) m. 24. *Plebejus icarioides evius* (Pasadena Cal.) f. 25-27. *Plebejus icarioides blackmorei* paratypes, m, f, m. 28. *Plebejus saepiolus saepiolus* (Mendocino Cal.) m. 29. *Plebejus saepiolus amica* (Victoria BC, syn. "insulanus") m. 30-31. *Plebejus saepiolus amica*=*insulanus* (text says paratypes) 2m. 32-34. *Plebejus saepiolus hilda*, paratype m, paratype f, m. 35. *Plebejus icarioides pheres* (San Fran.) m. 36-38. *Plebejus icarioides pembina*, m, f, m. 39. *Plebejus saepiolus* (looks like ssp. *gertschi*) f. 40. *Plebejus icarioides lycea* (Ariz.)(not "helios") m. 41-43. *Glaucopsyche lygdamus xerces*, m, form *polyphemus* m, form *behrii*=*mertila* holotype f.

Explanation of Plate LXVI. 1. *Plebejus icarioides pardalis* f. 2-3. *Glaucopsyche lygdamus columbia*, paratype m, m. 4-6. *Glaucopsyche lygdamus oro*, m, f, m. 7-8. *Glaucopsyche lygdamus couperi*, m, f. 9. *Glaucopsyche lygdamus xerces* form *antiacis* f. 10-11. *Glaucopsyche lygdamus xerces*, m, f. 12. *Plebejus samuelis* (det. Nabokov & Brown)(not a syntype of *scudderii*) m. 13. *Cupido amyntula* (ssp. *amyntula* or *maritima*?)(the name *kodiak* belongs to *P. saepiolus*)(Fort Yukon, Alaska) m. 14. *Plebejus atrapraetextus alaskensis*, f (Rampart, Alaska). 15. *Plebejus scudderii scudderii*=*subarcticus* paratype, m (Great Slave Lake region). 16. *Plebejus anna anna* (not a syntype of *annetta*)(Nordin, Donner Pass, Cal. according to Brown) m. 17. *Plebejus melissa melissa* lectotype (lectotype designation by Holland 1931 here) f. 18-20. *Plebejus melissa melissa*, m, f, m. 21-23. *Plebejus anna anna* (not syntypes or paratypes despite p. 264 & plate), neotype m, f (not syntype), m (not syntype or paratype). 24-25. *Plebejus acmon*-group, f, m. 26-28. *Plebejus alupini alupini* (not syntypes), m, f, m. 29. *Plebejus alupini alupini* (not *chlorina*, not syntype of *chlorina*) m. 30. *Plebejus chlorina monticola* (not a syntype of *chlorina*, contrary to text) m. 31-33. *Plebejus emigdionis*, m, f, m. 34. *Plebejus neurona* paratype m. 35. *Euphilotes battoides battoides* f. 36. *Euphilotes (battoides) glaucon oregonensis* paratype m. 37. *Euphilotes bernardino bernardino* paratype m. 38. *Euphilotes glaucon intermedia* paratype m. 39-40. *Euphilotes rita rita* paratypes, 2m. 41-43. *Euphilotes ancilla ancilla* (p. 266 says all 3 are paratypes), m, f, f. 44. *Euphilotes mojave mojave* (this would be a Holland 1931 lectotype designation, but Pelham Catalogue says this is holotype) holotype m. 45. *Philotes sonorensis extinctis* m. 46. *Euphilotes spaldingi spaldingi* paratype m. 47. *Glaucopsyche piasus sagittigera*=*catalina* m. 48. *Glaucopsyche piasus daunia* m. 49-50. *Plebejus podarce podarce*, f, m. 51. *Zizula cyna* (not a syntype contrary to text p. 271) m.

Explanation of Plate LXVII. 1-3. *Hemiargus hanno gyas*, m, f, m. 4-5. *Hemiargus hanno zachaeina*, m, f. 6. *Plebejus shasta* (could be ssp. *minnehaha*) m. 7-9. *Plebejus optilete yukona*, m, f, m. 10. *Celastrina neglecta gozora* (Honduras) m. 11-12. *Cupido amyntula herrii*, m, f. 13. *Celastrina neglectamajor* (Pittsburgh, Penn.) f. 14. *Ascia josephina krugii* (Puerto Rico) m. 15-16. *Ascia monuste phileta* (mating pair), m, f. 17. *Ascia monuste monuste* (syn. or ssp. *crameri*, holotype m, O.D. Ann. Carn. Mus. 1931 20:225-265, TL Surinam, according to J. Calhoun Holarctic Lepid. [1997]) holotype m. 18. *Ascia monuste raza* (S Baja Calif., see note 262 of Miller/Brown) paratype f. 19. *Phoebis agarithe* f. 20-21. *Colias pelidne skinneri* (p. 296 & plate say both are types, so neither is Holland lectotype designation) evidently syntypes (now paralectotypes), m, f. 22-23. *Phoebis statira floridensis*, m, f. 24-25. *Pieris (napi) hulda hulda* (not syntypes), m neotype, f. 26. *Pontia callidice occidentalis* form *calyce* holotype m. 27. *Euchloe hyantis* (p. 285 & plate say type, which would be a lectotype designation by Holland 1931, as Pelham Catalogue lists it. And Brown mentioned that it could possibly be a syntype if any exists. This is a tough decision, but because Brown's lot #4 of non-syntypes had the same "Hyantis Cal" label as this fig. 27 specimen, there is no proof that this is a syntype. And Brown wrote that it did not fit Edwards' description and lacked Edwards' "type" which Edwards wrote across two specimens that are evidently syntypes. So I accept only Brown's lectotype.)(doubtfully a syntype) m. 28. *Euchloe olympia* (not a syntype) m. 29. *Pontia callidice nelsoni* holotype m. 30. *Colias tyche boothii* (Pelham Catalogue suggests? that this is lectotype, which is in USNM) lectotype m.

Explanation of Plate LXVIII. 1. *Zizula cyna* holotype f. 2-4. *Echinargus ammon ammon* (not *filenus*, the Cuban ssp. of *H. hanno*), f, f, m. 5-6. *Lycaena phlaeas feildeni*, m, f. 7-9. *Plebejus glandon franklinii*=*suttoni*, syntypes (p. 269 says all three are types, and two of them are called type on plate, so not a Holland lectotype designation; it was named in this book and Miller/Brown & Pelham Catalogue say there is no holotype), syntype (Holland surely intended this to be holotype) m, syntype (Holland labeled it female so surely intended it to be allotype but it is male) m, paratype m. 10-12. *Lycaena epixanthe epixanthe* m, f, f. 13. *Colias philodice philodice melanic* f. 14-15. *Colias occidentalis christina* (not syntypes),

m, f (both coll. Geddes, probably Manitoba). 16-17. *Colias occidentalis occidentalis* (p. 293 & plate says both specimens types thus neither is Holland designation of lectotype) syntypes, m (atypical with orange uph spot, and little black uph smudge, and no upf black dash), f (could be christina). 18-19. *Colias harfordii*=barbara (p. 294 & plate says both specimens are types thus neither is Holland designation of barbara lectotype) syntypes, m, f. 20-21. *Colias harfordii* (p. 294 & plate says both specimens are types thus neither is Holland designation of lectotype) syntypes, m, f. 22-23. *Colias edwardsii edwardsii*=emilia (not emilia syntypes), m, f (this could be pseudocolumbiensis). 24-25. *Colias edwardsii edwardsii* (p. 295 & plate say both are types, thus neither is a Holland designation of lectotype), lectotype m, paralectotype f. 26. *Colias occidentalis christina* (paler ups form)(not astraea syntype) m (Banff, Alta.). 27. *Colias occidentalis sacajawea* (not astraea syntype) f (Judith Mts. Mont.). 28-29. *Colias palaeno palaeno* (Lapland), m, f. 30-31. *Colias palaeno chippewa*, m (now missing), f (lectotype of chippewa W. H. Edwards [not Kirby] & lectotype of helena W. H. Edwards). 32. *Colias nastes gueneei* (Southampton I.)(rossii looks like nastesXhecla & is a syn. of *C. tyche boothii*) m. 33. *Colias tyche boothii* form chione (p. 299 says type, so this would be a Holland 1931 designation of lectotype of chione, except plate says it is from Southampton Is., and p. 299 says “Chione originally described from Boothia Felix is not uncommon in Baffinland and is quite common on Southampton Island.”, thus this is not a syntype because those were from Boothia Felix; Pelham Catalogue says syntypes are in UMO & USNM) m.

Explanation of Plate LXIX. 1-2. *Parnassius eversmanni thor*, m, f. 3-6. *Parnassius phoebus apricatus*=golovinus, lectotype (plate says type, so a Holland 1931 designation of golovinus lectotype, and there is no holotype) m, m, allotype (properly paralectotype) f, f. 7-8. *Parnassius clodius* ssp. “claudianus” (ssp. pseudogallatinus from BC?—the ssp. of *P. clodius* are all rather worthless), m, f. 9-10. *Parnassius clodius* weak ssp. menetriesi, m, f. 11. *Parnassius clodius* weak ssp. altaurus (aberrant) m. 12-13. *Parnassius phoebus smintheus*=sayi (not syntypes), m, f. 14. *Parnassius phoebus hermodur*=invalid name “minusculus” (Miller/Brown note 186, an aberration) m. 15. *Parnassius phoebus hermodur* (ssp. somewhat uncertain)=alt. f. nanus m.

Explanation of Plate LXX. 1. *Battus devilliers* m. 2. *Papilio glaucus rutulus*=arizonensis (p. 319 & plate say type, thus a Holland 1931 designation of lectotype), lectotype m. 3. *Papilio glaucus canadensis* form arcticus, paratype (properly paralectotype, as there is no holotype) f. 4. *Papilio glaucus glaucus*=australis m. 5. *Papilio machaon oregonia* (p. 316 says paratype, but not a syntype) m. 6-7. *Papilio glaucus canadensis*, m (Longue Lac near Nipigon Ont.), f (Great Slave Lake). 8. *Papilio polyxenes coloro* (or *P. p. asterias* form pseudoamericus) (but exactly resembles my *P. polyxenes americus* from Colombia, so mislabeled?), m (Ariz.). 9. *Papilio indra pergamus* (p. 317 says type so evidently syntype; Pelham Catalogue says this is holotype, which is in AMNH) holotype m. 10. *Papilio aristodemus ponceanus* f. 11. *Eurytides celadon* m. 12. *Papilio polyxenes asterius*, form curvifascia, paratype m.

Explanation of Plate LXXI. 1-2. *Dryas iulia largo*, holotype m (Clench later named this “cillene” specimen the holotype of largo; Key Largo Fla.), f. 3. *Smyrna karwinskii* m. 4. *Argynnis* (*Speyeria*) *hydaspes hydaspes* (p. 94 & plate say type, which would make it a Holland 1931 lectotype designation, however p. 95 says “I figure one of Boisduval’s types of *A. hydaspes*, which was sent by the author to W. H. Edwards, and is now in my possession.”, indicating that there are several types, so this is just a syntype not a lectotype designation; dos Passos & Grey designated a male lectotype in USNM, but there is a female syntype, so fig. 4 may be that female syntype) syntype f. 5-6. *Argynnis* (*Speyeria*) *mormonia mormonia* (not syntypes), m, f. 7. *Polygonia faunus rusticus* form silvius (Yosemite Cal., uns has few spots)(p. 150 & plate say type, the original type used & labeled by Edwards, thus a Holland 1931 lectotype designation), lectotype f. 8. *Papilio ornythion* (however, it also looks identical to *P. astyalus bajaensis* m) m. 9-10. *Cercyonis sthenele paulus* (Provo, Utah), m, f. 11. *Anteos clorinde* m. 12-13. *Neophasia terlotii*, m, f (normal f=princetonia). 14. *Anthocharis julia browningi* paratype (Miller/Brown say HT in CM, Pelham Catalogue says syntypes in ANSP implying this is just syntype) m. 15. *Anthocharis midea midea*=flavida (a syntype later designated lectotype in 1954 according to Pelham Catalogue) lectotype m. 16. *Atrytonopsis ovinia edwardsi* m. 17-18. *Megathymus streckeri texana*=albicincta (p. 402 says both specimens are types but only fig. 17 is called type on plate, thus fig. 17 is a Holland 1931 designation of lectotype; however Pelham Catalogue & Miller/Brown says fig. 17 is holotype, in CM; O.D. says only that albicincta will be illustrated in this book) holotype m, paratype m.

Explanation of Plate LXXII. 1. *Megathymus ursus ursus* holotype m. 2. *Hypothyris lycaste* m. 3. *Oeneis polixenes yukonensis* holotype m. 4. *Oeneis polixenes subhyalina*=peartiae f. 5. *Oeneis melissa assimilis*=arctica holotype m. 6. *Oeneis uhleri cairnesi* holotype m. 7-8. *Boloria tarquinius natazhati* holotype m, probably paratype f (very bad print). 9. *Boloria tritonia distincta* holotype f. 10-12. *Megathymus streckeri texana*=leussleri (p. 401 says all 3 figs are types, but only this is called type on plate, thus a Holland 1931 lectotype designation, however Pelham Catalogue & Miller/Brown say it is holotype in CM) holotype m, m (paratype, as p. 401 says type), allotype f. 13-14. *Agathymus stephensi stephensi*, m paratype, f. 15-17. *Hesperia uncas lasus* (vic. Great Salt Lake), m, f, f.

Explanation of Plate LXXIII. 1-2. *Limenitis archippus floridensis*=eros (p. 165 says fig. 2 is type of eros, plate says fig. 1 is type of eros, thus this is not a Holland 1931 designation of lectotype), lectotype (designated by Brown) m, paralectotype m. 3-4. *Erebia rossii rossii*, m, f. 5. *Euchloe hyantis lotta* (p. 285 says paratype [properly syntype, as there is no holotype or lectotype]) m. 6. *Pseudocopaodes eunus eunus* holotype m. 7. *Phyciodes* (*Janatella*) *leucodesma* f. 8-9. *Euphydryas editha hutchinsi*, f, m. 10-11. *Hesperia lindseyi lindseyi* (p. 376 & plate say both specimens are type, thus

neither is Holland designation of lectotype) syntypes, m, f. 12. *Euphyes vestris* (not “*ruricola*”)(this specimen, called *ruricola* type on plate and paratype on p. 388, coll. in Calif. by Lorquin, was designated lectotype m of *ruricola* Bdv. [by Brown, Miller, & Clench, Trans. Amer. Ent. Soc. 106:77, 1980], but the *ruricola* O.D. describes a small yellow skipper like *Thymelicus lineola*, so this lectotype is invalid, not a syntype of *ruricola*. However this photo looks rather tawny! like *Polites themistocles*, but Brown Miller & Clench identified it as *vestris*. Syst. W N.A. Butt. p. 22 concludes [correctly I think] that *ruricola* is a nomen dubium.). 13-14. *Glaucopsyche lygdamus couperi*=*afra* (plate says both specimens are type, p. 262 says only fig. 14 is type, so neither is a Holland lectotype designation because fig. 14 is not a syntype), m (doubtfully a syntype, as text says only fig. 14 is type), f (not a syntype). 15-16. *Ancyloxypha arene*, m Duenas Guatemala, f S. Geronimo (Guatemala?). 17-19. *Satyrium fuliginosa fuliginosa* (not syntypes), m, f, f. 20-21. *Hesperia leonardus pawnee*=*ogallala* paratypes, 2m. 22-24. *Colias hecla hecla*, m (probably *Colias canadensis*), f, f. 25. *Eurema proterpia* form *gundlachia* m. 26-27. *Colias tyche boothii*, f, m. 28. *Colias nastes moina* f. 29. *Colias philodice philodice albino* f. 30. *Phoebis neocypris* m.

Explanation of Plate LXXIV. 1-2. *Boloria kriemhild* (p. 107 & plate says 3 specimens are type, thus none is a Holland lectotype designation) syntype (Pelham Catalogue says lectotype, which is in FMNH) lectotype m, same m uns. 3-6. *Boloria kriemhild* syntypes, f Utah (mis-labeled Ariz. on plate), same f uns, f (mis-labeled Colo.), same f uns. 7-8. *Boloria* evidently *euphrosyne* (=andersoni, photo of USNM holotype, evidently from Europe but mis-labeled “Brit. Colo.”), holotype m. 9-10. *Chlosyne harrisii hanhami* (plate says both are type, p. 127 says only fig. 10 is type, thus fig. 10 is a Holland 1931 lectotype designation [if these are syntypes, of course], and TL is locality of this female, which needs to be determined, evidently in USNM), m, f. 11-12. *Coenonympha tullia pseudobrenda* (this male designated lectotype of *brenda* by Brown, then later declared not a syntype by Austin & Gray), m. 13-16. *Myscelia cyananthe skinneri* syntypes (properly paralectotypes), m, m uns, f, f uns.

Explanation of Plate LXXV. 1-2. *Callophrys johnsoni* (p. 230 & plate say type, a Holland 1931 lectotype designation) lectotype, f, same f uns. 3. *Parnassius clodius menetriesi ab. immaculata* (Old Faithful Geyser, Yellowstone) holotype m. 4. *Colias hecla*=*albino* form *pallida* Skinner & Mengel (Greenland)(p. 298 & plate say type, thus a Holland 1931 lectotype designation), lectotype f. 5. *Battus philenor hirsuta* (p. 313 & plate say type, thus a Holland 1931 lectotype designation), lectotype f. 6. *Papilio glaucus canadensis form arcticus* (p. 319 & plate says type, thus a Holland 1931 designation of lectotype) lectotype m. 7-7a. *Achalarus toxeus*=*coyote* (p. 337 & plate say type, thus a Holland 1931 designation of lectotype; Pelham Catalogue agrees, though Miller/Brown say HT in CM) lectotype m. 8-8a. *Codatractus valeriana*=*mysie* (p. 339 & plate say type, thus a Holland 1931 designation of lectotype), lectotype m. 9-9a. *Pyrrhocalles jamaicensis* (p. 392 & plate say type, thus would be a Holland 1931 designation of lectotype, but p. 392 suggests it is the borrowed holotype) holotype m. 10-10a. *Oeneis calais caryi* holotype (says p. 197) m. 11-11a. *Oeneis uhleri nahanni* holotype m. 12-12a. *Atrytonopsis cestus* (not a syntype) m. 13-14. *Hesperia metea licinus*=*aberration horus* (Dallas, Tex.) holotype f. 15-16. *Erynnis tristis tatusi* holotype (see p. 355) m. 17-17a-18-18a. *Problema bulenta*, m, f. 19. *Oeneis melissa assimilis*=*simulans* holotype (see p. 200) m. 20. *Oeneis uhleri cairnesi* holotype (p. 197, evidently same specimen as LXXII fig. 6) m.

Explanation of Plate LXXVI (the names of figs. 1-3 and 21-22 need updating). 1-1a. *Caria domitianus* (or maybe *C. rhacotis*?) m. 2-2a. *Caria* (not *melicerta*, maybe *Symmachia*?) “type” m. 3-3a. *Caria* “ino” (looks very little like ino; all 3 *Caria* figs. are bad) m. 4-4a. *Apodemia multiplaga* paratype m. 5-5a-6-6a. *Apodemia phyciodoides* holotype m, allotype f. 7-7a-8-8a. *Calephelis nilus*=*perditalis* paratypes (Miller/Brown say HT USNM, but Pelham Catalogue say syntypes USNM, so these may be syntypes), 2f. 9-9a. *Calephelis virginiensis*=*louisiana* holotype (Opelousas, Louisiana) m. 10-10a. *Apodemia hepburni* m. 11-11a. *Calephelis wrightii* (evidently holotype, as p. 219 says these are Wright’s figs. and Pelham Catalogue says Wright figs. of ups/uns are holotype) holotype m. 12-12a. *Satyrium favonius ontario* m (bad figure). 13-13a. *Hypostrymon critola* (p. 241 text says copy of holotype figs.) m. 14. *Satyrium auretorum auretorum* holotype m. 15. *Satyrium sylvinus sylvinus* (p. 237 & plate say type, thus a Holland 1931 designation of lectotype; fig. 15 is the female syntype illustrated by Oberthür, & does not resemble the male syntype later designated lectotype by Emmel Emmel & Mattoon [Syst. W N.A. Butt. chap. 2 p. 9], who stated that Oberthür’s fig. was of the female syntype, thus this female is the valid lectotype designation by Holland 1931, and the later lectotype is invalid) f. 16. *Satyrium calanus falacer aberration heathi* holotype f. 17-17a. *Chlosyne endeis* m. 18. *Battus philenor philenor form? acauda*=*corbis* f. 19. *Parnassius phoebus apricatus* holotype (p. 310, Kodiak I.) m. 20-20a. *Oeneis alberta daura* (p. 196 says holotype from “Mt. Graham”) holotype f. 21. *Spalgis lemolea*=*ssignata* pupa. 22. *Spalgis epius* pupa. 23. *Feniseca tarquinius* pupa.

Explanation of Plate LXXVII. *Papilio glaucus glaucus*.

Index corrections/additions: *antiacis*, *Lycaena*, 261. *apama*, *Thecla*, 229. *arizonensis*, *Codatactus*, 333. *autolytus*, *Thecla*, 238. *calais*, *Oeneis*, 195. *calydon*, *Melitaea*, 125. *catalina* [really *Echinargus thomasi*], pl. XXX fig. 45. *couperi*, *Glaucopsyche*, 262. *critola*, *Thecla*, 241. *gilensis*, *Melitaea*, 129. *hanhami*, *Melitaea*, 127. *ianthe*, *Anthanassa*, pl. XVIII fig. 12. *inornata*, *Coenonympha*, 183. *kuskoquima*, *Erebia*, 203. *orcas*, *Hesperia*, pl. L fig. 15m. *oslari*, *Thecla*, 237. *taumas*, *Polites*, 381. *Thecla croesioides*, 225. *zampa*, *Systasea*, 344.

CORRECTIONS TO F. MARTIN BROWN'S PAPERS ON THE TYPES OF BUTTERFLIES NAMED BY WILLIAM HENRY EDWARDS (these papers appeared in Trans. Amer. Ent. Soc., volumes 90-113, 1964-1987, the Hesperidae papers coauthored by Lee D. Miller, and a few lectotypes authored by Paul A. Opler & Walfried J. Reinthal)

In the corrections that follow, I list the current name of the butterfly, then volume & page of Brown's text in Trans. Amer. Ent. Soc., then details of the correction. Some of the corrections are minor (typos, switched figures, etc.), but I have corrected many of the type localities, including six that were wrongly designated as "Archer Co. Texas" by Brown. The wrong specimen was illustrated as lectotype in one case. Other miscellaneous corrections are made.

Dozens of the lectotypes designated by Brown are invalid, because they were preceded by lectotype designations made in Holland's 1931 & 1898 Butterfly Book and 1915 Butterfly Guide, as noted above, and referenced below. Holland 1898 made 3 and Holland 1915 made 6 and Holland 1931 made 41 lectotype designations for Edwards' names. Brown credited Holland for 3 of these lectotype designations, credited dos Passos & Grey for 5, and Higgins for 1, while Brown incorrectly took credit for 24 lectotype designations, and Brown & Miller incorrectly took credit for 16, which were actually designated by Holland. But in most of these cases, Holland's and Brown's lectotypes are the same specimen, so the only change is who designated the lectotype. (Of course, the first designation of a specimen as lectotype is valid, and all later designations of that specimen or any other specimen are invalid.) Only in 13 cases are the later invalid lectotypes different specimens from Holland's.

Obviously, Brown's work was still valuable in doing the museum study and historical research to determine which specimens were syntypes and which were not, etc. Brown proved that numerous Holland "types" were actually not even syntypes.

When comparing the butterflies illustrated in Holland's 1931 Butterfly Book, with the same specimens illustrated by Brown, it unfortunately becomes obvious that many of the specimens have sustained considerable damage to wings, antennae, and sometimes to abdomen. Evidently repeated handling of the specimens, and perhaps dermestid beetles, have taken their toll. Thus public museums may not be the safest places to house butterfly specimens.

Papilio machaon bairdii form *hollandii*. 101:17, *hollandii* is not a "backcross", it is just a *bairdii* with yellow modifier genes making the side of the abdomen widely yellow. The abdomen of *bairdii* is basically black with 2 ½ rows of yellow dots, but varies from having yellow dots with no yellow wash, to more and more wash, and finally all yellow between the dots. 101:18, Holland 1931 designated same valid lectotype as Brown's invalid lectotype.

Papilio machaon brucei. 101:21, Holland 1931 designated a different valid lectotype pl. 40 fig. 4 female than Brown's invalid lectotype.

Papilio machaon oregonia. 101:22, neotype is female not male.

Papilio zelicaon form *nitra*. 101:26, Holland 1931 designated same valid lectotype as Brown's invalid lectotype.

Papilio glaucus rutulus=*arizonensis*. 101:28, Holland 1931 designated same valid lectotype as Brown's invalid lectotype.

Eurytides marcellus f. *walshii*. 101:29-30, fig. 10 & Holland's fig. is *walshii* holotype.

Euchloe olympia=*rosa*. 99:35, Holland 1931 designated a different valid lectotype pl. 32 fig. 39 male than Brown's invalid lectotype.

Euchloe hyantis hyantis. 99:38, 2nd paragraph line 6, "if any exist."

Anthocharis julia stella. 99:41, Holland figured "types" on pl. 32 fig. 35 male & 36 female. 99:43, Barnes & McDunnough 1916 (Contr. Nat. Hist. Lepid. N.A. 3[2]:62) wrote they "would restrict" the type to a male from Yosemite agreeing with O.D. & labeled type in red ink in Edwards coll. in CM; the word "would" means that this is not a valid lectotype designation, and that specimen was apparently not mentioned on 99:43 so may be lost.

Anthocharis julia julia. 99:44, the male fig. by Holland on pl. 32 fig. 34 is NOT "based on" the lectotype, because the black fw bar and hw spots differ greatly between the two. P. 45, *julia* & *thoosa* are different.

Anthocharis cethura pima. 99:47, Holland 1931 designated same valid lectotype as Brown's invalid lectotype.

Colias behrii. 99:52, Holland 1931 designated same valid lectotype as Brown's invalid lectotype.

Colias philodice eriphyle. 99:84, TL is actually spelled Lac la Hache.

95:179, 99:118, 106:68, 106:88. James Wilson (not Willis) Tilden.

Coenonympha tullia ampelos. 90:342, Holland figured *ampelos* on pl. 60 fig. 16 f, but his figs. on plate 25 figs. 21 & 30 were *insulanus* from Vancouver I.

Coenonympha tullia pseudobrenda. 90:346, Brown's (invalid) *brenda* lectotype was fig. by Holland on plate LXXIV figs. 11-12. But Austin & Gray 1998 (chap. 47 in Syst. W N.A. Butt) wrote that this was not a syntype of *brenda*.

Cercyonis oetus charon. 90:366, Holland's figs. were pl. 26 figs. 11 m & 12 f, & pl. 63 figs 25 f & 24 m.

Cercyonis pegala maritima=*texana*. 90:374, Brown correctly indicated that Holland 1931 validly designated this same specimen as lectotype.

Cercyonis pegala wheeleri. 90:387, Brown correctly indicated that Holland 1931 validly designated this same specimen as lectotype.

Oeneis (nevadensis) macounii. 90:387, Nipigon is in Ontario. 90:390, Holland 1915 The Butterfly Guide evidently designated the valid lectotype, a male on his pl. LXXVI that may be the same specimen as Holland's 1931 pl. XXVII fig. 3m, which Brown evidently treated as a syntype.

Oeneis nevadensis iduna. 90:392, Brown correctly indicated that Holland 1931 validly designated this same specimen as lectotype.

Oeneis polixenes brucei. 90:398, Holland 1931 designated a different valid lectotype pl. 27 fig. 7 male than Brown's invalid lectotype.

Erebia stubbendorffii ethela. 90:403, Holland 1898 The Butt. Book p. 211 designated a different valid lectotype pl. XXV fig. 18 female than Brown's invalid lectotype.

Gyrocheilus. 90:408-9 spells this *Gyrocheilus* (why?).

Argynnis (Speyeria) cybele carpenteri has not been found in the Sangre de Cristo Mts. of S Colo. as 91:240 claims.

Argynnis (Speyeria) nokomis. 91:251-2. Washington Co. Utah has ssp. near *apacheana*. The specimens from Lower Rio Florida and Hotchkiss were both females. Ojo Verde is only ~10 mi. SW of LaSal and is just a small desert wash & gauging station without permanent water that does not now and surely never had a colony of *nokomis*. 91:254 change *nokomis* lectotype to neotype.

Argynnis (Speyeria) zerene hippolyta. 91:267, Holland 1931 designated the valid lectotype that dos Passos & Grey later accepted as lectotype.

Argynnis (Speyeria) zerene bremnerii. 91:271, Holland 1931 designated same valid lectotype as dos Passos & Grey's invalid lectotype.

Argynnis (Speyeria) callippe macaria. 91:283, Holland 1931 designated same valid lectotype as dos Passos & Grey's invalid lectotype.

Argynnis (Speyeria) adiaete atossa. 91:293, Holland 1931 designated a different valid lectotype pl. 13 fig. 12 male than dos Passos & Grey's invalid lectotype.

Argynnis (Speyeria) atlantis atlantis. 91:298. Holland 1915 The Butterfly Guide designated the valid lectotype, a male on his pl. XIV, the same syntype specimen later fig. by Holland 1931 on pl. X fig. 9m which Holland 1931 & Brown called a paratype. This lectotype is from Hunter, Catskill Mts., Greene Co. NY, therefore dos Passos' later 1935 restriction to Hunter was unnecessary.

Argynnis (Speyeria) hesperis electa=nikias. 91:308. In *Papilio* (New Series) #8 I showed that *electa* belongs to *S. hesperis* (not *S. atlantis*) and corrected the *electa* TL to Twin Lakes, Lake Co., where Mead collected and *electa* occurs (it does not occur at Turkey Creek Junction).

Argynnis (Speyeria) mormonia bischoffii. 91:316 claims that Holland's pl. 11 fig. 7 male is a syntype, which is an error as p. 320 says an 1874 letter indicated that both syntypes were destroyed; so a neotype was designated. L. Paul Grey later corrected the TL to Anchorage.

Argynnis (Speyeria) mormonia opis, 91:322, Holland 1931 designated same valid lectotype as dos Passos & Grey's invalid lectotype.

Argynnis (Speyeria) mormonia erinna. 91:324, Holland 1931 designated same valid lectotype as Brown's invalid lectotype.

Boloria "epithore". 91:334, Holland's pl. 15 fig. 17 is *Boloria frigga*, so fig. 17 is definitely a "pseudotype" also.

Boloria astarte astarte. 91:340, Holland's figs. are pl. 18 fig. 14 f pseudotype and fig. 15 m victoria holotype.

Boloria titania butleri. 91:346, Holland 1931 designated same valid lectotype as Brown's invalid lectotype.

Euphydryas chalcedona colon. The O.D. on 92:361 (*Papilio* 1:45-46) was changed to Bull. Brooklyn Ent. Soc. 3:80 in Miller/Brown (Brown explains this change on 93:320-321). 92:363, Holland 1931 designated same valid lectotype as Brown's invalid lectotype.

Euphydryas chalcedona paradoxa=perdiccas. 92:364, O.D. is listed as *Papilio* 3:43 1883 in Gunder's revision, and listed as Bull. Brooklyn Ent. Soc. 3:80 1881 in Miller/Brown (Brown explains this change on 93:320-321). N. Kondla states that the O.D. gave TL as "Puget Sound". 92:366, nobody now believes that it ever occurred at Tenino as Brown wrote; this was another mislabeling by Herbert Morrison.

Euphydryas anicia brucei. 92:366, Holland's figs were pl. 59 fig. 3m-4f.

Euphydryas editha baroni. 92:372, Holland's pl. 16 fig. 7 is *E. chalcedona*, so is not a syntype *baroni*.

Microtia (Dymasia) dymas. 92:380. Holland 1931 designated a different invalid lectotype pl. XVI fig. 18 female than Brown's invalid lectotype. But actually Holland 1915 The Butterfly Guide designated the valid lectotype, a female on his pl. XX fig. 4f which Holland 1931 pl. XVI fig. 18f fig. and called type & Brown called paratype. It is from "Texas" (TL is still San Antonio).

Microtia (Texola) elada perse. 92:386. Actually Holland 1915 The Butterfly Guide designated the valid lectotype, a male on pl. XX fig. 3, which Holland 1931 fig. on pl. XVI fig. 19m as type & Brown later invalidly designated lectotype. Holland 1931 & Brown later invalidly designated the same specimen as lectotype.

Chlosyne elada ulrica. 92:385, Holland 1931 designated same valid lectotype as Higgins' 1960 invalid lectotype and Brown's invalid lectotype.

Chlosyne leanira fulvia. 92:390 TL declared to be Archer Co. Tex., but the bugs do not occur there so Archer Co. cannot be the TL, the TL was really “West Texas”. The type was coll. by Jacob Boll, as were five other taxa named by Edwards whose TL Brown declared to be Archer Co., specifically *Pyrgus philetas*, *Polites vibex brettoides*, *Yvretta carus*, *Amblyscirtes eos=comus*, and *A. eos=nilus*. All of these (except *comus*) were coll. by Jacob Boll, and all their TL’s were declared by Brown to be Archer Co. But none of these species actually occurs in Archer Co. (except *A. eos*), so the Archer Co. TL of these five bugs is obviously an error, so **I hereby correct and restrict the TL of all six (*fulvia*, *philetas*, *brettoides*, *carus*, *comus*, *nilus*) to the Davis Mts. in West Texas**. Note here that Jacob Boll has been confused with Jacob Doll at times (Edwards’ O.D. of *brettoides* mistakenly claimed that Doll not Boll collected it, and Brown himself mistakenly wrote Doll instead of Boll as the collector of *philetas*), but 106:45 makes it clear that Jacob Boll and not Doll collected all five, as Boll coll. in Texas 1876-1880 and died 1880, whereas Doll did not collect in Texas then and died in 1929 thus lived much later. (Maybe we can call this Boll-Doll confusion the bollworm—how droll.) 92:392, *fulvia* has 3 gen. in Colo.

Chlosyne sterope sterope=hewesii. 92:406, David Bauer thinks these two are synonyms (and they have the same TL).

Chlosyne sterope sterope. 92:407, Holland 1931 designated same valid lectotype as Brown’s invalid lectotype.

Poladryas minuta. 92:411 & 417, *minuta* and *arachne* are not sympatric in the foothills and lower mountains in SE Colorado, as Brown claims.

Poladryas minuta nympha. 92:417, Brown’s TL of vic. Fort Grant is impossible as only *arachne* occurs there; *nympha* occurs in Mex. (and a weak version of it N of Sonoita in extreme S Ariz.—bugs with a strong cream postmedian band like the lectotype occur only in Mexico). P. 419 states that H. Morrison collected material in Mexico just S of Arizona, therefore **I hereby correct the TL to NW Mexico including Sonora, Mexico**, where Morrison collected and the phenotype with cream ups median band actually occurs and where Holland (1931) described many specimens (the cream band does not occur near Fort Grant or in Colo.). This is another Herbert Morrison mislabeling of bugs, as “Southern Arizona.”

Phyciodes cocyta selenis=marcia. 92:432, Brown’s lectotype of *marcia* is invalid because it is not a syntype, and additionally it lacks diagnostic antennae, is female not male, and thus cannot be identified to species. This was amply discussed by J. Scott in *Papilio* (New Series) #10 & #13, and by Ronald Gatrell in *The Taxonomic Report* 4:1-19.

Phyciodes pallida pallida. 92:446 another Herbert Morrison mislabeling, of *pallida* from “southern Colorado.”

Phyciodes graphica=vesta. 92:461, TL is New (not Neu, though the original German must have been Neue) Braunfels. Holland’s figs. are pl. 17 figs 17-18-19.

Polygonia faunus hylas. 93:339, Holland’s figs. are pl. 19 fig. 7 (stated on p. 340 to be a pseudotype) & fig. 8 (stated on p. 341 to be possible syntype). 93:340 last line, change pl. 19 fig. 5 to pl. 19 fig. 8.

Polygonia faunus rusticus form silvius. 93:347, Holland 1931 designated same valid lectotype as Brown’s invalid lectotype.

Limenitis archippus obsoleta. 93:366, Holland figures pl. 7 fig. 5 as *obsoleta*, and mentions *hulstii* in text (and on plate in 1st ed. 1898), but does NOT mention *eros*. Brown notes that this specimen is holotype *hulstii*.

Asterocama celtis antonia. 93:377-9, Reinthal’s valid lectotype was fig. by Holland pl. 23 fig. 12 (note smudge on left fw apex and antennae and wings).

Asterocampa celtis montis. 93:380-383, Brown’s valid lectotype was fig. by Holland on pl. 23 fig. 11 (compare the two photos).

Asterocampa clyton flora. 93:389. Holland 1915 *The Butterfly Guide* evidently designated the valid lectotype, a male specimen on pl. L fig. 1, which may be one of the 2m2f syntypes in CM, and was not fig. by Holland 1931. The TL should be determined from the specimen in CM. Brown’s later invalid lectotype was fig. by Holland on pl. 23 fig. 1 (compare the photos).

Apodemia virgulti mejicanus. 94:117, Holland’s pl. 28 fig. 4 has unh orange like *mejicanus*, and fig. 5 is from Ariz. so is also *mejicanus* (*cythera* does not occur in Ariz.).

Apodemia virgulti duryi. 94:121, Holland 1898 *The Butt. Book* p. 230 designated same valid lectotype as Brown’s invalid lectotype.

Calephelis nemesi. 94:126-129, Brown writes that Holland figured *nemesi* on pl. 28 fig. 15, and lectotype photo does look like Holland’s fig. 15, but later 94:129 Brown claimed that pl. 28 fig. 15 is *australis*; evidently Brown thought the specimen is *australis*.

Lycaena nivalis nivalis=ianthe. 95:169, Brown must have meant to designate the TL as the E (not W) side of Lake Tahoe, as only the E side is in Nevada.

Lycaena florus. 95:172-174, Garnett’s (not Garrett’s) Rancho is really near Lundbreck at mouth of Crowsnest Pass (not near Didsbury), and Bird & Ferris (*Can. Ent.* 111:637, 1979) declared Brown’s lectotype invalid (because it is not a syntype) and designated another valid lectotype.

Lycaena cupreus cupreus. 95:176, Holland figured *cupreus* on pl. 64 (not 65) figs. 39-40. In *Papilio* (New Series) #12 I corrected the TL to Tioga Pass, Tuolumne Co. Calif.

Hypaurotis crysalus crysalus. 96:22-23, lectotype is female, not male.

Satyrium calanus calanus=wittfeldi. Holland 1915 The Butterfly Guide designated the valid lectotype, a female on pl. LXXI fig. 2 from Indian River Fla.[Georgiana, Brevard Co. Fla.], which was later fig. by Holland 1931 pl. XXIX fig. 19f, a different specimen than Brown's later invalid male lectotype.

Satyrium sylvinus putnami=itys. 96:40, Holland 1898 The Butt. Book designated same valid lectotype as Brown's invalid lectotype.

Satyrium sylvinus dryope. 96:49, "Saucelito" is now spelled Sausalito.

Ministrymon leda form ines. 96:53, Holland 1931 designated same valid lectotype as Brown's invalid lectotype.

Callophrys affinis affinis. 96:64-65, Holland's pl. 30 fig. 3f is Callophrys affinis perplexa from Calif.

Callophrys viridis. 96:66, Holland's pl. 30 fig. 2 has orange on unf & green unh so is probably C. a. perplexa.

Callophrys viridis. 96:66, line 2 should be p. 227 not p. 127.

Callophrys affinis apama. 96:69, Holland 1931 designated a different valid lectotype pl. 64 fig. 17 female than Brown's invalid lectotype.

Plebejus melissa melissa. 96:372-375, Brown's lectotype is invalid (and his TL restriction to vic. Twin Lakes and La Plata Peak is wrong also), because J. Scott (in Papilio [New Series] #12, 2006) showed that Holland 1931 Butt. Book already validly designated the melissa female on pl. 66 fig. 17 as lectotype. Scott restricted TL to Tinytown, Jefferson Co. Colo., where another syntype is from.

Euphilotes glaucon. 96:406-409, the male on Holland's pl. 30 (not 31 as listed on p. 408) fig. 39 is not Plebejus anna as Brown wrote on p. 408. It is Euphilotes, and thus probably is glaucon as Holland identified it. 96: 408 says it is [or was] in CM labeled "Glaucan male/Nev. Morr.", but Brown said that the specimen currently possessing that label (which has many pinholes) is anna, so evidently that label was switched from the Holland Euphilotes specimen (which is now lost? or mislabeled) after Holland photographed it, to an anna specimen. The male collected by Herbert Morrison cannot be a syntype (Brown noted it was a pseudotype) because Brown wrote that the source of the original types was "From 2m, 1f, taken by Mr. Henry Edwards."

Glaucopsyche piasus daunia. 96:412, Holland's fig. was pl. 66 fig. 48.

Glaucopsyche lygdamus. 96:425, Holland's pl. 30 fig. 35 looks like columbia, fig. 36 looks like couperi.

Celastrina nigra. 96:424-425, the O.D. was Edwards' Butt. N.A. where it was described and figured. W. T. M. Forbes later gave nigra species/subspecies status in 1960.

Celastrina argiolus. 96:430, Holland's pl. 67 fig. 13 is Celastrina neglectamajor.

96:432 misspelled Roy O. Kendall.

96:433 misspelled Vladimir Nabokov.

Zestusa dorus. 101:602, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Thorybes drusus. 101:605, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Cogia caicus moschus. 101:607, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Systasea zampa. 101:613, Holland's fig. was pl. XLVI, fig. 1.

Erynnis juvenalis clitus. 101:617, Holland's fig. was pl. XLV, fig. 8.

Pyrgus philetas. 101:623-624, holotype coll. J. Boll. Brown mistakenly wrote Jacob DOLL here, when he obviously meant Jacob BOLL, as here and on 92:391 Brown quoted Geiser as saying that Jacob BOLL (NOT DOLL) collected in N-C Texas 1876-1880; 106:45 confirms this. Brown restricted the TL to Archer Co. Tex., but the bugs do not occur there, thus TL Archer Co. Tex. is an error (Brown made this error on six names total), as noted above under Chlosyne leanira fulvia, so **I correct the TL of all six including philetas to Davis Mts. of West Texas.**

Pyrgus xanthus. 101:624, Holland's fig. was pl. XLVII, fig. 15.

Hesperopsis libya lena. 101:645, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

101:643 Desatoya Mts.

Carterocephalus palaemon. 103:262, Mendocino locale was no doubt correct, because ssp. or syn. "magnus" occurs there.

103:261, spelled mandan.

Copaeodes aurantiaca. 103, 270, Holland's fig. was pl. XLVII fig. 9, LI fig. 40.

Polites (Yvretta) rhesus & P. (Y.) carus figs. are reversed, on 103: pp. 275 & 276.

Polites (Yvretta) carus. 103:275-277, lectotype coll. Jacob Boll. Brown restricted the TL to Archer Co. Tex., but the bugs do not occur there. Thus TL Archer Co. Tex. is an error (Brown made this error on six names in total), as noted above under Chlosyne leanira fulvia, so **I correct TL of all six including carus to Davis Mts. of West Texas.**

Pseudocopaeodes eunus eunus. 103:277 lists the TL as bottoms of Kern River near Bakersfield, according to Morrison (who generally refused to give good locality information), but this TL is an error as the bug does not occur there (more mislabeled specimens from Herbert Morrison). **TL is hereby restricted to Victorville, the TL of wrighti.** Holland's fig. was pl. LXXIII fig. 6.

Stinga morrisoni. 103:280, Holland 1931 designated a different valid lectotype pl. LII fig. 15 male than Brown & Miller's invalid lectotype. Brown & Miller wrote that this may be a syntype but were not sure, but I am treating it as a syntype given by Edwards to his brother-in-law Theodore Mead, as happened with Problema byssus (see 106:63) etc.

Hesperia attalus. 103:282, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Hesperia meskei straton. 103:285, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Hesperia comma harpalus. 103:288 & 290. J. Scott (Papilio [New Series] #11, pp. 7-8, 2006) corrected the TL of harpalus & cabelus to the higher Sierra Nevada W of Carson City (far enough west to encounter the syn. "yosemite" phenotype found in the high Sierra). Holland's pl. LII fig. 20 is *Polites sabuleti*. On 103:291, Brown meant to label as 1) that part of 2nd paragraph reading "Cabelus...1975.", and he meant to label as 2) that part of 2nd paragraph reading "The third...upside down." 103:260, spelled cabelus.

Hesperia comma harpalus=cabelus. 103:291, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Hesperia comma "oregonia" (harpalusXidahoXhulbirti). 103:291-293. J. Scott (Papilio [New Series] #12 p. 69) showed that oregonia is a useless name that evidently applies to an intergrade zone mess like that in Trinity Co. Calif. illustrated by A. Shapiro (J. Res. Lep. 29:35).

Hesperia comma idaho. 103:293, J. Scott (Papilio [New Series] #11 p. 7-8) showed that idaho applies to the lowland ssp. widespread in W U.S., and restricted the TL to the lowlands at Doyle in Lassen Co. Cal. 103:295, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype. Brown & Miller (103:294) wrongly claimed that Barnes & McDunnough (1916, Contrib. Nat. Hist. Lepid. 3 #2:127) designated this specimen earlier as lectotype; B. & McD. actually wrote only that they "would restrict the type"; they did not actually restrict the type; Holland designated the valid lectotype in 1931.

Polites vibex praeceps=brettoides. Spelled vibex not vibe. 106:44-45, lectotype coll. Jacob Boll (not Jacob Doll as O.D. claimed). Brown restricted the TL to Archer Co. Tex., but the bugs do not occur there, thus TL Archer Co. Tex. is an error (Brown made this error on six names in total), as noted above under *Chlosyne leanira fulvia*, so **I correct TL of all six names including brettoides to Davis Mts. of West Texas**. 106:45, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Polites sonora siris. 106:46, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Polites origenes rhenae. 106:47, the rhenae TL must be Wet Mts. W of Pueblo, because it does not occur on the plains (where Pueblo occurs). **I hereby restrict the TL to Hardscrabble Canyon drainage S of Greenwood, in the foothills of the Wet Mts., Custer Co. Colo.**, where rhenae occurs.

Polites mystic dacotah. 106:52, Georgetown is a little too high altitude for this species, so the **TL is hereby restricted to Georgetown or eastward toward Idaho Springs**, in Clear Creek Canyon W of Denver, Colo.

Polites mardon. 106:53 last line, male fig. Butt. Book pl. 47 fig. 26 not 28. 106:54, Holland 1931 designated a different valid lectotype pl. LIII fig. 12 female than Brown & Miller's invalid lectotype.

Anatrytone logan lagus. 106:61, TL designated as Oak Creek Can., "Custer" Co. Colo. Actually there are two Oak Creek Canyons, in Fremont Co. (one in the foothills of Wet Mts., the other S of Cotopaxi in Arkansas Canyon), of which the former is more likely, while there is no Oak Creek Can. in Custer Co. But I have not found it in S-C Colorado where both canyons occur. Maybe it once occurred there?, as Neumoegen was presumably reliable (Herbert Morrison was totally unreliable on locality data).

Problema byssus byssus. 106:63, Holland fig. the lectotype on pl. 53 fig. 38 (not 28). And Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Ochlodes agricola agricola=milo. 106:69, TL was not Tenino Wash., as the species does not occur N of S Ore. Another mislabeled specimen from Herbert Knowles Morrison.

Poanes taxiles taxiles. 106:74, the TL is Oak Creek Canyon in Fremont (not Custer) Co. Colo., as noted above under *Anatrytone logan lagus*. **I hereby restrict the TL to the Oak Creek Canyon in foothills of Wet Mts. in Fremont Co.**

Euphyes vestris. 106:77, Emmel Emmel & Mattoon (1998) found that the *ruricola* O.D. does not match *vestris*, so this *ruricola* lectotype is evidently invalid.

Euphyes dion dion. 106:80-82. Allyn not Alyn in Fig. 28 legend. Lectotype was fig. on Holland's pl. LIII fig. 36 (not 38 as p. 82 writes).

Euphyes pilatka. 106:83. In 2), Holland's "Ind River" female must have had a label reading Butterfly Book/Pl. 53 fig. 35 (not 36), because fig. 35 shows pilatka while 36 shows dion.

106:85, Hans Herman Behr, not H. H. Berr.

113:30, Tryon (not Tyron) Reakirt. I read that Herman Strecker never returned any loans (that's called theft).

Atrytonopsis python python. 113:32, the male fig. by Holland pl. LIV fig. 4 is the holotype; Brown & Miller mistakenly figured the paratype female (the female on Holland's LIV fig. 5) as the holotype and obviously meant to illustrate the male (read Brown's Holotype paragraph).

Atrytonopsis pittacus. 113:33, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Atrytonopsis hianna deva. 113:37 fig. 6 is neotype not holotype, as noted on p. 38.

Amblyscirtes phylace. 113:40 restricts the TL to foothills of Wet Mts. probably along Hardscrabble Crk., but I did not find it in the Wet Mtn. foothills, and found it only at Pass Creek Huerfano Co., so the **TL should just be left as "So. Colo."** The fig. 8 holotype is female.

Amblyscirtes eos=comus. 113:43, Brown restricted the comus TL to Archer Co. Texas, an error (even though eos occurs there--Brown made this error on six names in total) as noted above under Chlosyne leanira fulvia, so **I correct the TL of all six names including comus to Davis Mts. of West Texas.** The valid TL is the locality of the neotype, but the neotype was labeled only "West Texas", not Archer Co. Texas, thus Archer Co. was NOT the locality of the neotype as Brown wrote. The original comus (lost in the ship sinking) was coll. by Mr. Belfrage rather than Jacob Boll, though the neotype was coll. by Boll.

Amblyscirtes eos=nilus. 113:44-45, holotype coll. Jacob Boll. Brown's restricted the nilus TL to Archer Co. Tex., an error (even though eos occurs there--Brown made this error on six names in total), as noted above under Chlosyne leanira fulvia, so **I correct the TL of all six names including nilus to Davis Mts. of West Texas.**

Amblyscirtes cassus. 113:52, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Notamblyscirtes simius. 113:54, Brown listed the TL as Oak Creek Can. Custer Co. Colo. (Neumoegen), and Pueblo, Pueblo Co. Colo. (Lintner), then mistakenly restricted it to Oak Creek Can. There is no Oak Creek Can. in Custer Co., while there are two in adjacent Fremont Co.; but Brown should have restricted it to Pueblo, Pueblo Co. Colo., because this is a shortgrass prairie species that doubtfully occurs in the mts. in Oak Creek Can., but is common on the shortgrass prairie around Pueblo. So **I hereby correct the TL to Pueblo.** Also, Holland 1931 designated same valid lectotype as Brown & Miller's invalid lectotype.

Panoquina ocola. 113:61, Brown could not find the museum female illus. by Holland pl. LIV fig. 22, probably because it is actually Lerema accius f.

113:67, Part II Section III was published in 1987 on pp. 113:29-69.

113:68, Corbet not Corbert.

Commentary on research on old names and types is needed here. Brown worked hard for 23 years (1964-1987), and must have spent at least \$20,000.00-\$40,000.00 in NSF government and personal funds, to fix the types of Edwards' butterflies, in a pile of published papers 3" thick. With that time and money he could have done a lot of good biological research, such as finding new hostplants, studying mate-locating behavior, etc. It's too bad that the ICZN Code's obsession with old names (the principle of priority) forces so much waste of time and money. In all other fields of science, old bad work is simply happily forgotten. But in taxonomy, old bad work hangs around forever like a rotten stench, and taxonomists are forced to dive into the old-name sewer and muck around in the murky sludge to dredge up horrid things (toxotaxa), and then sanitize them a little to make them pass minimum standards (see the cartoon at the end of Papilio [New Series] #18). The principle of priority is the diving board into the old-name sewer. If we could somehow get rid of the ICZN Code's obsession with priority and types, we could winch our taxonomists out of the old-name sewer and redirect them into useful biological work, instead of wasting their time on legalistic interpretation of old papers and old names. There are lots of butterflies that need work rearing and photographing eggs larvae pupae, studying the contact areas between dubious species, doing morphology to define genera objectively, etc., work that taxonomists could do if they weren't wasting their time correcting old bad work. We would be better off if the principle of priority were abolished. Without the principle of priority, confusing rotten names such as dumetorum could be instantly ignored and forgotten, as merit & popularity rather than priority would decide our scientific names. A few species might have several different scientific names in use simultaneously (which happens even today!), but this situation would be temporary, and soon the names would stabilize, and the names would be appropriate! Eliminating the principle of priority would carry over onto common names; currently lepidopterists retain horrible inappropriate names (such as Plebejus lupini—now alupini--whose larvae and adults have nothing to do with lupines) merely because they have priority and were in long use; with unshackled correct thinking, these misleading common names could be corrected, finally. If scientific names became unshackled from the principle of priority--free at last—they would greatly improve.

A CATALOGUE OF THE BUTTERFLIES OF THE UNITED STATES AND CANADA, with a complete bibliography of the descriptive and systematic literature.

Jonathan P. Pelham. 2008. Journal of Research on the Lepidoptera 40:1-652.

This outstanding book is very useful and welcome for butterfly experts. It updates the names of North American butterflies since the last catalogue/checklist by Lee Miller & F. Martin Brown in 1981, which was partially updated by a list of changes compiled by C. Ferris in 1989. The title is a misnomer, as the book actually is a catalogue of the NAMES, not the butterflies, as the book attempts to update and correct the ICZN status of all the proposed names (the biological status of hundreds of the actual butterfly taxa is arguably wrong, so one cannot really call it a catalogue of the living biological creatures). Thus the book represents a North American update of the Charles Bridges catalogues of worldwide names in 1988. The book is a good value, 652 pages for only \$25 (though if I were editor I would have put each name into just one paragraph and used 3 columns in index and two columns in the checklist [Appendix 3] and deleted blank lines in appendices, reducing the book to 400 p.).(The blank spaces can be used to write corrections & additions.) For each name, the book gives literature citation of the original description, type locality, designated types and museum location, and--the best feature of the book--the "original combination" of each name below the genus. Using the original combination, now we can know whether the bug was named as a "form" or "variety" to judge whether it is an infrasubspecific unavailable

name or an available name; and finally we can determine the original spelling and determine whether its ending has been sexually transmogrified by the suffix-changers to conform with ICZN Code requirements for making the sex of genus and species names match. Pelham has spent years of time and lots of \$ to examine original descriptions to record all this information. The book lists unavailable names after a double-cross, incl. homonyms after a double-cross symbol as in dos Passos' 1964 checklist, and lists infrasubspecific names in quotes after the same double-cross; I would have preferred a different symbol for infrasubspecific names (such as an odd-font I), but one can examine the original name combination and the other listed info to determine why the names listed after the double-cross are now considered unavailable for use by the ICZN Code. The book lists type locality and types in a separate line for each name; for lectotypes & neotypes the Code requires the TL to be the locality of lectotype or neotype, so it would have been nice for bugs with lectotype/neotype to write at the end of the type locality line "TL now locality of lectotype/neotype," and list the locality. Publications—especially journals—often had wrong dates on their covers, and Pelham has evidently determined corrected publication dates for many or most of these, as noted in the bibliography entries. The book is very thorough. My copy of Miller/Brown has corrections/additions on almost every page, so I compared those to the Catalogue and found that nearly all have been fixed/added. For instance, Miller/Brown mistakenly used the wrong name on six homonyms (arcas, palaemon, sophia, alticola, alaskensis, augustus), and Pelham corrected all of them.

P. VII says that the species-level status of the names was based on Opler & Warren (2002), which is unfortunate, because that obscure work included no subspecies, those two authors disagreed on species concepts, as one of them does not believe in subspecies and has never named any (people with that view inevitably raise all the distinctive ssp. to distinct species status); not a good foundation for the Catalogue. P. IX complains that vol. 12 of *Papilio* (New Series) lacked peer review (actually it had five times the usual number of reviewers as the regular vanity-press butterfly journals such as *J. Lepid. Soc.* & *J. Res. Lepid.*, and a review by a non-expert for those journals really results in just editorial improvement and not real improvement of the basic science), yet the Catalogue uses an unpublished thesis for its Hesperidae higher classification (though it is undoubtedly an improvement). Pelham did not have the time to read *Papilio* N.S. #12, as noted below where its findings are implemented. The higher classification for Nymphalidae uses the latest research such as new DNA phenograms such as Wahlberg et al. (2005), and seems mostly okay, though rearranged some, but I looked at various phenograms produced by that research that greatly switched positions of Papilionidae and Pieridae and Lycaenidae with small iterations of methods, which does not give one much confidence in the results, so I wonder how long the seemingly-overconfident classification down to tribes and subtribes in this Catalogue will last. Nymphalidae subfamilies are arranged contrary to important gross larval morphology (scoli etc.); Satyrinae and Morphinae should be placed near the base somewhere after Libytheinae & Danainae (see J. Scott & D. Wright 1990, *Butterfly Phylogeny & Fossils*, chap. 5 pp. 152-208 in Otakar Kudrna's *Butterflies of Europe*, vol. 2; recent DNA research has evidently proved that Apaturinae apparently lost the body scoli and belong among the Nymphalinae). Riordininae is obviously a subfamily of Lycaenidae, not a family, by characters of adults and immatures, despite some DNA data.

The Catalogue seems to be reluctant to change the status of numerous names of species and subspecies, and usually maintains the 1981 Miller/Brown status without implementing recent revisionary changes of status. Many ssp. that have been proven to be synonyms are still listed here as ssp. *Lycaena thoe* is the correct name as noted, yet *hyllus* is still used. *Sterope* is the correct name, yet *acastus* is still used. ("Repeating an error doubles it".) *Dumetorum* is the correct name for *C. sheridanii* as Warren wrote, yet *sheridanii* is still used (I like this refusal to use *dumetorum* [even though the Code states that existing usage must be maintained until the Commission rules on name suppression], because *dumetorum* has caused enough trouble by whacking *viridis* and abandoning *perplexa* and now threatening to overrun *sheridanii*, and I have petitioned the ICZN to suppress the name *dumetorum* because it upsets stability and has caused mental-meltdown confusion across W.N.A., witness A. Shapiro's 2007 field guide; my petition also protects the name *sheridanii*). Some well-studied published changes in the literature are ignored, while the book makes some drastic changes with little or no explanation, apparently based on little or no information. There are nearly 300 names that I think are given the wrong status (several hundred synonyms treated as valid subspecies [this is good news for splitters, as no matter how identical is a synonym, someone somewhere will list it as a valid ssp. until the end of eternity], some ssp. are placed in the wrong species, some valid ssp. are treated as synonyms, etc.) (the Burke Museum must not have a very good comprehensive collection of butterflies from throughout North America). And there are many cases in which the ssp. within a species have been arranged in unnatural jumbled sequence (*Apodemia* "mormo", *Celastrina*, *Boloria*, *Speyeria*, *Euphydryas*, *Lycaena arota*, etc.) (p. VIII says they were arranged by latitude/longitude rather than systematic relationship). I mention few of those hundreds of cases of wrong status or placement below, as there are too many of them to be corrected here, but most of them are mentioned in the other book reviews in this *Papilio* (New Series) #19 issue.

So, we need a new Biological Catalogue of N.A. Butterflies (not names). A Biological Catalogue would classify the wild biological creatures by morphological/behavioral/genetic etc. relationship, and reassign the nomenclature as necessary (as much as allowable in the Code) in order to best fit the real biological units, and would also include non-Code names of those forms that are important in the lives of the butterflies (for instance forms involved in seasonal adaptation, forms such as *f. alba* that provide *Colias* females in cold environments with extra nitrogen, forms involved in mimicry, important genetic forms, etc.). Pelham's Catalogue of names merely lists such forms according to how the original name combination

& description was worded, as synonyms or as infrasubspecific unavailable names etc., without mentioning whether the name is of a seasonal form or sexual form or whatever, so there is no way to tell in this Catalogue whether the name is one of those biologically-important forms. Such forms that I used in my 1986 book are listed in the Catalogue without mentioning their biological function, and are called *nomina nuda* (actually the ICZN Code says that form names are no longer regulated by its rules, so such form names that were proposed after 1960 without types etc. need not have even been listed in this Catalogue).

But for nomenclatural matters, this is an outstanding book.

The book uses original spellings for the endings of species/ssp. names (rather than matching species gender to genus gender), which is commendable in my opinion, but is frequently contrary to the latinization rules forming half the Code. Sometimes a latinized ending is also listed and called “species-genus gender conformation”, but I’m sure an expert scholar of latin could concoct a thousand more. A paper by G. Lamas is listed to justify not latinizing the endings. (I don’t like changing the endings, and my names were not intended to be latin and were concocted just to sound good, so I am annoyed when people change the endings of my names.)

The worst part of the Catalogue is Pelham’s attempt to name five new subspecies. He attempts to name all five by using unavailable names of homonyms or aberrations and raising them to subspecies status. ICZN Code Art. 13 says “To be available, every name published after 1930 must satisfy the provisions of Article 11 and must/ 13.1.1. be accompanied by a description or definition that states in words characters that are purported to differentiate the taxon, or/ 13.1.2. be accompanied by a bibliographic reference to such a published statement..., or/ 13.1.3. be proposed expressly as a new replacement name (*nomen novum*) for an available name, whether required by any provision of the Code or not.” Pelham did not publish a description or definition of any of these five names, so none of the five satisfy 13.1.1. Note that the words “the taxon” in 13.1.1. mean the taxon being named, namely the subspecies in these five cases; and the glossary defines “taxon” as a “population, or group of populations” (a taxon is not an individual such as an aberration—the glossary does define “infrasubspecific taxon”, but that means a taxon such as a subspecies that is a whole population or group of populations to which someone wrongly gave an unavailable “form” or “variety” name). Here are the five proposed names:

1) On p. 375 Pelham proposed *Chlosyne nycteis pastoron* to replace the unavailable homonym *obsoleta*. *Ssp. pastoron* is now a valid available ssp. name, because Pelham referenced the original description of *obsoleta*, which has a description and definition, which satisfies 13.1.2., making the name available.

2) On p. 118-9 Pelham proposed *Papilio indra parvindra* to replace the unavailable homonym *pygmaeus*. *Ssp. parvindra* is now a valid available ssp. name, because he referenced the original description of *pygmaeus*, which has a description and definition, which satisfies 13.1.2. making the name available. Nomenclaturally-irrelevant note: *pygmaeus* is a subjective synonym of *panamintensis* according to Steve Spomer, who has reared and studied all the *P. indra* ssp., so *parvindra* is also a subjective synonym.

For the next three names, Pelham attempted to name three new ssp. by raising the status of aberrations to subspecies status. The original description of an aberration does not qualify as the description or definition of the taxon such as a subspecies, because by definition aberrations are abnormal individuals and are thus different from the population and thus the subspecies from which they came, and aberrations are not populations or groups of populations. And the Code requires a description or definition of the taxon being named: the subspecies. Thus the original description of an aberration cannot be used to satisfy the description or definition of a taxon required by 13.1.1., or the bibliographic reference to the description or definition required by 13.1.2. Aberrations are unavailable names, thus cannot fulfill 13.1.3. either. Moreover, article 1.3.2. says “Excluded from the provisions of the Code are names proposed for teratological specimens as such”, which indicates that a description of an aberration cannot be used to satisfy the reference requirement of 13.1.2 (thus it is impossible to elevate a name that does not even enter into nomenclature; there is no name to elevate). So, raising an aberration to ssp. status mandates satisfying 13.1.1. or referencing 13.1.2., by requiring Pelham to write or reference a “description or definition that states in words characters that are purported to differentiate the taxon”, the taxon being the subspecies being named, not a freak individual within the population. Unfortunately, Pelham fails to write the “description or definition that states in words characters that are purported to differentiate the taxon”, and fails to provide a reference to such a published statement, for all three proposed ssp., as follows.

3) On p. 366 Pelham proposes the name *Euphydryas colon svilhae* as a new ssp., by attempting to raise the aberration “*Euphydryas chalcedona* race *perdiccas* tr. f. [transition form is the name Jean Gunder used for the hundreds of aberrations that he named, as Gunder explained in 1924-1928 & 1932 *Entom. News*, and W. D. Field explained in 1934 *Can. Ent.* 66:253-257, and C. dos Passos explained in 1938 *Amer. Mus. Novitates* #999] *svilhae*” to ssp. status. Unfortunately Pelham does not provide a description or definition, and only states that it is from Olympic Mts. of Wash., and references Guppy & Shepard (2001), which mapped all the bugs from W-C Wash. and Olympic Mts. to SW & W-C BC as ssp. *paradoxa*. Thus there is no description or definition, as Guppy & Shepard did not distinguish the Olympic Peninsula bugs as different, and merely referred to them as part of the *paradoxa* range. We do not know how Olympic Mts. *svilhae* is distinguished from the bugs southward or far northward, as there is no description or definition to differentiate the taxon, thus *svilhae* does not satisfy 13.1.2. or 13.1.1. (or 13.1.3.), and is an unavailable *nomen nudum*. [Nomenclaturally-

irrelevant note: I have examined 7m7f of Hurricane Mts. bugs, which look very similar to colon & wallacensis, most like the latter in wing pattern & size; they do not look like a different ssp.]

4) On p. 167-8 Pelham proposes the name *Anthocharis sara sulfuris* as a new ssp., by attempting to raise the aberration “*Anthocharis sara race julia tr. f. sulfuris*” to ssp. status. Unfortunately Pelham does not provide a description or definition, and only states that it is from SE BC, E Wash., NE Ore., N Ida. and W Montana, and references A. Warren (2005). But Warren (2005) did not say that this is a separate ssp. or new taxon, he described NE Ore. bugs as near already-named taxa, he just wrote that adults in the Warner Mts. of SE Ore. are phenotypically similar to ssp. *stella*, and wrote that NE Ore. has individuals resembling ssp. *browningi* flying with ssp. *stella*, and wrote that many in Baker & Wallowa Cos. of NE Ore. approach the phenotype of ssp. *browningi*, and wrote that ssp. *stella* phenotypes also occur in the Klamath River Can. in S-C Ore., and wrote that high-elevation Cascades populations cannot be assigned to either ssp. near *stella* or ssp. *flora*. Warren did not mention BC or Wash. or Ida. or Montana at all. So we do not know what bug Pelham was proposing, so *sulfuris* does not satisfy 13.1.2. or 13.1.1. (or 13.1.3.), thus is an unavailable nomen nudum.

5) On p. 381-2 Pelham proposes the name *Chlosyne palla blackmorei* as a new ssp., by attempting to raise the aberration “*Melitaea palla a. female blackmorei*” to ssp. status. Unfortunately Pelham does not provide a description or definition, and states only that it is from SE BC, E Wash., NE Ore. & W Montana as discussed by Warren (2005). But Warren (2005) did not mention BC, and only mentioned that dark females can be found from central Ore. to W Mont., and mentioned that females in NE Ore. “may be dark, dusky, or pale, and in some populations, dark forms predominate”, and he wrote that dark females are common in Calif. *eremita* and are found in Calif. ssp. *palla* and in SW & C Ore., and wrote that NE Ore. also has populations in the Columbia Gorge that look similar to *sterope* and are invariably mixed in with museum series of *sterope*. Thus we do not know what bug Pelham was proposing, so *blackmorei* does not satisfy 13.1.2. or 13.1.1. (or 13.1.3.), thus is an unavailable nomen nudum. Additionally, Pelham notes the range of his proposed ssp. *blackmorei* is SE BC, E Wash., NE Ore. & W Montana, yet the Gunder *blackmorei* aberration is from outside of that range in SW BC, which would seem to invalidate any use of the aberration in Pelham’s attempt to describe/define ssp. *blackmorei* for 13.1.2. [Nomenclaturally-irrelevant note: Such a subspecies should be named by persons knowledgeable about it, because the SW BC bugs probably are darker than the others.]

Thus, four of the new subspecies names Pelham proposes are worthless, and three are nomina nuda. The editor and peer reviewers of this Catalogue should have found these errors and corrected them; parts of the catalogue evidently escaped quality peer review. It is bad practice to raise aberrations to ssp. or species status, and Art. 13A recommends that a diagnosis should also be provided. I once used the name of an aberration for a new subspecies (*Euphydryas chalcedona hennei*), but my original description provided a complete description and diagnosis of the ssp., plus the type locality, and designated holotype & a hundred paratypes in museums (contrary to p. 363 which failed to provide that data and wrongly stated that *hennei* Scott is an aberration raised to ssp. status—these TL & type details are added below). And before someone renames a homonym, it is also accepted taxonomic practice to notify the author (of the name *pygmaeus* and *obsoleta*, though the author of the latter is now deceased) and give them a chance to rename the taxa.

Details of the corrections that I have found are are listed below. I cannot find all the errors; decades will be required to find most of them, because people almost never read the details about aberrations and other obscure names. (Many corrections were contributed by N. Kondla, some by C. Guppy and others.) The type localities of many butterflies especially from Colorado have been fixed, to match the historical itineraries of the collectors with the actual distribution of the butterflies, as noted in Brown’s Edwards’ types corrections above.

P. 16 I think *Thessia* is just a subgenus at best in *Achalarus*, as Steinhäuser defined it on genitalia (wings are nearly identical to *Achalarus albociliatus*) and the crude genitalia figs. in Evans of *jalapus* & other *Achalarus* aren’t any more diverse than are the species within *Thorybes*.

P. 18 *aemilia*, Holland 1898 p. 325 designated lectotype of pl. XLVI fig. 39 male.

P. 18 *dobra* is just an aberration (here’s the difference between names and biology again: nomenclaturally it is an available name for a ssp., biologically it is an aberration).

P. 18 subgenera *Erynnis* & *Erynnides* are perfectly valid in *Erynnis* (see Burns 1964 & *Papilio* [New Series] #14), and the book uses subgenera elsewhere.

P. 24 *lena*, the lectotype is Holland 1931 pl. L fig. 36 male V.

P. 31 *scudderi* does not occur in Graham Mts., so **I hereby restrict TL to Huachuca Mts., Ariz.**, where it also occurs.

P. 32 *zarucco/funeralis* belongs beside the *persius*-group, see *Papilio* (N.S.) #12.

P. 33 *borealis*, delete District of Mackenzie which no longer exists.

P. 34 *pallida*, spelled *Potrero* for 4 other names (*canescens*, *celata*, *nelphe*, *pithyusa*).

P. 38 *lagunae* holotype was not fig. by Scott (1975).

P. 38 ***caespitatis* TL is correctly still Marin Co.**, because Emmel et al. had no valid reason for changing it (obviously authors cannot change the TL willy-nilly for no reason, so the first justifiable TL is valid; without this principle people could change the TL every year just for fun [Pelham seems to agree with this principle in text of *eremita* on p. 381 & *fuliginosa* on p. 203]; see the corrections of Syst. W N.A. Butt. in this *Papilio* [N.S.] #19 issue).

P. 38 emend *macdunnoughi* to *mcdunnoughi*; a lapsus calami in art. 32.5.1. as it was named for James McDunnough.

- P. 39 apertorum is just a spring form not a ssp. (biologically it is form pseudoxanthus Scott).
- P. 40 Archer Co. TL is error for philetas (the Archer Co. errors were corrected in Brown's Edwards' types corrections in this Papilio [N.S.] #19 issue). Corrected TL is Davis Mts. in west Texas.
- P. 40, Heliopyrgus would seem to be a subgenus of Heliopetes, as the genitalia of Heliopetes purgia is similar.
- P. 43 mackenziei, delete District of Mackenzie which no longer exists in Canada.
- P. 44 Most of the Megathymini are given the wrong species/ssp. status.
- P. 44 polingi, Holland 1931 p. 361 says pl. LI fig. 20 male is paratype.
- P. 49 albocincta, holotype fig. Holland 1931 pl. LXXI fig. 17 male.
- P. 52 garita, based on Reakirt's itinerary, the **TL is hereby restricted to Mts. W of Denver Colo.**, the only place he collected.
- P. 52 edwardsii, **TL hereby restricted to Hardscrabble Can., Wet Mts., Custer Co. Colo.**
- P. 58 erna is more of a form than a ssp., its population intermediate toward linda.
- P. 58 aenus, **TL hereby clarified to Hardscrabble Can., Wet Mts., Custer Co. Colo.**
- P. 59 carolina, Holland 1898 p. 367 designated lectotype of pl. XLVI fig. 36 male.
- P. 60 Archer Co. TL is error for comus & nilus. Corrected TL is Davis Mts. in west Texas (see above in Brown's Edwards' types corrections in this Papilio [New Series] #19 paper).
- P. 61 I didn't find phylace in Wet Mts. including Hardscrabble Road, so **TL should just be "Southern Colo."**
- P. 61 Nastra larvae look almost identical to Atrytone arogos, so why are they placed in different tribes?
- P. 67 eunus Kern River bottoms TL is error as it does not occur there. TL restricted above to Victorville (TL of wrightii).
- P. 67 muertovalle is a valid paler ssp. from Death Valley to Needles, see Papilio (N.S.) #18.
- P. 70 manitoba, TL now Lac la Hache.
- P. 70 borealis, Nain is in mainland Labrador, not Nfld., in the province of "Newfoundland and Labrador."
- P. 70 colorado, **TL is evidently Kenosha Pass or Guanella Pass, Colo.** Holland 1931 p. 371 says pl. LII figs 1-2 are types, but fig. 2 is not called type on plate and fig. 1 is Hesperia nevada, thus neither are likely to be syntypes.
- P. 71 idaho, Brown & Miller 103:294 wrote in error that Barnes & McDunnough 1916 [Contrib. Nat. Hist. Lepid. 3 #2:127] designated the same specimen as lectotype that Holland 1931 & Brown & Miller designated as lectotype, which is from "East Calif."; B. & McD. actually only wrote that they "would restrict the type", leaving the actual restriction to Holland 1931.
- P. 71 cabelus, Austin's TL is error as bug is absent in the Central Valley W of the Sierras. Holland 1931 designated lectotype as pl. LII fig. 17 male.
- P. 71 harpalus, in Types, delete "18mD (paralectotype)".
- P. 74 MacNeill (1964) wrote that the lindseyi type has label "Napa Co. Calif."
- P. 76 Archer Co. TL is error for carus. Corrected TL is Davis Mts. in west Texas (see above in Brown's Edwards' types corrections in this Papilio [New Series] #19 paper).
- P. 78 mardon, Holland 1898 Butt. Book did not state that the fig. was type, so did not designate a lectotype. However, Holland 1931 Butt. Book wrote on p. 381 that both figs. on pl. XLVII fig. 26 male & pl. LIII fig. 12 female are types, but on plates only LIII fig. 12 female is called type, thus the latter female is a Holland 1931 lectotype designation, and the former male is just a syntype (now paralectotype). Both were mislabeled "Mt. Hood".
- P. 80 rhenia **TL is Wet Mts. W of Pueblo, Colo.**, see corrections of Brown's Edwards types in this Papilio issue.
- P. 80 dacotah TL restricted above to Georgetown or eastward toward Idaho Springs, as Georgetown may be too high.
- P. 81 Archer Co. TL is error for brettoides. Corrected TL is Davis Mts. in west Texas (see above in Brown's Edwards' types corrections in this Papilio [New Series] #19 paper).
- P. 81 vibex, Miller/Brown wrote TL certainly on the mainland, meaning ?Florida, & they wrote that brettus TL is Georgia (name based on Abbot drawing).
- P. 85 byssus, Holland 1931 designated pl. LIII fig. 38 (not 28) male lectotype, and pl. XLVI fig. 20 (not 41) female is paralectotype.
- P. 85 kumskaka, Holland 1931 pl. LIII fig. 39 male is Problema bulenta, not kumskaka, so is doubtfully a paratype.
- P. 86 spelled monofacies (also on p. 459).
- P. 87 aaroni, Gatrell wrote the lectotype is in CMNH not ANSP.
- P. 87 taxiles TL & p. 91 lagus TL are Oak Creek Can. in Fremont Co. not Custer Co., which is dubious for lagus anyway as the bug isn't there now.
- P. 88 howardi, Holland 1931 called pl. XLVI fig. 38 male type on p. 391 and nothing on plate, & called pl. LIV fig. 1 female paratype on p. 391 & plate; & called pl. XLVI fig. 14a nothing on p. 391 & plate; thus fig. 38 male is Holland's 1931 valid lectotype designation; Gatrell wrote that fig. 38 is a syntype, which is now lost (pl. XLVI fig. 38 male Poanes howardi in 1931 book is the same bug as Phycanassa howardi in 1898 book [Gatrell wrote pl. 46 then wrongly wrote LXVI, it is XLVI]); Gatrell designated a 2nd lectotype in CMNH with "type 7092 label, which is invalid; Holland's lectotype is valid even though lost.
- P. 88 morrisoni, Holland 1931 designated lectotype pl. LII fig. 15 male (pl. XLVI figs. 26-27 are not called types on plate).

- P. 89 **pratincola TL is correctly still Broderick, Yolo. Co. Calif.** as Scott defined it, because Emmel et al. changed it for no good reason (see the corrections for Syst. W N.A. Butt in this Papilio issue).
- P. 91 snowi, Ute Pass is Teller Co. Colo.
- P. 91 Anatrityone was actually resurrected by Scott (1992, Papilio [N.S.] #6) based on numerous larval pupal & adult traits; Burns later repeated this based on a few weak adult traits.
- P. 92 simius TL corrected to Pueblo, Pueblo Co. Colo. by Scott in corrections of Edwards types in this Papilio (N.S.) #19 issue. Change 1898 to 1931, as Holland 1931 Butt. Book designated lectotype on p. 369. Pelham Catalogue wrote that Holland 1898 Butt. Book designated lectotype, but he meant to write 1931 as simius is on p. 369 in 1931 book and on p. 341 in 1898 book which was thoroughly revised in 1931. Also, Holland 1898 did not say any specimen was type thus made no lectotype designation.
- P. 93 emended to mcguirei, not macguirei.
- P. 94 dukesi, Holland 1931 fig. a “paratype” on pl. LIV fig. 26 male.
- P. 97 python, Holland’s 1931 pl. LIV fig. 4 male is holotype (Brown & Miller mistakenly fig. the female in Holland fig. 5 & called it the holotype, they obviously indicated the male was holotype & intended to illustrate the male).
- P. 102 emend altaurus to alturus, a lapsus calami for Lake Alturus.
- P. 103 apricatus, Holland 1931 fig. holotype pl. LXXVI fig. 19 male.
- P. 103, St. Ilja is Mt. St. Elias according to p. 101.
- P. 104 golovinus, Holland 1931 designated lectotype on pl. LXIX fig. 3 male.
- P. 105 manitobaensis, Exshaw is the Alta. town, not Exsham.
- P. 105 if minusculus is from Colo., move it to sayii.
- P. 106 hermodur is a perfectly valid ssp. that is darker and genetically distinct when reared in lab, and requires no reevaluation as it is older than all the remaining sayii synonyms. The **TL is hereby restricted to alpine zone in Sangre de Cristo Mts. or Monarch Pass, in S Colo.**, where the bug is limited in southern Colorado.
- P. 108 Troidini was transferred to Parnassiinae by C. Hauser (1993), a view evidently rejected by later authors.
- P. 111 floridensis, named by Holland 1898 Butt. Book p. 307, holotype by monotypy, holotype fig. in that book & Holland 1931 on pl. XLIV fig. 2 (not 3) male; named as “Papilio ajax winter form floridensis” thus is infrasubspecific, so change = to double-cross. Delete “cited G. Edwards...Birds I:pl.34” which was not mentioned by Holland, though maybe Pelham is stating only that G. Edwards mentioned this name floridensis.
- P. 111 walshii, holotype fig. Holland 1931 pl. XLIV fig. 4 male.
- P. 112 aliaska, Pelham claims that Holland 1931 designated lectotype on p. 314 pl. 41 fig. 1 male, which if true would make hudsonianus a syn. of aliaska as Holland’s p. 314 says that fig. 1 is hudsonianus from Rupert’s House on Hudson Bay. However, Holland also says on p. 314 that he has another type from Scudder’s original material from Alaska; thus Holland called 2 specimens types, so neither is a valid lectotype designation. And fig. 1 hudsonianus is not a syntype: Holland’s fig. is from Hudson Bay, and Scudder’s aliaska syntypes are from Alaska, thus pl. 41 fig. 1 is not an aliaska syntype so cannot be lectotype. We revert to Miller/Brown, who say “TL Nulato Alaska. HT should be in MCZ.”
- P. 113 umbellatarum Fabricius must be 1738, not 1938.
- P. 114 dodi is a syn. of ssp. brucei; both are polymorphic but usually yellow, the black form varying from uncommon in the south (<5% at Glenwood Springs brucei TL) to rare northward in Canada.
- P. 114 dodi, Kondla (1981) also defined TL as Dorothy, Alta.
- P. 114 hollandii, Roaring Forks is near Carbondale, Garfield Co. Colo.
- P. 115 calverleyi, Holland 1931 Butt. Book fig. holotype on same plate XLI fig 6f.
- P. 117 rudkini, “Papilio bairdii form rudkini, f. nov.” was named as an available subspecific name by Comstock 1935 Bull. So. Cal. Acad. Sci. 34:143-144 by art. 45.6.4., because it was named as a form before 1961 and the original description described a subspecies, and Comstock labeled it a “race” on 34:143 and a “species” on 34:144. The Chermocks did not name rudkini. And even if Comstock’s text had described rudkini as infrasubspecific, it would still be a ssp. name by 45.6.4.1. because later authors raised it to subspecific rank (Pelham believes that 45.6.4.1. is valid according to p. 17).
- P. 117 comstocki. The original combination was actually “Papilio rudkini form Comstocki f. nov.” Form comstocki is an infrasubspecific name because it was described as a form like clarki.
- P. 118 phyllisae, spelled Butterbredt Peak
- P. 125 arcticus, Holland 1931 designated lectotype on pl. LXXV fig. 6 male as just “type”.
- P. 127 alexiars is a ssp. of P. glaucus (even though it has wing pattern & genitalia of rutulus) and has mtDNA similar to glaucus (F. Sperling) & was treated as a ssp. of glaucus by J. Scriber (J. Res. Lepid. 27:96-103 & 27:222-232 & Oecologia 71:94-103), & by Tyler, Brown, & Wilson (1994).
- P. 134 दौरا, add these entries: TL: “Virginia”; Types: holotype may be in BMNH (ex van Lennep colln.).
- P. 135 eugenia is not in U.S., and solana & maybe pallidula surely belong to sidonia, also not in U.S.
- P. 138 is centralis in U.S.?
- P. 142-3, why two alba names, is one a white male?

- P. 144 eriphyle is limited to the central interior BC according to C. Guppy, so all 5 synonyms do not belong to it though some may be closer to eriphyle than to philodice. TL now called Lac la Hache.
- P. 144 kootenai. Because N BC is in the range of ssp. vitabunda, the **TL is the holotype locality of Kaslo, on the Kootenai River, currently on the edge of a reservoir built since the holotype was collected.**
- P. 145 has two pallida Cockerell entries on page, both surely the same albino female form.
- P. 146-8 sullivanii was clearly proved to belong in *C. occidentalis* in O.D., *C. christina* also belongs in *C. occidentalis*, and *Papilio* (N.S.) #12 placed *wasatchia*=*pseudochristina* in *C. occidentalis* and placed *astraea* & *altiplano* into *C. edwardsii* & placed *pseudocolumbiensis* into *C. columbiensis*.
- P. 148 edwardsii was named by Behr, not Edwards, as the original description clearly states.
- P. 147 alberta, TL is just S Peace River Valley, where eurytheme was absent until the last 10 years.
- P. 148 Miller/Brown wrote that pallida is preoccupied by pallida Staudinger, *Cat. Lepid. Europe* 3 (1860), but it's infrasubspecific anyway.
- P. 149 elis, there is no holotype as O.D. did not designate one. N. Kondla states that the type series was almost certainly a mixed series of elis & minisni. Holland 1931 did not designate a lectotype, so there are only syntypes.
- P. 150 hela, the Eskimo Point in "NWT" (not the one near Churchill Man.) is now in Nunavut.
- P. 150 Scott (1986) Butts. N. A. elevated *Colias canadensis* to species level.
- P. 151 is olga in Alaska? It is a syn. of herzi in Gorbunov book, a syn. of kolosovae in Grieshuber & Lamas 2007 *Mitt. Münch. Ent. Ges.* 97:163.
- P. 152 chione, Holland's 1931 pl. LXVIII fig. 33 male is not syntype as it is from Southampton I., and Holland's p. 299 says chione was originally described from Boothia Felix.
- P. 152 *C. Ferris* Bull. *Allyn Mus.* #96 said both rossii & gueneei are hybrids *nastes*X*hecla*, but rossii is orangish & gueneei not orangish, so rossii is a syn. of boothii.
- P. 152 moina, the Eskimo Point in "NWT" (not the one near Churchill Man.) is now in Nunavut
- P. 154 gigantea, above Fort York is in Manitoba (now York Factory Manitoba), not Ont.
- P. 155 ssp. minisni Bean (1895) *Psyche* 7:228 TL Laggan Alberta is valid available name & skinneri its synonym, as N. Kondla states that Bean described minisni as a full species with a description (but Cris Guppy disagrees that the words constitute a description) & wrote it is "clearly allied to pelidne"; minisni is not based on elis f; some people consider minisni different from skinneri; Grieshuber & Lamas accepted minisni in their global *Colias* list.
- P. 156 chippewa, add Types: lectotype is same specimen as lectotype of helena.
- P. 160 sennae TL is very uncertain and must be fixed by someone.
- P. 165 Anthocharini.
- P. 170 It's hard to believe that *Euchloe ausonides* & *Pieris protodice* are in separate tribes, their larvae are so similar.
- P. 171 rosa, Holland's 1931 valid designation of lectotype pl. XXXII fig. 39 male is not the same specimen as Brown's later invalid lectotype, as the wing pattern clearly differs.
- P. 172 hyantis, Holland's fig. had no type label & does not match description, thus is doubtfully a syntype & Brown's lectotype designation of a different specimen should be considered valid.
- P. 175 terlooti is correct not terlooi, this was a simple lapsus calami of Baron Terloot's name, a wrongful substitution of an i for a t.. A lapsus calami must be corrected according to 32.5.1. Pelham has misinterpreted the Code here; every lapsus calami--every misspelling--must be automatically corrected. (The word spelling automatically involves a comparison of the current letters to the known correct usage.)
- P. 178 the marginalis ssp. a to f all seem to belong to *P. hulda* as ssp./synonyms. Passosi is at present a nomen dubium, and C. Guppy will designate lectotype to fix the name.
- P. 181 mcdunnoughi is correct emendation for James McDunnough; macdunnoughi was a lapsus calami thus must be corrected (Art. 32.5.1.).
- P. 182 hyatti is a synonym.
- P. 186 crameri, pl. LXVII fig. 17 male was called holotype by J. Calhoun *Holarctic Lepid.* 1997.
- P. 190 arethusa, author's last name was Wolley Dod, not Dod.
- P. 190 the mislabeled cupreus types are identical to lapidicola thus lapidicola is a syn. of cupreus; cupreus TL corrected to Tioga Pass, Tuolumne Co. Calif. (the Lake Co. Ore. TL listed in the Catalogue was based on a comedic chain of four compounding errors); artemisia is a valid ssp.; all by J. Scott in *Papilio* (N.S.) #12
- P. 194 clara, Holland 1931 p. 253 designated lectotype of pl. XXX fig. 26 female.
- P. 194 rutila is a homonym of European *Lycaena dispar* rutila Würneburg 1864 *Beitr. Schmett.* 1:391, 494.
- P. 195 I and others have been using the correct name thoe, and the readers should too. The bad hyllus Cramer painting resembles females of thoe, dispar, hippothoe, alciphron, and thersamon; Kořak thought it was thersamon, and it resembles *L. thersamon* more in its scalloped hw and the distinctive upf submarginal dots, and especially resembles alciphron in those traits, though the white unh resembles dispar & thoe best, and it does have a large black submedian spot in fw cell CuA₂ (but this spot is painted badly as it is larger than any *Lycaena* I have seen); this uncertainty as well as its TL of Smyrna in Asia Minor (where alciphron occurs) is sufficient reason to render it an unusable nomen dubium.

- P. 195 thoe. On Gue'rin-Me'neville's plate 81 the name was listed as thoe Boisduval, so Boisduval 1832 is the author of the name published in the 1832 Gue'rin-Me'neville work.
- P. 195 epixanthe, **TL is N.J.** as it doesn't occur at Philadelphia.
- P. 196-7 three authors revised *L. dorcas/helloides* in *Papilio* (N.S.) #12: Scott, C. Kondla, & C.Guppy showed that *L. florus* is a distinct species; Scott named *micurion*; Scott placed *arcticus*, *megaloceras*, *dospassosi*, & "castro" as ssp. of *L. florus* (Scott has petitioned ICZN to suppress the name *castro*, see *Papilio* [N.S.] #18 which fixes TL of *castro*); *hulbirti* & *sternitzkyi* belong to *megaloceras*.
- P. 198 charlottensis, holotype fig. Holland 1931 pl. LXIV fig. 47 male.
- P. 199 Miller/Brown say *minyas* is misspelling of *minijas*.
- P. 204 coolinensis with its mislabeled types is syn. of *acadica*, see *Papilio* (N.S.) #12. *Papilio* (N.S.) #20 restricts *coolinensis* **TL to "NE North America or Great Plains."**
- P. 205 sylvinus, Holland 1931 designated lectotype of Boisduval's female (Oberthür's fig.) on pl. LXXVI fig. 15 female; the later designation of male is invalid.
- P. 206 itys, Holland 1898 p. 243 designated lectotype of pl. XXIX fig 17 female, the same specimen as Brown's invalid lectotype.
- P. 206 titus, Gatrell fixed TL on 4(6):9 not 4(6):7.
- P. 207 calanus **TL obviously Florida**, the range of the taxon.
- P. 207 wittfeldi, the valid lectotype was designated by Holland 1915 *The Butterfly Guide*, a female on pl. LXXI fig. 2, later fig. by Holland 1931 pl. XXIX fig. 19f, from Georgiana, Indian R., a different specimen from Brown's invalid lectotype.
- P. 211 I still place *ilavia* as a ssp. of *favonius*, *polingi* as a distinct sp., because of wing shape, wing pattern, & stigma.
- P. 214 *perplexa* belongs in front of *affinis*.
- P. 215 spelled *albipalpus*.
- P. 215 I have petitioned ICZN to suppress *dumetorum* and it surely will be, because it is causing monstrous confusion.
- P. 216 *Papilio* (N.S.) #19 designates *paradoxa* lectotype & fixes TL etc.
- P. 218 *muiroi* & *thornei* are more ssp. of *gryneus*, even considering A. Shapiro's research.
- P. 222 *hadros*, paratype fig. Holland pl. LXIV fig. 46 male.
- P. 232 *ines*, Holland's 1931 designated lectotype on p. 240 pl. XXIX fig. 35 female, which may now be in USNM or AMNH.
- P. 233 *clytie*, holotype fig. Holland pl. XXX fig. 6 female.
- P. 239 the *sissoni* type and lowland Calif. bugs are a syn. of *E. U.S. comyntas*, and *pacnowe* is a valid ssp. from C Ore.-BC (*Papilio* [N.S.] #12).
- P. 240 ssp. *immaculata* is more valid as a ssp. than several other *E. amyntula* ssp. listed.
- P. 241 Four to six *Celastrina* ssp. must be reshuffled into their proper species.
- P. 241 *lucia*, there is a holotype by monotypy, because O.D. said "One specimen..."
- P. 245 *astenidas* TL is bad, the name is unusable nomen *dubium* until TL is fixed.
- P. 246 At least 6 *Euphilotes* ssp. are placed in the wrong species.
- P. 246 *speciosa*, Holland 1931 p. 267 says only that pl. XXXII fig. 2 female is a paratype.
- P. 256 *incognitus* not *incognita*.
- P. 256 *Orcus* holotype is an unspotted-unh aberration, evidently of ssp. *columbia* (another example of the difference between valid available ICZN names, and valid taxa; Miller/Brown listed it as an aberration).
- P. 256 *Cyclargus* is surely a syn. of *Echinargus*.
- P. 257 Scott in *Papilio* (N.S.) #12 revised *Plebejus idas/melissa* and named one new ssp.: the Ericaceae-feeding *P. scudderii* includes *aster=empetri*, *nabokovi*, *alaskensis* (but C. Guppy & C. Schmidt independently have since found that *alaskensis* eats legumes so evidently belongs to the next sp. *P. atrapraetextus*); *P. atrapraetextus* includes *ferniensis*, *longinus*, *sublivens*, *benwarner*, *annetta*, & *fridayi* [& *alaskensis*]; p. 259 Warren did not find *benwarner* in Ore., and the *benwarner* *gnathos* is not as short as that of *P. anna* (see TL dot 57 on *gnathos* morphology fig. 4 of C. Nice et al. *Molec. Ecol.* 14:1749); *P. anna* includes *anna=ricei*, *lotis*, *azureus*, & *vancouverensis*; the TINYTOWN TL of *melissa* is not an error, as Brown found syntypes from there and *Fairplay* was just Brown's first guess which he later abandoned for another county, and Warren knows nothing about this situation; the *pseudosamuelsis* Red Mtn. Inn. TL is in Lake Co. not Pitkin Co.
- P. 259 *lotis* is extinct, so needs the cross symbol.
- P. 260 *fridayi* syntypes are in CNC (N. Kondla photos), and Comstock (1927) Butt. Calif. illustrated two evident syntypes from TL Mammoth on pl. 53 figs. 21-22 mentioned in O.D. Evidently collected at high altitude (*ivallda* & *malcolmi* are alpine species with same Mammoth TL).
- P. 261 *Plebejus saepiolus* & *P. icarioides* ssp. were revised by Scott in *Papilio* (N.S.) #12.
- P. 269 Schmidt et al. in *Papilio* (N.S.) #12 assigned *aquilo* Bdv. 1832 (TL North Cape in Scandinavia, m & f syntypes in USNM) to Europe following Higgins & Riley etc.

- P. 270 Steve Kohler's finding of two species of "A. glandon" in W Mont. suggests different species status for glandon & megalo.
- P.271 virginienensis. On Gue'rin-Me'neville's plate 81 the name was listed as virginienensis Boisduval, so Boisduval 1832 is the author of the name published in the 1832 Gue'rin-Me'neville work.
- P. 271 louisiana, holotype fig. Holland 1931 pl. LXXVI fig. 9D & 9aV male.
- P. 272 I examined genitalia etc. of many rawsoni, freeman, & arizonensis to assign them the different status given in the 1986 Butt. N.A. book.
- P. 275 mormo & dumeti & mormonia, C. Gillette found A. mormo & Eriogonum hosts absent at supposed TL Davis Creek Park, so **TLs should be just Washoe Co. Nev.**
- P. 275 The Apodemia "mormo" taxa are all jumbled, see Papilio (N.S.) #18.
- P. 276 add dialeuca Opler & Powell 1962 J. Lepid. Soc. 15:167-8 TL Sierra San Pedro Martir, 5 mi. NE La Encantada, Baja Calif. Norte, holotype CAS, as it does occur in Calif.
- P. 277 mejicanus, bug does not occur near Mazatlan, so **TL is Sonora Mex. to S Ariz.**
- P. 277 duryi, Holland 1898 Butt. Book p. 230 designated lectotype of pl. 28 fig. 10 female (from 5 mi. E Mesilla...).
- P. 277 A. phyciodoides was surely mislabeled from Ariz., as K. Roever told me that the original collectors (the Dorfner? brothers as I vaguely recall) collected often in Sonora where the bug actually occurs.
- P. 277 arizona O.D. J. Res. Lepid. 26(1-4):130-2.
- P. 278 two of the five Libytheana are given wrong status.
- P. 281 Ackery & Vane-Wright 1984 synonymized strigosus to thersippus.
- P. 282 one could add after halia the name cleobaea (Godart) HELICONIA, Enc. Meth., 9:222 [1819], TL "Brasil; Antilles", STs may be in MNHN, although p. 437 says it is extralimital, so someone must have restricted its TL?
- P. 284 nigerrima, this bug lacks the median white band like sinefascia, and was fig. in Cockerell's Zoology of Colorado.
- P. 286 after angustifascia, add syn. sinefascia Barnes & Benjamin Bull. So. Cal. Acad. Sci. 25:15 (1926), TL near Prescott Ariz. restricted by F. M. Brown Trans. Amer. Ent. Soc. 93:371-372 (1967), HT in CM, raising in status of W. H. Edwards' aberration name. The holotype is an aberration of L. weidemeyerii that lacks the central white band (a second similar specimen was found E of Central City, Gilpin Co. Colo. in 2008, and nigerrima is a third).
- P. 287 burrisonii (not burrisoni) original description was clearly of a subspecies, resembling rubrofasciata whose range now overlaps that locality [however C. Guppy has not found rubrofasciata at low or higher altitude near Landsdowne=Armstrong but believes it may exist at higher altitude; he currently considers it a local ssp. of lorquini]; the word hybrid was not mentioned in O.D.; it is currently a nomen dubium. Ssp. itelka is different. C. Guppy will designate a neotype to fix burrisonii. (By the way, a name based on hybrids is an available name only if the taxon [population] is a hybrid [in other words, intermediate][Art. 17.2]; Art. 23.8 says name of individual animals of hybrid origin cannot be used for either of the parents, which is logical because there is no way to decide which parent taxon to attach the name to.)
- P. 287, eros, Holland 1931 clearly labeled pl. LXXIII fig. 1 male as the type of eros.
- P. 288 pseudodorippus, Holland 1898 wrote that he possesses Strecker's type, so Holland's 1931 fig. was surely of this specimen, so Holland 1931 designated it lectotype, except this aberration is surely holotype by monotypy.
- P. 289 G. Carpenter & B. Hobby 1944 Trans. Roy. Ent. Soc. London 94:311-346 said the generotype of Limenitis (populi) is close to bredowii in genitalia & venation.
- P. 289 hulstii, holotype fig. Holland 1931 pl. VII fig. 5 male (pl. 73 fig. 1 male is lectotype of eros).
- P. 290 Agraulis would seem to be a subgenus of Dione.
- P. 290 Warren suggested bredowii & eulalia are sympatric thus separate species but his writeup indicates that bredowii & californica could be conspecific.
- P. 297, is eunomia younger than aphirape & tomyris as the dates suggest is possible?, was there a first reviser action or ICZN suppression?
- P. 297 nichollae TL was defined by Pike in two papers as N end of Wilcox Pass; Mary Nicholl did not define TL as her paper was 20 years before O.D. And specimens from Mt. Assiniboine coll. by Dean Nicholson of Cranbrook BC are ssp. dawsoni according to N. Kondla.
- P. 300 pardopsis, holotype is female.
- P. 300 saga, bug is not in Nfld. so **TL is mainland Labrador**, in the province of "Newfoundland and Labrador."
- P. 301 in laurenti, Brighton is the actual town.
- P. 301 lehmanni, holotype fig. Holland 1931 pl. LV fig. 9 male.
- P. 304 lapponica is older than freija; was lapponica suppressed by ICZN?
- P. 305, boisduvalii, N. Kondla notes that O.D. plainly writes that it was described from countries to the east of Europe, and nothing was said of Labrador or North America. There are no syntypes available, so this is a toxotaxon and boisduvalii should be removed from the North American fauna, as there is no real evidence that it was American. Labrador bugs are ssp. grandis.

- P. 306 ranieri is the correct spelling, because Guppy & Kondla showed that ranieri and Mt. Ranier were used many times in the O.D. therefore the sole rainieri spelling in the title is a lapsus calami [probably changed by some editor] that must be corrected to ranieri (Mt. Rainier was once spelled Mt. Ranier); and their choice of ranieri over rainier constitutes first reviser selection of ranieri under art. 24.2.3. anyway; they noted the TL is in Pierce or Lewis Co.
- P. 306 sangredecristo is a ssp. in Colo. (TL) where all the females are melanic; the same phenotype is a polymorphic female form in weak ssp. ingens on Beartooth Plateau of NW Wyo. (however N. Kondla notes that in the Beartooth area ingens flies at lower altitudes and sangredecristo-like bugs fly at high altitude, which bears investigation).
- P. 309 columbia, Holland 1931 p. 88 designated lectotype pl. XIV fig. 3 male. TL now Lac la Hache & Quesnel Lake [near head of Quesnel River].
- P. 310 coerulescens, lectotype fig. by Holland 1931 pl. LV fig. 3 male ups.
- P. 310 L. P. Grey informed me that valesinoidesalba belongs to ssp. apacheana.
- P. 310, J. Scott 1982 J. Res. Lepid. 20:63 wrote: "The neotype of Speyeria nokomis was caught by Mrs. Cockerell at Beulah, New Mexico, and sold to E. Oslar, based on correspondence from T.D.A. Cockerell to F. Benjamin of the Smithsonian." Maybe Oslar could have coll. nokomis in Ouray Co., but it's doubtful. (By the way, I think the several old specimens of ssp. nokomis from Bent & Mescalero in Sacramento Mts. N.M. (fig. R. Holland J. Lepid. Soc. 62:173)(the Bent ones with no year and mixed-up month and dubious collector source) also are Oslar-mislabeled Mrs. Cockerell Beulah specimens (the Sacramento Mts. evidently has ssp. coerulescens not nokomis, as I have seen and photographed 1m of the 2m coll. by K. Roever in 1973 which is ssp. coerulescens not ssp. nokomis).
- P. 312 snyderi, Holland 1931 p. 97 said pl. XVIII fig. 6 male is paratype, which isn't a lectotype designation.
- P. 312 halcyone, spelled Hartsel.
- P. 312 bremnerii, **TL is** locality of lectotype; Holland's 1931 lectotype was labeled **Vancouver**.
- P. 314 platina, Holland 1898 p. 117 designated pl. XVIII fig. 7 male the lectotype. The TL is the locality of this specimen, though if it is vague the locality dos Passos & Grey gave would probably be usable.
- P. 314 liliana, Holland 1931 p. 98 says pl. XIII fig. 11 male is "type", a Holland 1931 lectotype designation (presuming the specimen is a syntype); Pelham Catalogue says that this specimen is possibly a fig. of the dos Passos & Grey lectotype, which is in AMNH, but dos Passos & Grey 1947 don't mention that.
- P. 316 spelled chilcotinensis (& p. 483).
- P. 318 emend macdunnoughi to mcdunnoughi; a lapsus calami in art. 32.5.1.
- P. 319 atlantis, valid lectotype was designated by Holland 1915 The Butterfly Guide, a male on his pl. XIV, later fig. by Holland 1931 pl. X fig. 9m, from TL Hunter, Greene Co. NY.
- P. 321 dennisi dos Passos & Grey Dec. 12, 1947, Am. Mus. Novit. (1370):20 [Novit. 1247 has no p. 20 & no mention of dennisi] is the valid ssp. name and helena 1957 is the synonym.
- P. 322 hydaspe, Holland 1931 figured pl. LXXI fig. 4 female syntype according to his p. 95.
- P. 323 rhodope, Caribou TL is correct, L. Grey 1989 J. Lepid. Soc. 43:4-5.; Kondla (2001) noted TL is locality of lectotype, Caribou wagon road, 2500 ft.
- P. 323 conquista, Scott 1982 J. Res. Lepid. 20:63 was the first to publish that conquista as well as zerene were caught by A. Klots in Wyo. and mislabeled NM. **TL conquista is hereby designated NW Wyo.**
- P. 323 spelled bischoffii.
- P. 325 artonis, Holland's 1931 pl. XII fig. 13 male was not a syntype according to Brown 91:326. **TL is Ruby Mts. in Elko Co. Nev.,** not Wells.
- P. 327 antonia lectotype fig. Holland 1931 pl. XXIII fig. 12 male (identical to Brown's fig.).
- P. 327, valid ssp. montis lectotype fig. Holland 1931 pl. XXIII fig. 11 male (identical to Brown's fig.).
- P. 328 flora, the valid lectotype evidently designated by Holland 1915 The Butterfly Guide, a male on pl. L fig. 1, evidently one of the 2m2f syntypes in CM; TL should be determined from its locality in CM. Later invalid lectotype fig. Holland 1931 pl. XXIII fig. 1 male (identical to Brown's fig.).
- P. 339 dentigera is dubious in U.S., based on TL of it & zerynthia.
- P. 340 orion belongs under dious as both TLs are Surinam.
- P. 345 subpallida, spelled Westcliffe.
- P. 347 thomsonii is spelling in the O.D.
- P. 347 hyperborea, Seitz' figure is of holotype by monotypy.
- P. 349, dos Passos had the fine intention of naming the darker northern ssp., but he unfortunately chose a wrong TL for marsyas & created a syn. of satyrus, as Okanogan Co. bugs are closer to TL Colo. satyrus (look at the photo of BC male in BC book p. 249, that's as pale as they get even in Ariz. & Colo.) than to N.B. transcanada, which have the uns dark & reddish in tint (much different), thus transcanada is a distinct ssp. & neomarsyas is a syn. of satyrus as Butt. Cascadia etc. treated it.
- P. 349 c-argenteum, Kirby figured the holotype, a holotype by monotypy as text said "A single specimen..." Kirby's c-argenteum holotype is an imprecise painting, whose small black spots caused it to be placed into the synonymy of Polygonia progne, but has some features that might lead some people to consider it to belong to Polygonia faunus rather

than to *P. progne* (N. Kondla notes especially the shape of the silver mark), which would upset stability of nomenclature if it were considered to be the same species as *faunus*, as it is older than *faunus*. However, Art. 23.9.1. says that prevailing usage must be maintained when 23.9.1.1. the senior synonym [*c-argenteum*] has not been used as a valid name after 1899 (I am not aware of any work that treated it as the valid name for *P. progne* or *P. faunus* since 1899, because it has been treated as a synonym of *P. progne*), and 23.9.1.2. the junior synonym [*faunus*] has been used as the valid name for the taxon in at least 25 works published by at least 10 authors in the immediately preceding 50 years including a span of not less than 10 years (*faunus* has been used by more than 100 scientific papers and butterfly books including those books reviewed in this *Papilio* [N.S.] #19 issue, those uses uniformly spread during that time with accelerated publication rate of popular books containing *faunus* lately; there are about 30 uses in *J. Lepid. Soc.* alone during this time, 30+ uses in other journals, 40+ in books, all by more than 50 authors, etc.). Therefore (23.9.2.) the name **faunus is valid as a nomen protectum that has precedence over c-argenteum, and I now take this action to permanently suppress the nomen oblitum c-argenteum for use to name the Polygonia faunus species.**

- P. 350 *oreas*, **TL hereby restricted to Redwood Park, Contra Costa Co. Calif.**, a necessary restriction because ssp. *near-silenus* occurs in far N Calif.
- P. 351 *faunus*, spelled Greene Co. N.Y.
- P. 351 Ssp. *cenveray* is a valid ssp., larger & paler than *arcticus* according to Guppy & Shepard (2001; with *cenveray* being their ssp. *rusticus*) & photos/specimens of *arcticus*; and N. Kondla finds that *arcticus* & *cenveray* are separated in range by another ssp.
- P. 352 K. Brown *Lepid. News* 1988 #2 p. 39-40 found *amathea* & *fatima* intergrading in the field in Darien Panama (relevant better evidence than lab mtDNA that has been proven to be rather useless for the study of phylogeny) and wrote that *fatima* is a ssp. of *A. amathea* Linnaeus (1758).
- P. 357 *gilletti*, Holland's fig. is pl. LVI fig. 22 male.
- P. 357 the *Euphydryas editha* ssp. are mostly jumbled in the wrong groupings (see *Papilio* [N.S.] #12).
- P. 359 *aurilacus*, Holland's 1931 fig. looks different from Gunder's fig. of holotype; one of them is probably holotype, and Holland's is probably paratype.
- P. 360 *rubicunda*, Gunder's 1929 "type" would seem to be lectotype designation.
- P. 361 *monoensis*, Holland 1931 p. 123 says pl. LVII fig. 26 male is holotype (the plate says it is paratype), but it looks a little different from Gunder's true fig. of holotype.
- P. 361 *gunnisonensis* TL Owl Crk. Pass road 2 mi. E of U.S. 550, 3.4 mi. NE Ridgway...
- P. 362 *beani*, Holland's 1931 fig. pl. XVIII fig. 13 male looks different from Gunder's type so isn't Gunder's lectotype; however Holland 1898 p. 141 said pl. XVIII fig. 13 male is Dr. Skinner's original type thus Holland 1898 designated this the valid lectotype and Gunder's is evidently paralectotype.
- P. 363 *hennei* Scott, add TL: Box Can., San Diego Co. Calif.; add Types: holotype in EMEC. My O.D. of *hennei* contained a full description/definition of adults & genitalia along with TL and a hundred types, it did NOT have the same data as Gunder's aberration *hennei*.
- P. 363 *irelandi* TL is Boy Scout Camp, Camp Wolfboro, trail near Alta Peak,...
- P. 364-5, D. Bauer (1975 *Butts. N. Amer.* [ed. W. Howe] p. 176-177) wrote that *sperryi* is a synonym of *dwinellei* and has the same moist meadow habitat etc.; Bauer should know as he was the last author to have considered *chalcedona* & *colon* to be separate sp.; the J. Emmel et al. p. 828 citation was just a checklist listing *sperryi* in *E. colon* without explanation, so was evidently wrong.
- P. 365 *colon*, lectotype designated by Holland 1931 p. 117 & fig. pl. XVI fig. 5. **TL is required to be locality of lectotype which is "Columbia River"**.
- P. 368 *alena* TL was further restricted in *Papilio* (N.S.) #12 p. 31, to summit Brian Head, Iron Co. Utah.
- P. 370 *E. bernadetta* is a separate species (see *Papilio* [N.S.] #12) that was found to be sympatric with *anicia* in Alta.-Mont.-Wyo. by S. Kohler & N. Kondla etc., yet the Catalogue wrongly lumps it into *anicia* and inexplicably raises *colon* to species status despite every author since Bauer lumping it into *chalcedona* (though J. Emmel reports *E. colon* near *wallacensis* on *Symphoricarpos* sympatric with *E. chalcedona macglashanii* on Castilleja in Pine Creek Can. in Warner Mts., in 2007 Season Summary of News of Lepid. Soc. [details were not supplied]).
- P. 370 *gilensis*, holotype f fig. Holland 1931 pl. LVIII fig. 18 female.
- P. 370, ssp. *wenatchee* is misspelled & is a very distinctive ssp. which is blackish like *colon* (*veazieae* is whitish like *bernadetta*).
- P. 371, 3 of the 5 *arachne* ssp. are misplaced, and all are ssp. of *P. minuta* (see *Papilio* [N.S.] #12).
- P. 371 *gunderiae*, Holland 1931 would have designated lectotype fig. pl. LVIII fig. 10 female, however it is an extreme aberration and is a unique specimen that is holotype by monotypy, as the O.D. states "The type is a female." TL is Beulah, Colo., coll. Herman Nash.
- P. 371 In the Edwards types material elsewhere in this *Papilio* (N.S.) #19 issue I corrected the TL of *P. minuta* nymph to NW Mexico including Sonora, Mexico, where the lectotype phenotype actually occurs (it does not occur in Graham Co.). Austin did not determine nymph's synonymy, he just named another ssp. For nymph, Holland 1931 p. 128

- called pl. XVI fig. 21 male type in text but not on plate, and called pl. LVIII fig. 14 female type on p. 128 & on plate but it cannot be lectotype as Brown wrote it is not a syntype, thus neither is a Holland 1931 lectotype designation; Holland 1898 did not designate any specimen lectotype either.
- P. 372 is a surprise, I didn't know I named *pardelina*! (I didn't--Higgins called it a form before 1961 and clearly described it as a subspecies as he described the entire series of them as equally distinctive all yellower etc., so Art. 45.6.4. clearly means that his name was a subspecies the instant he described it, thus the two Articles 10.2 & 45.5.1 concerning infrasubspecific names do not apply. So replace Scott 1986 by Higgins 1960).
- P. 372 *Papilio* (N.S.) #12 clearly proved that all the *fulvia* ssp. belong within *C. leanira*.
- P. 373 *montana*, spelled *Popocatépetl*.
- P. 374 *fulvia* TL Archer Co. Texas is error as noted in my corrections of F.M. Brown's Edwards types papers in this *Papilio* (N.S.) #19 issue (6 names have this erroneous TL). Corrected TL is Davis Mts. in west Texas.
- P. 374 *fulvia*, Brown designated the valid lectotype not Holland 1931 (Holland called 2 specimens types on p. 131 & pl. LIX, thus neither is a lectotype designation).
- P. 376 *gorgone* TL=neotype locality=River Road, Burke Co. Ga.
- P. 378 *hanhami*, Holland 1931 p. 127 says only pl. LXXIV fig. 10 female is type, which is a lectotype designation as Pelham writes it is a syntype. The TL is the Manitoba or Minnesota locality of this lectotype, which must be determined from the label on specimen evidently in USNM.
- P. 378 spelled *inghami*.
- P. 379 move *damoetas* & its ssp. *windriver* & *altalus* here, all are ssp. of *C. whitneyi*.
- P. 381 *perse*, the valid lectotype was designated by Holland 1915 *The Butterfly Guide*, a male on pl. XX fig. 3, which Holland 1931 fig. pl. XVI fig. 19 & Brown later invalidly designated lectotype.
- P. 383, *Dymasia* & *Texola* should be lumped into *Microtia* based on DNA (Wahlberg & Zimmerman 2000, *Cladistics* 16:347-363).
- P. 383 *dymas*, the valid lectotype was designated by Holland 1915 *The Butterfly Guide*, a female on pl. XX fig. 4, later fig. by Holland 1931 on pl. XVI fig. 18.
- P. 384 *callina* female was designated lectotype by Holland 1931 & figured on his pl. 59 fig. 21, but was called the holotype by Higgins. It is the holotype as it has the X on its label written by Boisduval. The Catalogue is wrong in stating that "the phenotype of the lectotype is typical of central Mexican *T. e. elada*". I have examined *C. elada* from across its range in U.S. & Mexico. *T. elada ulrica* occurs in central Mex. (Michoacan, Jalisco, Queretaro, Mexico, Tamaulipas, Nuevo Leon states) & NE Mex. & Tex., characterized by an evenly-reticulate pattern of black lines over the uniformly orange-brown ups (with smaller upf postmedian tawny spot band) that is totally unlike the *callina* type. *T. elada* does not occur in Calif. thus the Sonora label does not refer to Los Angeles Calif. One might think the locality could mean Sonora state in Mexico because ssp. *perse* from there & Ariz. has a wider ups orange-brown median band that tends toward the *callina* type (ups rather uniform lighter orange-brown). However specimens from S Mex. (Matatlan & Acatlan in Puebla, & Monte Alban in Oaxaca) are very similar to the *callina* type with wide postmedian ups band which is frequently whiter on upf, and the Matatlan female is almost exactly like *callina* holotype. So it seems clear that *callina* fits the bugs in S Mex., not central Mex. The TL of *elada* "Mexico" is bad and the *callina* type label "Sonora" definitely seems to be an error, which could cause misapplication of both names. Therefore to ensure stability and fix this mess it is necessary to fix the TLs of both *callina* and *elada* (neotypes cannot be designated because both names already have lectotypes). So, **to fix the identity of *elada* and *callina* I hereby declare the TL of both to be that area of southern Mexico where the *callina*-type phenotype is common, which with my current knowledge is "southern Mexico from Matatlan & Acatlan in Puebla to Monte Alban in Oaxaca"**.
- P. 386 move *P. picta* next to *P. phaon*.
- P. 387 *barnesi*, Holland 1898 p. 156 designated pl. XVIII fig. 5 male the lectotype, the same specimen as Brown's invalid lectotype.
- P. 388 *arizonensis*, TL restricted to Ariz. by Scott 1994, *Papilio* (N.S.) #7.
- P. 388 *tharos*, TL now Van Courtland Park, New York City.
- P. 390 *Phyciodes diminutor* (with *incognitus*) is a separate species sympatric with *P. coccyta* (*Papilio* [N.S.] #13).
- P. 391 *harperi* is an aberration of *P. b. saskatchewan*, not *P. b. batesii*.
- P. 400 *yukonensis*, the O.D. designated no holotype, thus Holland 1931 designated pl. LX fig. 21 male as lectotype.
- P. 401 *sweadneri* is a valid ssp. with paler wings.
- P. 402 *furcae*, **TL is S Rim Grand Can.**, as it is absent on N Rim.
- P. 402 *mackenziei*, N. Kondla notes that Davenport explicitly designated a holotype, though he evidently did not illustrate it.
- P. 406 the ssp. of *C. pegala* are mostly given the wrong status, and *maritima* & *carolina* are valid ssp., as noted by R. Gatrell & elsewhere in this *Papilio* (N.S.) #19.
- P. 406, A careful reading of the info in Gatrell's paper indicates that the correct name is *M. eurytris* C. Maynard 1891 (Manual N.A. Butt., Boston, De Wolfe, Fisk & Co.) who raised *eurytris* to ssp. status, and *eurytris* Fabricius is a

misspelling of the homonym eurytus. Gatrell treated *M. e. eurytris* & *M. e. viola* as a separate species from *M. cymela*, but I have 4 topotype eurytris from Berkeley Co. S.C. & 2 topotype cymela from Colleton Co. S.C. that Gatrell sent me, which look the same and are darker on ungs than Maryland cymela, so I think more work is needed to show that these are different species.

- P. 406-7, status of *maritima*, *alope*, *carolina*, *ochracea* fixed in *Papilio* (N.S.) #19
- P. 407 *incana*, lectotype fig. Holland 1931 pl. LXIII fig. 18 male (as boopis).
- P. 407 *texana*, I hereby declare *texana* a synonym of *maritima* (**first reviser action**) to make *maritima* the correct name for those who think they are synonymous (they look similar).
- P. 409 *meadii*, TL is now Bailey, North Fork South Platte River, Park Co. Colo.
- P. 409 *damei* was placed as a ssp. of *C. pegala* for a long time, and in Catalogue is placed as a ssp. of *meadii*; it actually belongs to an area of intergradation between *C. sthenele* & *C. meadii*, and properly belongs to *C. sthenele* as a reddish form gotten from intergradation/introgression from *meadii*. The TL "Grand Canyon Arizona" is rather imprecise, but the *damei* O.D. wrote that upf orangish is limited to rings around the ocelli (sometimes almost obsolete) and unf orange is much reduced compared to *meadii* and largely replaced by brownish, which fits the bugs flying at Cape Royal near the E end of N Rim of the Grand Canyon and not the variable bugs near W end, so **damei TL is hereby designated as Cape Royal on E end of N Rim of Grand Canyon** (the O.D. paper also described *Coenonympha tullia furcae* from Grand Canyon, which it is known only from the S Rim & S canyon wall, which might suggest that *damei* came from S Rim also; however J. Garth's Butt. Grand Can. NP describes S Rim [Desert View] "*damei*" as having bright red on upf & unf which means that he was describing *C. meadii*, not *damei*, so the S Rim probably has only *C. meadii*). *C. sthenele* & *meadii* form a stenochospecies, as they are one species in N Ariz.-S Utah, but are two species in Montana (see *Papilio* [N.S.] #12). *Damei* is older than all three *C. meadii melania* & *C. meadii mexicana* & *C. sthenele masoni*, so it would replace one or two of those names no matter which species *damei* is assigned to. It is doubtful that *mexicana* is the same as Utah-Ariz.-Colo. pops. of *meadii*, so *mexicana* should be restored to ssp. status by those who think *mexicana* differs from Colo. *meadii* (*mexicana* seems to be a syn. of *melania*, as both have slightly more unh ocelli than ssp. *meadii*). If *damei* is transferred to *C. sthenele*, it is older than *masoni* and would replace it, and a few reddish-unf females of *masoni* are known which could be called form *damei*. But Ken Davenport considers that *paulus* differs from *masoni* by having a slight reddish tint (due to introgression from *meadii*), thus *damei* should evidently be placed into *paulus* rather than *masoni*, and is younger than *paulus* so won't replace it. Thus to maximally preserve stability, *damei* should be considered to be just a synonym of *paulus*, where it would cause no trouble. Or, *damei* could be considered by some to be an oranger ssp. *C. sthenele damei* that is distinct from both *C. s. paulus* & *C. s. masoni*. I haven't seen the *damei* types, but according to the O.D. and the later placement of *damei* into *C. pegala*, the amount of red is small, so it would be *C. sthenele* form *damei*, which fits its placement as a syn. of *paulus*. (Here's a new strategy for preserving stability. As Pelham placed *damei* Barnes & Benjamin as a ssp. of *C. meadii*, I'm surprised he didn't also make a new entry under *C. sthenele*, as a double-crossed infrasubspecific "*C. sthenele paulus* form *damei* Scott 1986, Butts. N.A. p. 241", with the usual Pelham commentary "Proposed as a form with no nomenclatural standing; a nomen nudum." [This *damei* Scott is of course a homonym of the original *damei* Barnes & Benjamin, but that is no problem as it is infrasubspecific so homonymy does not apply.] Here I will just propose two more infrasubspecific nomina nuda forms, *C. sthenele masoni* form *damei*, and *C. meadii* form *damei*, which are infrasubspecific thus can be used within *masoni* and within *meadii melania* without replacing *masoni* and *melania*. The reader should add those two new infrasubspecific nomina nuda to the Catalogue. Such infrasubspecific nomina nuda represent a whole new strategy of preserving nomenclatural stability. The *damei* phenotype occurs within *C. sthenele paulus*, *C. sthenele masoni*, and presumably *C. (sthenele) meadii melania=mexicana*, so we need these infrasubspecific nomina nuda to keep the name *damei* from killing *masoni* and *melania* and *mexicana*. Now, readers can use the name *damei* in all three taxa, and if anyone questions the priority issue, they can just say that this *damei* is an infrasubspecific nomen nudum which does not have priority. In this role of stability savior *damei* is a "schizophrenic taxon"--a split personality--because the same name is used in several different taxa with differing availability status in them. {<<Papilio Bonus: In *Papilio* [N.S.] #18 I wrote about "pretend type localities" and "jumping subspecies" that are necessary to use in order to properly reduce three or more names to just two for the endpoints of a cline, while still obeying the principle of priority. Now we have "schizophrenic taxa" to preserve stability. Bob Pyle wrote in Butt. Cascadia that the Emperor wyomingo may have no clothes. Bob was mistaken there, but his witty remark definitely applies to a schizophrenic taxon, as we have to leave off the clothes [nomenclatural types etc.] to create a nomen "nudum" infrasubspecific name spelled the same as a ssp. or species, in order to preserve stability. Okay Bob, now let's laugh like hyenas at the pathetic gyrations that are made necessary by the ICZN Code to make words from the old-name sewer properly apply to real biological creatures. And then we can be sad that there's so much time and energy wasted on the effort. Then let's hope that in a generation or two when the existing latin-loving nomenclatural pedants depart, we can have ICZN rules that emphasize biological units first, nomenclature last. A good starting point would be eliminating the principle of priority.>>}) A third [unnecessary] solution to preserve stability, would be to petition the ICZN to suppress the *damei*.
- P. 410 *charon*, Holland also illus. paralectotypes on pl. XXVI figs. 11 male & 12 female.

- P. 410 pocus, TL now Lac la Hache, BC
- P. 411, spelled Girocheilus in F. Brown's Edwards types series, & Geirochilus by Edwards; the original spelling was Gyrocheilus?
- P. 412 kuskoquima, syntypes fig. Holland 1931 pl. LXI figs. 21 male & 22 female.
- P. 413 emend mackinleyensis to mckinleyensis, a lapsus calami (O.D. correctly spelled McKinley Park).
- P. 414 brucei is not a ssp., it is an ocelli-less form of E. epipsodea.
- P. 414 brucei, Cashier Valley & Cashier & Cashier Mts. & Bullion Mtn. were all at head of Hall Valley, Park Co. Colo., see F. Brown 1964 Trans. Amer. Ent. Soc. 90:396-7.
- P. 414, epipsodea O.D. designated no holotype, only 2 syntypes; Ehrlich 1955 designated lectotype by fig. a syntype from BMNH with the word type in fig. legend.
- P. 415 alaskensis, Holland's 1931 lectotype pl. LXI fig. 13 male is called type on p. 206 & plate.
- P. 416 ethela, Holland 1898 The Butt. Book p. 211 designated lectotype of pl. XXV fig. 18 female, from Trout Crk. in Yellowstone Park.
- P. 416 demmia TL defined as Chicago Basin, La Plata Co. Colo., by F. Brown with D. Eff & B. Rotger 1957, Colo. Butts.
- P. 416 youngi, Holland 1931 designated lectotype pl. LXI fig. 28 male. Gorbunov 2001 treated lafontainei as a ssp. of E. kozhantshikovi, and discussed Siberian specimens of E. dabanensis with [E. dabanensis] youngi valvae.
- P. 419 brucei, Holland's 1931 pl. XXVII fig. 7 male is syntype and Holland validly designated it as lectotype, despite the July 3 date (none of the syntypes were labeled Aug.); (Holland 1898 made no mention of type thus made no lectotype designation); **TL** is locality of lectotype which is **Bullion Mtn., Hall Valley...**
- P. 419 balder, Pelham's restriction of TL to Nain, Labrador, is wrong. Boisduval (1834 Icones...; copy obtained by N. Kondla who sent copy to Scott) said balder is from "Cap-Nord, au Groenland et en Islande [=Iceland].", which is his type locality, and merely wrote that he has received some specimens from Nain that do not differ from the balder that he described, so it's not apparent that he considered the Nain bugs to be syntypes of balder. Butler (1868) attributed authorship of the 1832 balder name to Boisduval and not to Gue'rin-Me'neville. Gue'rin-Me'neville (1832) (copy from J. Pelham) figured balder on plate 80 (the male ups looks like it could be European) and the bottom of the 1832 plate 80 clearly labeled it Satyrus balder. Boisd. thus attributed authorship of it to Boisduval; the 1844 text of Gue'rin-Me'neville's work listed the habitat as Cap Nord (and did not mention Nain Labrador either) and cited it as "Satyrus Balder. Boisd. Icones, etc." which seems to be an attribution of authorship to Boisduval. Considering all this, authorship is Boisduval (1832) in the 1832 work by Gue'rin-Me'neville, as Article 12.2.7. suggests that plate 80 made the name balder available by indication by illustrating it with a new name, and according to article 50.1. the authorship of balder is Boisduval as Gue'rin Me'neville stated. It has traditionally been used for European populations as a synonym of jutta. Its TL is Cap Nord on the northern tip of Scandinavia, which is near Scandinavian taiga populations of jutta. So, **I hereby correct the TL of balder to the TL of jutta (Lapland, Norway and Sweden), and any neotype of balder that is designated by someone revising O. jutta in the future should be designated as the specimen that is designated as the name-bearing type of the name jutta** (the original types of both are lost). So cross off the a, because balder is a European bug that appears to be a synonym of jutta. Balder is a toxotaxon nomen dubium, whose types cannot even be compared to Labrador specimens, because there are no specimens, just the painting. It is balderdash! Europeans created this toxotaxon, so the rotten name is properly theirs.
- P. 420 balderi is obviously a lapsus calami misspelling of balder so must be emended to balder. The 1837 publication merely has unnamed paintings of male ups and uns, and this text (my translation from German): Eumenis balderi. (Papilio nymphalis, Oreas nubila.) Fig. 981. 982. Male. Smaller than Eumenis aello*, it is the same as the excellent "l.c." figured male example and is on upperside, only through pale ground color and the dark other color, different from the greatly interrupted spot series. The underside of the forewings, except for the surroundings of the eyespots, is almost entirely pale-brown and going toward the hindwing, not like the cited species, is white exposed. Labrador. From Mr. Sommer. *Aello Huebner. Sammlung europaische Schmetterlinge Pap. 519. 520. Male. (end of text) This text has no types, and nothing to indicate that it is a description of a new species. It is just figures and description of a male Oeneis jutta from Labrador that the author identified as balder and then misspelled balderi, a lapsus calami as it was printed just a couple years after balder was named by Boisduval 1832 in Gue'rin-Me'neville. The mathematical odds that balderi is independent of balder is $(1/26)^{**6}$ (because there are 26 letters in alphabet and 6 letters in the root balder) which is essentially zero. It is not an available name. Neither balder nor balderi nor ssp. jutta applies to the Nearctic fauna. Why are taxonomists so eager to dredge up horrible worthless names and apply them to nicely stable critters?? That is a gross defect of taxonomists (which I have done a few times), and it must stop.
- P. 420 leussleri, District of Mackenzie no longer exists.
- P. 421 gibsoni, Holland 1931 p. 200 says pl. LXIII fig. 10 male is holotype being named in book, and Miller/Brown say HT in CM; however Pelham Catalogue says O.D. was in 1930 in Ann. Carn. Mus. 20:51, which evidently designated no holotype, which makes the fig. 10 male a Holland 1931 lectotype designation as Pelham states.
- P. 421 semplei, Holland 1931 designated lectotype pl. LXIII fig. 1 male.
- P. 421 simulans, holotype fig. Holland 1931 pl. LXXV fig. 19 male.

- P. 423 hanburyi, Mackenzie District no longer exists.
- P. 423 calais, Holland 1931 figured holotype on pl. LXII fig. 7 female; Miller/Brown wrote type may be lost.
- P. 423, all the ssp. placed in *O. chryxus* now belong in *O. calais*, except *O. chryxus* which is a separate species (see *Papilio* [N.S.] #12).
- P. 423 the true *chryxus* type is the male fig. in O.D. which is the valid holotype by monotypy; Shepard's lectotype is invalid and is a female of *O. calais altacordillera* (*Papilio* [N.S.] #12).
- P. 423, ivallda, Holland 1931 designated lectotype female on pl. LXII fig. 4.
- P. 424 Porter & Shapiro proved only that ivallda & stanislaus are conspecific.
- P. 424 oslari, **TL hereby defined as South Park between Kenosha Pass & Fairplay, Park Co. Colo.**
- P. 424 iduna, lectotype was designated by Holland 1931 pl. LXXII fig. 6 male; TL is required to be locality of lectotype which is "Mendocino" [County].
- P. 424 macounii, the valid lectotype was designated by Holland 1915 The Butterfly Guide, a male fig. pl. LXXVI, which may be the same specimen fig. by Holland 1931 pl. XXVII fig. 3, which Brown evidently treated as a syntype.
- P. 425 uhleri, **TL is Georgetown or eastward toward Idaho Springs**, as Georgetown may be too high altitude.
- P. 425 cairnesi, Holland 1931 figured uns of cairnesi type on LXXIV fig. 20m, & ups of same male on pl. LXXII fig. 6 male.
- P. 425 varuna, Holland's paralectotype is pl. LXII fig. 14 (not 11) female.

The 47-page checklist is rather useless as all the info is in the text, though one can better see the higher classification (families to tribes & genera) treatment here without the clutter of the species & ssp. names. P. 451 *Celaenorhinus* misspelled. P. 488 New Subtribe.

The bibliography of taxonomic references is long and is bound to be useful. Most of those references were not cited in the Catalogue's main text, and many of the letters after the year of the references that were cited do not correspond to the letters in the bibliography. But those references will be a great source for people researching the literature, especially if you know or suspect that someone worked on a bug but can't find the citation.

Index: P. 611 *Anaea/Anaedidi/Anaeni* are p. 392. P. 611 *anna* (*Diaethria*) is p. 431. P. 613 *asteria* is p. 431. P. 618 *colon* is p. 365. P. 621 *drusus* (*Chlosyne*) is p. 376. P. 643 *rainieri* (*Boloria*) is p. 306.

Overall, this Catalogue is very useful as a compendium of names. Pelham obviously spent years of effort on the task. (Based on Butt. Cascadia and Warren's Butt. Oregon, Pelham has a lot of biological information such as hostplants etc. that he has gathered, which has never been published. Maybe now he will have time to publish it.)

THE BUTTERFLIES OF NORTH AMERICA, A NATURAL HISTORY AND FIELD GUIDE.

James A. Scott. 1986. Stanford University Press, Stanford, Calif. 583 p., 64 color plates.

This is my own book, which took about 7 years to write. Following are additions and corrections. Of course some of the taxonomy is obsolete, as progress marches on, so I don't list most of those changes. A revision would update it all, slim down some parts, add 100+ more ssp., include photos of egg/larva/pupa of nearly all species, drop "field guide" from the title, and redo the plates in taxonomic sequence, using edited photos from nature like the Kaufman book. But I will require more remuneration for such a book than I received from this one, so I doubt that it will happen.

Literature citations for each bit of information from the literature were not included in the book because they would have required hundreds of pages of space and would have made the book impossible. However, the sources of hostplants and most other biological information are listed in my card file; this file consists of about 3000 cards, each full card containing ~1-8 references. The file should be published, but publication would require many months of keyboarding and great expense with no hope of financial return. Unfortunately the ineffective publisher failed to get the book stocked in the chain bookstores, so it did not sell very well, in contrast to low-quality little field guides that are stocked in chain bookstores and sell well.

Plates. In the film negative for each plate is a little image of the Kodak Gray Card (standard 18% reflection), a standard used by professionals to maintain print quality, but the incompetent printer/publisher failed to use that card to calibrate the colors (the lazy morons evidently just scanned the butterflies and ignored the gray card), so the colors on most plates are a little off (often too red). Plate 3, photo of "Feniseca" larva is not that & is mysterious, & looks only partly like actual Feniseca; D. Wright thinks it is a Syrphid fly larva that preys on aphids. Pl. 12, 35c is *C. occidentalis sacajawea*. Plate 14, 128a is *O. calais caryi*. Pl. 16, 109d is the new bug which is at least a new ssp. of *sthenela* from Humboldt Co. with more zigzag unh postmedian line etc. Pl. 18-19, *evarete* was changed to *genoveva*, then changed back again! Pl. 27, 230c-d are ssp. *camillus*. Pl. 36 *C. hesseli* is on *Vaccinium corymbosum* flowers. Pl. 38, 405c is *Pleb. atrapraetextus fridayi*. Pl. 41, 615a-b are *Codatractus uvydixa* from Nuevo Leon Mex. Pl. 43, 658a-b are *Erynnis baptisiae* from E Kansas. Pl. 43, 677c is ssp. *lena*. Pl. 50, 472e is ssp. *colorado*. Pl. 63 on 634a ups, add male to read 634a male ups. The 394b male letters should be moved leftward, prior to ups. 2002 butterflies are illustrated on the plates (10 photos of eggs, 45 of larvae, 30 of pupae, 1917 of adults). Note: All pinned adults used for photos now have a label that reads "fig. James Scott book". All pinned adults used for photos are in J. Scott collection except for 39 specimens in these collections: USNM (*Eurytides celadon* 1m1f, *Papilio ornythion* 1 f ups, *P. astyalus* 2f ups, *Phoebis orbis* 1m1f, *Diaethria asteria* 1m, *Anartia lytrea*

chrysopelea 1m1f, Epiphile adrasta 1f, Hamadryas atlantis 1m1f, Marpesia eleucea 1m1f, Libytheana carinenta motya 1m1f, Apodemia multiplaga 1m, Allosmaitia pion 1m, Strymon limenia 1m1f, Choranthus radians 1m1f, C. haitensis 1m, Euphyes berryi 1m1f, Erionota thrax 1m, Bolla brennus 1m1f, Polygonus savigny savigny=manueli 1m); Carnegie Museum (Eurema chamberlaini 1m2f, Chlorostymon maesites 1m1f); Dale Jenkins (Hamadryas iphthime 1m1f); J. Richard Heitzman (Libytheana carinenta motya 1f); Roy O. Kendall (Apodemia multiplaga 1m).

Part I. P. 1, schmetterling and tagfalter are names in modern German. In old dutch the word botervlieg does exist; boter=butter, vlieg=fly, but botervlieg is not used in modern dutch in which butterfly is now dagvlinder. P. 5 first line, change "wasps" to flies. p. 12. fig 4e should read first-stage larvae of some Lycaeninae. P. 13 4th line from bottom, should read fact fused, making ten visible in males, nine in females, and ten in larvae and pupae. P. 18 2nd parag. line 3, some metalmarks also zigzag erratically. P. 19 par. 4, wing muscles are also attached directly to the wing bases in dragonflies. P. 20 end of line 12, add hyphen on "dur" to read dur-. P. 22 some Lycaenidae also have a transverse dorsal cleavage line on pupal head. P. 27 3rd parag. line 2-3, delete and Asterocampa, 140-142. P. 28 line 2, add coloro after P. polyxenes. P. 29, in all or most species that overwinter as eggs, the first stage larva actually is fully developed within the hibernating egg (David M. Wright, pers. comm.). P. 37 last line change Petersen to Peterson. P. 40 and p. 419, Megathymus streckeri female noises are probably just due to normal wing movements so are nothing unusual. P. 45 Monarchs in S Mex. usually fly northward about 21 March. P. 59 3rd line from bottom should read lay large clusters of eggs... P. 60 2nd parag. line 2, most Colias also oviposit on leaf uppersides, and the Papilio glaucus group oviposits on upperside of leaf tips. 6th parag. line 2 change his Vanessa larvae to her Vanessa larvae. P. 61 Harkenclenus and Lethe eurydice also lay eggs in small clusters. P. 68 parag. 2, sap feeding adults generally land on the tree trunk above the sap then crawl down to it, to avoid being stuck to the sap. Par. 5, only the skippers with a long proboscis--long enough to extend to the end of the abdomen--eject a drop of fluid from the end of the abdomen onto bird dung or salty soil and then suck up the fluid (in other butterflies the proboscis is too short to reach to below the end of the abdomen). P. 73, 337. Heliconiini synthesize cyanogenic glycosides from the amino acids valine and isoleucine (A. Nahrstedt & R. Davis, 1983, Comp. Biochem. Phys. 75B:65-73), so these chemicals are not (or are rarely) obtained from their hostplants. Par. 3, Charles Remington (pers. comm.) states that Papilio machaon-group larvae are perfectly edible to birds. See p. 190 species 29 for recent demonstration of mimicry of millipedes. P. 74 par. 4, tropical spiders reject unpalatable butterflies such as Danainae and Heliconiini by odor rather than wing color (paper in Science magazine about 1987). Par. 6, the Hesperinae cremaster is typically attached to a silk pad within the leaf nest, so is not available for use as a spear. P. 76, change "perforated cupolas" to "ring glands" four places on page (the proper technical name for these structures is lenticle, the common name is ring gland, and other synonyms are porrenkupeln, ring-pores, etc.). P. 78 par. 2, the black form of P. machaon is recessive in England (see E. Ford, Butterflies) where there are no models, but it has become dominant in N. Amer. where it mimics Battus. P. 81 end of par. 2, add "and Strymon martialis & acis." (tails are longer in 7 Fla. hairstreaks too). The pale silvery blue color of arctic G. lygdamus kournakovi and C. argiolus lucia is also convergent. A. cethura pima and E. (hyantis) guaymasensis (Sonora) are both yellow in the southwest. P. 82 2nd to last line, change populations to animals. P. 86, now that pollution is lessening around London, the moths are becoming paler again. P. 87, crocodiles evolved just prior to dinosaurs, and birds are just one group (Therapoda) of dinosaurs (Robert T. Bakker, The Dinosaur Heresies, 1986, Wm Morrow, N.Y. 481 p.). Hedyloidea should be added to tree just above the 100 mark on trunk. P. 94 parag. 1 end, some Zygaenidae, larvae and adults, are remarkably resistant to cyanide. P. 95 parag. 4, if the moth is far from the bat it flies directly away from the bat. P. 96 3 lines from bottom, a Madagascar Spingid has a 25 cm proboscis; Charles Darwin predicted its existence even before it was discovered, based on the very long length of the flower it was later found to get nectar from and pollinate. P. 107 Colias hecla and meadii and Neophasia menapia and terlootii also show geographic replacement, but new evidence proves that Colias alexandra and occidentalis do not. P. 112 Trapezitinae occur in New Guinea also. P. 114 parag. 3, trumpets may have heralded the start of Linnaeus' field trips, rather than his return? P. 115 W. Edwards was president of a coal company.

Part II: P. 127 in fig. 49 legend, change "except for species [76] and 343." to "for some species". Libytheana actually lack Va so erase the word Va and the tiny circle next to it on the species 274 drawing of Fig. 49. P. 128-9, V1 is actually absent on prothorax of all first stage Lycaenidae larvae so erase it from bottom of left-hand rectangle of species 276 & 358 drawings. P. 130 parag. 5 line 4, delete "usually" because hardened rings are on all Lycaenidae larvae. P. 131 Hesperidae also have long setae on prothorax (paragraph 5). P. 132 Apaturinae have secondary setae on A10 (paragraph 28). P. 137 paragraph 28 add usually to read hairs usually very short; in paragraph 31 should read though they are seldom used. P. 139 the Lycaenid cremaster is often attached to a silk pad. P. 144 Erebia pupae are underground in Britain. P. 145 paragraph #55 should cite species 22-24 (not 21-24). P. 148 second to last line should be J. Eliot (not T. Eliot). P. 151 4th line from bottom, Acraeinae is on p. 228 not 338.

Part III: **Subspecies concept.** I would now recognize 100 or more subspecies than I did in this book. I hadn't seen some of the ssp. then. The goal was to include a ssp. in the book only if it is distinguishable from other ssp. by the ordinary person; thus numerous named ssp. listed by Miller and Brown (1981) that are distinguishable only by locality are not included. Some of the weak subspecies may be statistically significantly different from other subspecies, but may not be biologically significant; there is a big difference. With a sufficiently large sample size (such as 50 or 100 or more) almost

every population can be proven to be statistically significantly different from another population. Statistics was developed to detect slight differences in yield of vegetable crops, where a 1% greater yield may translate into thousands of dollars over the whole state or country; but this small difference is of doubtful interest to any Lepidopterist. Therefore, the only criterion for acceptance of a subspecies has to be some criterion of absolute difference (the most universal and least biased being distinguishability by an ordinary person). People complain about names changes because they cling to the names they first learned; they complained about my name changes, and now they're complaining about other people's changes. Changes are inevitable when they involve improvements or corrections, but I lumped too much in many groups (for instance *Euphydryas editha rubicunda* is very distinctive and should have been included even in the book's superlumped treatment of *editha* ssp.). So I now consider valid the old ssp. that were missed, and new distinctive ssp. **Form names.** The ICZN ruled that infrasubspecific names no longer have formal status after 1960. Miller & Brown (1981) interpreted this rule to mean that all form names should be ignored completely and listed as synonyms. However, it is hypocrisy to accept as valid numerous very weak subspecies which can hardly be distinguished even by an expert, and yet ignore forms which are far more distinct in appearance and which are involved in important biological phenomena such as seasonal adaptation (such as the spring form of *Colias eurytheme* whose color and size is vastly different), mimicry (such as the black female form of *Papilio glaucus*), or genetic polymorphism (such as the white females of *Coliadinae*). The ICZN ruling, that form names are no longer subject to the rules of nomenclature, is actually a great opportunity to improve the form names. Therefore, in the book a form which has a similar appearance and the same genetic/environmental cause in all the species in which it occurs is given the same name in all. This system is necessary because there are several dozen names for white females of *Coliadinae* species alone, for instance; all are given the name *alba*. Subsequent authors have mostly not used form names, but progress is slow and they will use them eventually. **Common names.** Changes of some common names are also inevitable when the former name was misleading. For instance, *Eurema nicippe* was called the Sleepy Orange, but anyone who has seen this species fly realizes that it flies fast and often quickly changes direction; thus the name is grossly misleading, hence the book name Rambling Orange, which isn't great (maybe Erratic Orange would be better). *Plebejus lupini* (now *alupini*) was called the Lupine Blue, but the bug has nothing to do with lupines so the common name was expunged. The name Lupine Blue obviously belongs to *Plebejus icarioides*, which eats dozens of lupine species and nothing else. I corrected all the misleading & bad common names. Science must be a search for truth, not a repetition of lies. The ignoring of my name fixes by NABA & others is very disappointing; what can we call a writer who deliberately uses a misleading lying name—a moron? The person who repeats an error merely to avoid change is committing a greater wrong than the person who created the error in the first place, because the repeater knew it was an error, whereas the creator thought it was correct. Justifying a fault doubles it.

P. 157, line 7, replace *sevata* by *josephina howarthi*. P. 163, *E. philolaus* ranges south to Costa Rica. P. 164, *xuthus* last seen on Guam in 1968 (*Envir. Ent.* 22:265). P. 167, the white line on the southern ssp. map (mentioned on 5th line from bottom) is too weak and should run from near Reno Nevada to SW Nebraska. Ssp. *oregonia* also occurs in N Elko Co. Nev. Peace R. Alta. populations have been named ssp. *pikei*, which often have rather stubby fw, & eyespots closer to Alaska. 4 lines from bottom, several black form *bairdii* are now known from SE Alta. P. 168, 5 lines from bottom, should read C and S Alta. P. 169, Umbelliferae hosts in Yukon-Alta. are for hybrid *P. machaon* X *zolicaon*. P. 171, *joanae* is a ssp. of *P. machaon*. *P. zolicaon* males probably become habituated to, and thus ignore, the obvious male pheromone; thus females can no doubt distinguish males from females, but males may not be able to distinguish females from males unless a female pheromone is present. W Mont.-NW Wyo. *polyxenes* records are probably misidentifications of *P. zolicaon* form *nitra*. Ssp. *asterius* is misspelled in book. P. 175 replace *nuttallii* by *graveolens*. P. 176. *P. andraemon* was introduced to Jamaica. P. 177. *P. deVries* (1986 *J. Res. Lepid.* 24:292) states that records of Piperaceae for *P. crespontes* belong to *P. thoas*, and that records of Rutaceae for *P. thoas* refer to *P. crespontes*; perhaps deVries is correct, though the photo of *P. crespontes* in his "Butterflies of Costa Rica" book looks like a misidentified *P. thoas* to me (Douglas Mullins states that it looks like *P. paeon thrason* to him). P. 179 *P. androgeus* was common in S Fla. from 1976 to 1983, absent since. P. 180, NW Neb. has mostly *P. glaucus glaucus* evidently. Female pupae from the cross *glaucus* X *canadensis* also stay in diapause. Ssp. *alexiares* also has black females. P. 181 parag. 3 11 ssp. *alexiares* also lacks the upper spot. P. 183 the *Litsea* host of *P. pilumnus* was reported as "laurel", which is probably *Litsea* but may be another Lauraceae plant locally called laurel. P. 184 *P. victorinus* is a ssp. of *P. menatius*. P. 188 replace *montanulus* by *maximus*. P. 190 *eversmanni* is not on Honshu. P. 192 *E. albania*, *mazai*, & *jethys* are all variants of *E. melite* according to Gerardo Lamas. Parag. 2 *Dismorphiinae* pupae also have cone on head and enlarged wings. P. 193 *C. boothii* & *thula* are ssp. of *Colias tyche*. P. 194, second *alba* should not be in boldface. *Lupinus arcticus* is the only recorded host of *Colias canadensis*; the other 5 "hecla" hosts belong to *C. hecla*. P. 195 see *Papilio* (N.S.) #12 for *alexandra/harfordii/occidentalis*. P. 198 in second-to-last sentence of *C. eurytheme*, nondominant traits do not sort out in the hybrids. *C. philodice* is now native to Bermuda (found in 1986 by David M. Wright). P. 199 line 4 change green to dark green. *C. interior* Wash. D.C. dot is placed wrong and should be in extreme W Md. Move the *C. interior* dot from Steens Mtn. in S Ore. to *C. pelidne*. P. 200 last line, ssp. *scudderii* is in Utah also. P. 201 *C. behrii*. The Munz & Keck flora uses *Vaccinium nivictum* for *V. "cespitosum"* in Calif. P. 202 the female form of *Anteos clorinde* resembling the male should be called form *alamacho*, as in *Phoebis philea*. P. 204 *P. sennae*, second to last line, mating should read coupling. P. 206 *Phoebis neocypris*, not *intermedia*. Some *E. nicippe*

hosts are shrubs. P. 209 the name *longicauda* is older than *gundlachia*, but form names no longer have formal recognition so the latter could be used because it is better known, however *longicauda* is more appropriate. E. *proterpia* strays N to Kans. P. 211 parag. 3, *Physaria* is almost never eaten by *Pierinae*. *Thlaspi arvense* is also introduced from Europe and American *Pierinae* have not yet adapted to it. P. 216 *Pieris rapae* is possibly resident in Iceland. P. 217 *P. napi* occurs on Pribilof Is. Alaska. P. 220-223, species 79-82 should be in genus *Pontia* (R. Robbins 1986 J. Lepid. Soc. 40:88). Last line p. 220, spots in protodice females are brown or black. P. 221 *P. protodice* patrols all day. P. 225, 369 *Pseudotsuga taxifolia* is a synonym of *menziesii*. P. 226 *Catacticta nimbe* is now known in Sonora Mex. P. 227 par. 4. *Polygonia*, *Asterocampa* (abd. segment 10), and *Ithomiinae* first stage larvae also have secondary setae. P. 228 line 10 delete "and perhaps *Aristolochiaceae*". *Danainae* has 157 species. P. 229 change first word "except" to even. In par. 4, the ancestral hostplant of *Danainae-Ithomiinae* was probably the plant group *Parsonsia*, which contains both cardiac glycosides and lycopsamine. P. 230 *Asclepias syriaca* was the main hostplant of overwintering Mex. Monarchs. Form *alba* is due to a recessive gene (J. Stimson & L. Meyers 1984, J. Res. Lepid. 23:153-160). L. Brower has found that there are only two flights Apr.-June in Fla. (where summer temperatures are lethal), 3 flights in Wis. P. 231 first line of *D. gilippus*, should read upf cells; these white spots are sometimes small, sometimes absent in cell CuA₂. P. 232 *D. gilippus* adults diapause in the Costa Rica mountains during the dry season. P. 236, Douglas Mullins has raised Aug.-Sep. *nabokovi* from May-June *pyracmon* thus proving conspecificity (and he also reared *C. nayarit* from *C. windi*!); larvae and pupae are straw-yellow like *pertepida*. *C. pertepida* has two flights southward. P. 238-9 *N. mitchellii* in N.C. is not really a distinct sp.; it is slightly darker beneath with the unh eyespots often oblique and longer than *Mich. mitchellii* (probably due to introgression with *N. areolata*), but the differences are not great enough for ssp. status. Ssp. *areolata* near the N.C. colony often have shorter than usual eyespots, as do S Dade Co. Fla. adults (the short-spotted *areolata* are now called *Neonympha helicta*). N.C. *mitchellii* has two flights M May-E June and L July-Aug. (ssp. *helicta* has two slightly later flights there). P. 239, *H. hermes* adults diapause Nov.-Jan. in N. Fla. (T. Emmel). Ssp. *benjamini* is a syn. or weak ssp. of *inornata*. Ssp. *mackenziei* also occurs NE Alta. P. 240 *C. haydeni* should read Larva (half grown) yellowish-green, with a supralateral and a subdorsal white stripe, and a white lateral stripe edged by purple-pink, with two purple-pink tails. *C. pegala* spp.: R. Gatrell (1985, So. Lepidopterists Bull. #2) sank *abbotti* to *pegala*, and sank *alope* to *pegala* because *alope* is in the blend zone of *pegala-carolina*. We should use: *pegala* (coast at least from Ala. to Fla. and N.C.) has large yellow patch with only one upf eyespot on males; *carolina* (mts. S.C.-N.C.) paler yellow patches & 1-2 ocelli; *maritima* (= *texana*) ("alope") (most of E U.S.) orangish-yellow fw patch with 1-2 ocelli. P. 242. ssp. *charon* occurs Ariz. also. P. 243 *E. rossii* from Que.-Lab. is based on "Cape Mugford Labrador". Ssp. *mckinleyensis* is the proper name rather than *erinny*. P. 246 *E. anyuica*=*oculta* from Yukon only differs slightly in long series from western adults, so is not a valid ssp. P. 249 *O. bore* is known from S B.C. NW of Jasper Park Alta. P. 251 *O. alpina* is also in Hudsonian Zone bogs. P. 252 *O. polixenes*, the S B.C. black spot on map represents a misidentified *O. bore* (Jon Shepard) so remove from map. *O. polixenes* is not yet known from Mont. (line 6 of species text). Add *O. philipi* (C. Alaska-Yukon, taiga bogs, flies in odd years) is usually dark-brown with 2 mm-wide white submedian & postmedian unh bands, whereas alpine pops. are usually paler; it has generally called a distinct species (?), Tom Kral states that normal paler *polixenes* and *philipi* fly together everywhere from bogs to tundra and intergrade completely, and he found the two forms in copula. P. 254 *A. "troglodyte"* *aidea* looks rather different from Caribbean-Fla. *troglodyta* spp., and could still belong to *troglodyta* (some female *aidea* have a hooked fw like the others), but it seems best to treat *A. troglodyta* and *A. aidea* as separate species. In the Caribbean, the gnathos is more hooked in ssp. *troglodyta* and *minor*, but the wing pattern is very similar, and N. Riley combined all Caribbean-Fla. ssp. into *troglodyta*. P. 255 change *alicia* to *reinthali* (peninsular Fla. NE to SW tip S.C.). P. 262 *L. weidemeyerii*: Rocky Mountain Admiral would be a better common name. Change *Salix subcoerulea* to *Salix drummondiana*. P. 263 the yellow-orange *L. bredowii* larva was a diseased abnormal individual (fig. by R. Pyle Audubon Field Guide to N.A. butterflies). P. 267-8, the cracking sound of adult *Hamadryas* is made with modified wing veins around the forewing discal cell. P. 275, *amathea* and *fatima* are ssp. (K. Brown Lepid. News 1988 #2 p. 39-40). P. 277-80 many places, keep the names *evarete* & *genoveva* the way they are. T. Turner & J. Parnell (1985, J. Res. Lepid. 24:142-153) switched the names, then Butt. Venezuela recently switched them back. Turner & Parnell give valid additional differences of valva spine number, testis color, larva and pupa color between the two. P. 280-281, the winter range of *Vanessa* is somewhat uncertain; thus *atalanta* may not overwinter so far north. *V. cardui* overwinters as adults, evidently on the Colo. plains rarely, where one was caught in Jan. in Colo. and migratory movement is rare in Sept.-Oct. P. 286, *P. gracilis* eggs laid singly, cremaster pad orangish-white. P. 288 *P. progne* cremaster pad pink; adult flight is slightly weaker than in *P. gracilis*; ssp. *progne* perches in gulch bottoms too. P. 289, 284, *N. milberti* and *P. comma* also roost with fw far forward & antennae between them. P. 290 *N. vau-album* larva is not green in Utah. P. 291 *N. antiopa* migrates out of the Central Valley Calif. in spring and migrates back in the fall (A. Shapiro). *hyperborea* is merely an aberration. P. 296 ssp. *rubicunda* is very much larger than *nubigena* so is a valid ssp. if this size is genetic. P. 300 *C. theona* hibernates as third-stage larva. P. 301 *C. melitaeoides* & *C. eumeda* are distinct sp. P. 302 3rd-4th lines from bottom should read If two recessive genes *r* are present. P. 304 *nycteis* hibernates as 4th stage; ssp. *reversa* is in SW Man not S Man. P. 308 replace *callina* by *arizonensis*. P. 309 *Phyciodes cocyta*=*morpheus* occurs in Sioux Co. Neb. My 3 papers in *Papilio*(N.S.) #7,10,13 totally revised *Phyciodes*. P. 314. *P. phaon* par. 4, change "native" to "frequent" in Ark.-Mo. P. 316 3rd line from bottom should read size as or larger than the female. P. 318 *B. eunomia* is

unsilvered in Labrador and Seward Pen. Alaska. P. 319 *B. selene* courtship should read: The male may hover above the female, and may flutter after landing behind her (sometimes he butts her while he flutters); then his wings are partly spread while hers are closed while they join. P. 319 ssp. *grandis* & *chariclea* fly in the same area of taiga bogs just NW Fairbanks Alaska, a sympatry that could be because they are separate species, but further collecting showed that *chariclea* is rare there most years and *chariclea* flies mostly earlier than *grandis* there, so this does not necessarily prove they are separate species (see Butt. Canada). 5 lines from bottom, 1st-stage-larvae also hibernate in Colo. P. 321 *B. improba* unnamed is ssp. *nunatak*. P. 322 Siberian *B. "alberta"* are evidently *B. erda kurenzovi*. *B. astarte* unnamed is only slightly if any different so is not a distinct ssp. P. 323 Canadian arctic Baffin I. etc. *tarquinius* missed, but looks like it is same sp. as *natazhati*. P. 327 Great Plains *S. idalia* colonies are still widespread and strong. P. 330 line 9 of *S. atlantis*, change "reddish" to redder-brown in two places. P. 331 ssp. *dorothea* is also in Chuska Mts. of NE Ariz.-NW New Mex. P. 334 *S. coronis coronis* & *semiramis* intergrade in Sequoia N.P. P. 340 *zorcaon* is a syn. of *H. isabella eva*. P. 342 4th line from bottom, add "occasionally", to read "Males occasionally locate female pupae", because in nature most matings involve flying females. P. 343 the elevation of abdomen and extrusion of the abdominal glands is probably a rejection posture even though the repellent chemical may not be available until after mating. P. 348 *Apodemia "mormo"* is 3-4 sp. (G. Pratt & G. Ballmer J. Lepid. Soc. 45:46-57; Scott *Papilio* [N.S.] #18). P. 350 *A. nais* males perch in gulches/depressions from about 12:30-2:30 p.m. The main habitat of *A. phyciodoides* is oak woodland (M. Smith 1986 J. Lepid. Soc.). P. 351. *A. multiplaga* ranges south to Costa Rica. P. 353 *C. wrighti* has three flights Mar., June-July, Sep.-Oct. P. 356 *Feniseca* hibernates as pupa. P. 359, replace *E. minijas* by *E. toxea*. P. 360 *titus* is in genus *Satyrium*. P. 362 parag. 6 last sentence should read The orange upf patch is small or absent in most areas (ssp. *strigosum*) including coastal S.C.-Ga. (unh marginal orange spots larger, ssp. *liparops*), very large in C Fla. (ssp. *floridensis*; marginal orange and blue unh spots larger). P. 363 form *heathi* is frequent in Carbon Co. Wyo. P. 364 *S. auretteum* occurs in Lake Co. Ore. P. 366 move the end of the *S. fuliginosa* black streak extending from NE Nev. and ending in C Nev., northward so that it ends in N end of Eureka Co. The C Nev. records of *M. leda* are all strays. *Thereus zebina* is properly named *Rekoa marius*. P. 366, 413, 414, 475. *Prosopis juliflora* is a synonym of *P. glandulosa* in most recent floras. P. 370 ssp. *margaretae* has much longer tails and a slightly-less contrasting unh median band. *Cyrrilla* is error. *C. polios* lays eggs on or near the base of leaf buds (rarely on flower pedicels). P. 371 change *augustus* to *augustinus* two places and in plates and index, delete (=augustinus). *C. mossii* is not in Neb. P. 373, *Sedum texana* is a Tex. host of *C. xami*. *C. hesseli angulata* occurs in Fla.-N.C. P. 374 lines 2-3, change form *smilacis* to form *arufa*. At end of sentence add Ssp. *smilacis* (coastal S.C.-Ga.) resembles *gryneus* but unh median line slightly straighter. Change "Cedros I. Mex." to "the similar ssp. *cedrosensis* on Cedros I. Mex has less white unh postbasal bars". Ssp. *gryneus* and *siva* intergrade in Dawson Co. Neb. also. 14 lines from bottom change one green-unh population to ssp. *juniperaria*. P. 378 ssp. unnamed is *pseudodumetorum*. P. 379 *C. s. paradoxa* was inadvertently named by Scott in this book (its appearance in *News of Lepid. Soc.* several years earlier did not satisfy the Code), see *Papilio* (New Series) #19. *E. heermanni* should read *heermannii*. P. 380-1, *Fixsenia* belong to *Satyrium*. P. 383 *S. melinus* abdomen is orange only on summer males. 4-5 lines from bottom, the desert scrub bug in Big Bend Tex. apparently is a separate species *Strymon serapio* (unh resembles *bebrycia*, unh has blue), Mar.-Sep. P. 386 two sp. of *Electrostrymon hugon*=*sangala*=*endymion* & *joya*=*canus* have been found in S Tex. P. 387 *L. phlaeas hypophlaeas*=*americana*. Near bottom, in Utah Uinta Mts. has near-*artemisia* & Wasatch Mts. has *artemisia*. P. 388 change *hyllus* to *thoe*. Warner Mts. has *L. nivalis warnermontana*. P. 389 *L. florus* (really a species) should be used rather than the older name *L. h. castro*, because I have appealed to ICZN to suppress *castro* (see *Papilio* [N.S.] #12, 18). P. 390 ssp. *epixanthe* occurs in N.Y., Mass, N.J. southward. P. 391 change ssp. *nevadensis* to *obscuramaculata*, and change index *nevadensis* (*Lycaena*, sp. 382) to *obscuramaculata* also. P. 392 *L. x. editha* from Alta. is based on only one old dubious record. P. 394 *L. marina* overwinters in S Nev. P. 395 *B. exilis* arrived in Hawaii 1978. P. 397 ssp. *gozora* doubtfully occurs in U.S., or may not be distinct from ssp. *echo*. "Argiolus" is now 6 sp. P. 401-3. *E. battoides*: *Euphilotes bernardino* is a separate species and includes ssp. *allyni* and *martini*. *Euphilotes* are totally changed & plunged into chaos since the book. P. 405 *E. rita*: no Mex. records but probable. P. 406 *idas* is now 3-4 sp. (see *Papilio* [N.S.] #18). P. 408 *P. melissa* males often perch on low spots of trails or gulches to await females, but often do not stay long (*P. glandon* occasionally does this too). *Astragalus mollissimus* is a synonym of *A. drummondi*; *Psoralea* is not a host. P. 409 see *Papilio*(N.S.) #12 for ssp. of *saepiolus* & *icarioides*. P. 411-2 ssp. *acmon* is a species, the other "acmon" ssp. are *P. alupini* ssp., & *P. chlorina* & *P. c. monticola* is a separate sp. P. 414 2nd line, should read Greater Antilles. P. 415 *Pyrrhopyginae* also spread wings at rest. P. 419 the clicking sounds are just incidental products of flight. P. 420 *Aegialini* last sentence, *Aegiale* larvae also are in tequila. P. 425-6 *Carterocephalus* and *Piruna* are in subfamily *Heteropterinae*. P. 425 *C. palaemon* mature larvae hibernate (see M. Brooks & C. Knight, *A Complete Guide to British Butterflies*, Jonathan Cape, London, which has larva & pupa photos). P. 427 6th line from bottom, should read south to Fla. keys. P. 432 *Stipa columbiana* is error. P. 433 *A. prittwitzi* also flies in E Aug. (probably 3 generations or more); females vary from male-like to mostly brown on ups. *Hylephila phyleus* is a stray in Bermuda. P. 435 *Stinga* has 1 gen., Sep. is error. P. 436 *H. juba* eggs stay white (become only very faintly pinkish), larvae hibernate about 1/3 grown. *H. comma* is mislabeled in W Tex.; ssp. unnamed is *mattonorum* & has greenish-ochre-brown unh that often lacks spots. Replace *Andropogon saccharoides* by *A. scoparius*. P. 438 *H. leonardus* males also often perch among clusters of flowers to await females. P. 441 *H. dacotae* males at low density perch mostly on ridgetops or on top of slopes. Only some *dacotae* sites are

on shorelines of Ice Age lakes. P. 442 *H. nevada*: C Nev. and extreme S Nev. records are errors (misidentifications). P. 443 3rd line, change *Festuca ovina* to *Festuca idahoensis*. The W Grand Mesa *P. draco* intermediates are actually *P. sabuleti*. 7 lines from bottom change *Eragrostis trichodes* to *Sporobolus airoides*. P. 445 *P. mystic* males perch in grassy swales and gulches. *P. themistocles* eggs develop reddish dots. P. 446 replace *scoparius* by *gerardii*. P. 447 *W. egeremet* males perch all day on plants 1-4 m above ground in little forest clearings. P. 449 *Problema byssus* ssp: upf base blackish-yellow in SE U.S. (ssp. *byssus*), yellow on Great Plains (ssp. *kumskaka*). P. 450 *P. bulenta* has only one flight in Md., June 20-July 14. *O. yuma* males also patrol slowly among hostplants to seek females. P. 451 replace *Blepharoneuron tricho.* by *Muhlenbergia montana*. P. 452 *taxiles* is distinct sp.; replace *Puccinellia airoides* by *Glyceria striata*. P. 455 *Euphyes dion* ssp. *bayensis* (Bay St. Louis S Miss. to SE Tex., oranger ups, stigma slightly narrower) was named as a distinct species but intermediates to ssp. *dion* occur at TL. P. 458 *A. hianna deva* (stigma absent to present & 3-parted) has only one flight, May-June. P. 459 *A. python margarita* is a syn. of ssp. *python*. *A. ovinia* perches in afternoon as well as morning (K. Roever). P. 460 *A. aenus megamacula* occurs only in S Ariz. (ssp. *aenus* in C-N Ariz.). P. 462 *A. unname* is *A. elissa*, which occurs near Patagonia southward to Atascosa Mts., and in Cochise Co. Ariz. (coll. Biedeman 1903, Ent. News. 15:344) but is not known specifically from the Mule Mts. It is a temporary resident in S Ariz. (absent because of freezes from 1979 to about 1983), and does not fly in May. *Celia & belli* are distinct sp. P. 463 *A. tolteca prenda* is apparently only a temporary resident in S Ariz., moving north from Sonora Mex in late summer of some years. P. 464 *A. vialis* has 2 gen. in Neb. *A. phylace* flies at low to middle altitudes in the Chiricahua Mts. Ariz. (*A. fimbriata* at higher altitudes there), and (p. 465) sometimes flies with *A. fimbriata*. P. 465 parag. 4 capitalize Subtropical. P. 466 *C. ethlius* is a stray in Bermuda. P. 478 *A. pseudocellus* last found in Ramsey Can S Ariz. June 28 1936. P. 480 Ssp. *mexicana* has brown unh. P. 481 *Codatractus valeriana*= *mysie* (Ariz.-W. Mex.) has large white fw spots; the plate 41 photos are *Codatractus uvydixa* from E Mex. with small or absent fw spots. P. 485 *S. pulverulenta* now known in Baja Calif., Mex. P. 491 *E. horatius* unh submarginal spots usually paler than *E. p. meridianus*. P. 492 *E. persius* occurs in N Man. also. P. 493 Paul Klassen lists *E. lucilius* records from W-C Man. to Churchill Man., which I treated as misidentified *persius*; these have been reexamined; there are evidently valid records of *E. lucilius* from Gillam, Thompson, Mile 349 Churchill RR, Sandilands Man., but not from W-C Man. *E. persius* Salicaceae hosts were no doubt errors based on *E. icelus*. P. 494 *E. baptisiae* flies with *lucilius* also to Tenn. P. 495 *P. albescens* (now considered a species?) also occurs in S Nev., but is probably dubious in Kans. P. 499 *P. mejicanus* pupa is like that of *catullus* but hairs slightly longer (delete "slightly hairier"). P. 501 *P. araxes* flies in L July also; adults rest and bask with wings spread laterally; adults sometimes land on open water (!) with wings spread to sip water.

Appendixes: P. 505 for Bermuda, add *Vanessa virginiensis** & *Colias eurytheme**, delete asterisk after *Colias philodice* & *Calpodus*, delete *Lethe*. Hawaii, add *Polites sabuleti*. Add Pribilof Is. Alaska *Pieris napi* (N). P. 506 4th line from bottom, add "or screw clamps" after copper tube. P. 509 a #0 insect pin should be deheaded and backed into match stick. P. 510, to make a holder for feeding adults, cut 8-10 hacksaw blade slits into the edge of 1/4" thick X 1" X 4" plywood, insert a spring-steel forceps holding adult into each slit (this works great!). The plywood can be the edge of a nice feeding trough that holds the honey-water food. P. 511, to mate adults in the lab, a female can be held with forceps and backed toward a caged male; or the female can be anesthetized and placed near the male. Immatures can be overwintered in the refrigerator if sphagnum moss is placed in closed container, or little drops of water are added occasionally to prevent dessication. P. 512 replace "Dupont Duco Cement" with "acetone-based clear cement."

Reference Matter: P. 529, Cotton "Grass" is really a sedge; its name should have been Cotton Sedge. P. 532 in Graminae, change "passim" to "most of". P. 546 in "life zone" entry delete "topography". P. 561 add to index: *elissa* (*Amblyscirtes*, sp. 545), 462, pl. 46). P. 566 add to index: instar, 21. P. 578 add to index: *simplonia* (*Euchloe*), 214. P. 581 *viardi* is in genus *Itaballia* not *Itabella*.

THE BUTTERFLIES OF NORTH AMERICA. William H. Howe (ed. & painter) & 20 contributors. 1975. Doubleday & Co. Inc., Garden City, New York. 633 p.

This book was a vast improvement on Holland's 1930 Butterfly Book, which except for the nice plates contained little information. It is now out of print and is somewhat out-of-date, so I just correct the plates. There wasn't much biology in this book, so I later decided to write my own book.

The ~2000 paintings themselves—by William Howe—are some of the best butterfly paintings ever done. Each plate was painted on a single cardboard, so the individual butterflies could not be rearranged, accounting for the jumbled positioning of some species as individual butterflies were added to later plates as afterthoughts. Following are corrected identifications (some of the minor recent name-changes and lumping of weak ssp. are not updated). Pl. 1#20 is *Poladryas minuta* near-nympha. Pl. 2#12 may be *septentrionalis* but looks more like *areolata* from SE U.S.; #24 is *N. ridingsii stretchii*. Pl. 3#1-2&16 are *C. pegala maritima*; 3&5&8-9 *C. p. nephele*; 15 is *C. p. gabbii*. Pl. 4#3-4 *C. p. nephele*; 7&11 *C. oetus oetus*; the *Coenonympha* are all *C. tullia*; 14-15 *C. t. ampelos*; 18-19 *C. t. mono*; 22 *C. t. pseudobrenda*; 28 *E. mancinus*. Pl. 5#3&6 *C. p. nephele*; 16 *E. stubbendorffii ethela* (N.A. has *E. stubbendorffii* not *E. theano*); 18-19 *C. t. subfusca*. Pl. 6#5 *O. nevadensis nevadensis*. Pl. 7#11 looks like it may be *O. philipi*; 15-16 *O. polixenes luteus*. Pl. 8#14 *O. calais strigulosa*; 16 *O. calais caryi*. Pl. 9#5-6 *O. polixenes subhyalina*; 12 *O. bore fordi*. Pl. 10#1-2 *C. pegala pegala*; 8

E. mancinus; the *Coenonympha C. tullia*; 16-18 *C. t. inornata*. Pl. 11#3 *A. troglodyta floridensis*; 8 *A. clyton flora*; 9&23-24 *A. celtis reinthali*; 21-22 *A. celtis jeffermont*. Pl. 12#3 *L. weid. weidemeyerii*; 9-10 *A. clyton texana*. Pl. 13#2 *L. weid. angustifascia*; 3-4 *L. portlandia portlandia*; 6-7 *A. celtis montis*; 8-9 *A. clyton texana*; 10 *L. anthedon*; 13 *L. anthedon borealis*. Pl. 14#8 *L. lorquini itelkae*; 10-11 *Adelpha eulalia*. Pl. 15#1 *M. dorcas amymone*; 2 *M. zerynthia=coresia*; 5 *Cyclogramma bacchis*; 10 *P. minuta arachne*; 11 *D. serina=egaea dyonis*. Pl. 16#5&7 *P. marg. marginalis*; 6 *P. faunus rusticus*; 16-17 *P. oleracea frigida*. Pl. 17#11 *P. gracilis zephyrus*; 15 *P. faunus rusticus*. Pl. 18#3 *P. faunus cenveray*; 5 *P. oreas near-silenus=threatfuli*; 6 *P. oreas oreas*; 11 *Junonia genoveva*; 14 *J. evarete nigrosuffusa*; 15 *A. amathea fatima*. Pl. 20#3-4 *S. aphrodite near-whitehousei*; 12-13 *S. aphrodite manitoba*. Pl. 21#6-8&15 *S. zerene platina*. Pl. 22#3-4 *S. zerene gunderi*. Pl. 23#6-8 *S. coronis snyderi*; 11-12 *S. coronis halcyoneXsnyderi*; 15 *S. coronis semiramis=hennei*. Pl. 24#6-7&9-10 *S. hesperis dennis*. Pl. 25#11-12 *S. zerene picta*; 14-15 Castle Lake. Pl. 26#3 *S. hesperis dorothea*; 4-5 *S. hesperis chitone*; 6-7 *S. atlantis sorocko*; 9-10 *S. hesperis tetonia*; 11-12 *S. hesp. hesperis*; 13-14 *S. hesperis electa=nikias*; 15 *S. hesperis cottlei=dodgei*. Pl. 27#1-2 *S. hesperis greyi*; 3-4 *S. hesperis tetonia=wasatchia*; 5-6 *S. hesperis viola*; 7-8 *S. hesperis beani*; 9 *S. hesperis cottlei=dodgei*; 10 *S. hesperis irene*; 11-12 *S. hesperis dennis*; 15 *C. sterope vallismortis*. Pl. 28#1-2 *S. call. callippe=comstocki*; 3-4 *S. call. calgariana*; 5 *S. call. semivirida*; 9-10 *S. call. nevadensis*; 11-12 *S. call. macaria*; 13-14 *S. call. rupestris=inornata*; 15 *S. call. semivirida*. Pl. 29#1-2 *S. call. calgariana*; 3-4 *S. call. juba=laura*; 9-10 *S. call. meadii*; 11 *S. call. juba=sierra*; 12-13 *S. call. calgariana*. Pl. 30#3-4&14-15 *S. hydasphe rhodope*; 6 *S. hyd. rhodope* (from Wyo. not NM); 7-8&13 *S. hydasphe minor*; 11-12 *S. hyd. hydasphe*. Pl. 31#6-9 *S. hesperis beaniXtetonia*; 10 *S. hesperis schellbachi*; 11-12 *S. hesperis lurana*; 13 *S. atlantis pahasapa*; 14 *S. egleis utahensis*; 15 *S. hydasphe rhodope* (from Wyo., not NM). Pl. 32#1-2 *S. aphrodite whitehousei*; 3-4 *S. hesperis nausicaa*; 11-12 *S. morm. mormonia*; 15&17 *S. mormonia eurynome*; 16 *S. aphrodite manitoba*. Pl. 33#1-3&15 replace *napaea* by *alaskensis*; 8 *B. improba improba aberration youngi*; 13-14 *B. titania butleri*; 16 *B. titania arctica*. Pl. 34#1 *B. eunomia caelestis*; 3 *B. eunomia tricularis*; 7-8 *B. freija freija*; 9 *B. selene myrina*. Pl. 35#2 surely Moraine Park; 5-6 *B. bellona bellona*; 15 *B. titania grandis*. Pl. 36#4&7 *E. chalcedona hennei*. Pl. 37#9-10 *E. editha quino=augusta*; 13-14 *E. anicia alena*; 15-16 *E. anicia maria* (*C. Ferris* in *Butt. Rocky Mtn. States* p. 332 says the *maria* & *windi* IDs were switched, but he was wrong as the *windi* looks correct to me & *Todd Stout* finds that *maria* is blackish); 17 *E. anicia eurytion=windi*. Pl. 38#1-2 *E. anicia wecoeut*; 4-5&9&12 *E. editha editha*; 7-8 *E. editha quino*; 10-11 *E. editha rubicunda*; 16 *E. bernadetta macyi*. Pl. 39#1-2 *E. anicia eurytion=windi*; 5-6 *E. bernadetta rorina*; 7-8 *E. anicia anicia*; 15&18 *E. bernadetta veazieae*. Pl. 40#1 *C. gorgone*; 9 *C. sterope dorothyi*; 10-11 *P. minuta arachne*; 12 *Chl. acastus dorothyi* male *Durkee, Ore. May 16 1940 LACM*; 13 *Chl. sterope sterope* male *John Day River, Gilliam Co. Ore. June 2 1961 DLB*; 14 *Chl. sterope vallismortis=robusta*; 15&18 *Chl. whitneyi whitneyi*; 16 *Chl. sterope sterope*. Pl. 41#1-4 *C. sterope acastus*; 6 *C. palla dark ssp.*; 7 *C. palla palla=eremita*; 9-10 *C. whitneyi damoetas*; 15 *C. leanira fulvia*; 16 *C. leanira alma*; 17-18 *C. leanira cerrita*; 27-28 *Microtia (Texola) elada perse*; 29 *Microtia (Dymasias) dymas chara*. Pl. 42#6-7 evidently are correct *E. anicia maria=effi*; 8 *E. colon wallacensis=nevadensis*; 11-12 *S. mormonia eurynome*; 13-14 *S. morm. opis=jesmondensis*. Pl. 43#3 *S. cybele leto*; 12 *Chl. palla palla=eremita*; 13-14 *P. minuta arachne*. Pl. 44#7 *P. phaon jalapeno*; 9 *P. batesii batesii*; 10 *P. pulchella owimba*; 12 *P. pulchella near-owimba*; 14 *P. texana texana* (redder); 18 *P. frisia tulcis*. Pl. 45#3 *S. nokomis apacheanaXnokomis*; 4 *S. n. near-nokomis*; 5-6 *C. sterope sabina*; 11 *E. colon wallacensis=nevadensis*; 12-13 *C. harrisii harrisii*; 14 *P. graphica*; 17-18 *C. hoffmanni segregata*. Pl. 46#3 *S. hesperis dorothea*; 9-10 *E. editha edithana*; 13 *P. minuta minuta*; 14 *E. bernadetta macyi*; 17 *E. editha nabigena*. Pl. 47#1-2 *Libythea carinenta bachmanii*; 7&24-25 *A. nais chisosensis*; 14-15 *L. carinenta larvata*; 29-30 *Cal. rawsoni rawsoni*; 41-42 *A. palmeri marginalis*; 45-46 *A. virgulti deserti*; 47-48 *A. virgulti mejicanus*. Pl. 48#2&17-18 *A. dialeuca dialeuca*; 3 *A. virgulti mejicanus*; 4&6-7 *A. mormo mormo*; 5&8&13 *A. virgulti virgulti*; 9&14 *A. virgulti virgulti blacker Baja variety*; 15-16 *A. cythera tuolumnensis*; 19-20 *A. cythera cythera*; 28-30 *Hypostrymon critola*; 34-35 *Satyrium acadica*. Pl. 49#2 *Eumaeus toxeus*; 4-5 *A. halesus halesus*; 12 *Satyrium semiluna semiluna*; 16-17 *C. gryneus nelsoni*; 18-19 *C. gryneus siva*; 20-21 *C. xami texami*; 22-23 *C. gryneus loki*. Pl. 50#1-2 *C. aug. augustinus*; 19-20 *C. niphon niphon*; 22 *E. laeta quaderna*; 23-24 *Satyrium titus titus*. Pl. 51#2 *C. gryneus swadneri*; 10 *Rekoa marius*; 13-14 *C. cecrops isobeaon*; 15-16 *S. istapa*; 17-18 *Udara blackburni*; 23-24 *Ministrymon azia*. Pl. 52#3-4 *S. melinus melinus*; 5-6 *S. bebrycia*; 23-24 *S. Satyrium titus watsoni=winteri=campus*. Pl. 53#3-5 replace *fotis* by *mossii*; 6 *C. gryneus muiri*; 11 *Satyrium favonius ilavia*; 12 *Electrostrymon hugon=sangala*; 13-14 *C. affinis perplexa*; 15 *C. sheridani [viridis] comstocki*; 16 *C. sheridani [viridis] lemberti*; 17-18 *C. affinis homoperplexa*; 21-22 *C. [sheridani] viridis viridis* (the name *dumetorum* is *toxotaxon sewage*). Pl. 54#1-2 *Satyrium fav. favonius*; 3-4 *S. favonius ontario*; 17 *S. sylvinus putnami*; 18 *S. sylvinus megapallidum*; 26 *L. cupreus artemisia*. Pl. 55#1-2 *L. dione*; 5-6 *L. phlaeas hypophlaeas*; 7-8 *L. thoe*; 11-12 *L. florus dospassosi*; 18 *L. heteronea heteronea* (or *klotsi*); 19-20 *L. nivalis browni*; 26 *L. rubidus ferrisi*. Pl. 56#5-6 *L. arota virginensis*; 30-31 *P. icarioides pembina*. Pl. 57#4 *Phil. sonorensis extinctis*; 7-8 *Euphilotes spaldingi pinjuna*; 11-12 *Cel. lucia lucia*; 13-14 *Cel. ladon*; 15 *Cel. neglecta echo*; 16 *Cel. lucia sidara*; 19-20 *Cel. neglecta*; 25-26 *Hem. thomasi bethunebakery*; 27-28 *H. isola*. Pl. 58#3-4 *P. chlorina chlorina*; 5-6 *P. chlorina monticola*; 9-10 *G. lygdamus incognitus*; 12 *G. lygdamus oro*; 13-15 *G. lygdamus xerces*; 19-20 *P. podarce*; 25-26 *Euphilotes bernardino*; 27-28 *E. glaucon centralis*; 29 *E. ancilla*; 34-35 *B. exilis pseudofea*. Pl. 59#1-2 *P. icarioides incognitus* (closer)X~~pheres~~; 7-8 *P. saepiolus amica*; 10-11 *P. aquilo labrador*; 15 *P. shasta pitkinensis*; 16 *P. shasta charlestonensis*; 24&29 *P. scudderi empetri*; 27-28 *P. melissa melissa*; 30-31 *P. atrapraetextus near-longinus*; 32&37 *P. scudderi aster*; 33-34 *P. atrapraetextus longinus*; 35-36 *P. atrapraetextus alaskensis*;

38-39 *P. atra. atrapraetextus*. Pl. 60#1-4 *P. anna anna*; 5-6 *P. anna lotis*; 7-8 *P. atrapraetextus sublivens*; 9-10 *P. (atrapraetextus?) samuelis*; 13 *P. saepiolus rufescens*; 14 *P. saepiolus aehaja*; 15-16 *P. saepiolus near-gertschi*; 17-18 *P. icarioides fulla*; 19-20 *P. icarioides pembina*; 29-32 *P. acmon*. Pl. 61#3-4 *E. ancilla columbiae*; 15-16 *E. glaucon intermedia*; 17-18 *E. glaucon oregonensis*; 19-20 *E. glaucon glaucon*; 21-22 *E. bernardino martini*. Pl. 62#1 *Pap. troilus* male; 3 *P. glaucus rutulus*; 6 *P. glaucus glaucus*. Pl. 63#3 *P. zelicaon f. nitra*; 9-10 *P. machaon bairdii*; 11 *P. polyxenes coloro*; 12 *P. machaon oregonia*. Pl. 64#1-2 *P. polyxenes coloro f. clarki*; 3 *P. machaon brucei*; 4 *P. p. coloro f. comstocki*; 6 *P. polyxenes kahli*; 7 *P. zelicaon*. Pl. 65#2 abdomen tip should show space; 5 *P. astyalus pallas*. Pl. 67#8 *P. machaon brucei*. Pl. 68#1-2 *P. phoebus smintheus*; 3-4 *P. phoebus hermodur*; 5-6 *P. phoebus apricatus*; 7 *P. phoebus pseudorotgeri*; 13-16 *P. clodius menetriesi*. Pl. 69#5 *P. phoebus apricatus*; 6-7&9-10&13-14 *P. phoebus smintheus*. Pl. 70#19 *P. oleracea*; 20 *P. hulda pseudobryoniae*; 24 *P. marginalis venosa*. Pl. 71#4-5 *K. lyside*; 14 *A. julia flora*; 16 *A. julia columbia*; 22 *E. hyantis lotta*; 24-25 *E. olympia*. Pl. 72#2 *E. दौरा palmira*; 5 *E. arbela boisduvaliana*; 6 *E. salome*; 21 *E. lisa*. Pl. 73#3&6 *C. scudderii gigantea*; 7 *C. occidentalis sacajawea*; 8 *C. tyche thula*; 12 male; 13 *C. p. minisni?* Pl. 74#6&9 *C. (tyche?) canadensis*; 7-8 *C. tyche boothii*; 14 *C. occid. chrysomelas*; 17 *C. occidentalis near occid. (Xwasatchia)*. Pl. 75#9-10 *C. philodice*; 19 female; 25 male. Pl. 76#13 *A. julia flora=alaskensis*; 14 *A. julia browningi*; 18 *E. hyantis andrewsi*; 21-22 *P. sisymbrii sisymbrii*. Pl. 77#5-6 *A. virgulti duryi*; 9-10 *P. batesii anasazi*; 15&17-18 *O. polixenes subhyalina*; 16 *C. sthenele paulus f. damei*. Pl. 78#1-2 *C. tullia insulanus*; 3-4 *C. tullia californica*; 5 *P. pulchella camillus*; 11 *P. hulda pseudobryoniae*; 12 *C. tullia kodiak*; 13-14 *C. tullia mackenziei*; 17 *P. clodius baldur*; 19 *C. occidentalis wasatchia*; 24 *Allosmaitia strophius*. Pl. 80 nearly all the ssp. are synonyms. Pl. 81#1 *A. remingtoni valverdiensis*; 2 *A. mariae mariae*; 4&7 *A. aryxna freemani*; 9-10 *A. n. neumogeni*; 11-12 *A. aryxna baueri*. Pl. 82#10 *M. cofaqui*; 12&15 *A. n. neumogeni*. Pl. 83#1-2 *A. neum. neumogeni*; #3&6 *M. yuccae coloradensis*; 17 *P. panoquinoides errans*. Pl. 84#2 *Lerodea arabus dysaules*; 5-6 *Notamblyscirtes simius*; 12-13 *A. aenus erna*; 14 *A. aenus linda*; 19 *A. tolteca prenda*. Pl. 85#3-4 *A. hianna deva*; 5-6 *A. hianna loammi*; 16 *A. python*; 23-24 *E. vestris vestris*. Pl. 86#5-6 *pilatka*; 9 *Quasimellana eulogius*; 13-14 *Poanes melane melane*. Pl. 87#3-4 *Paratrytone snowi*; 18-19 *Anatrytone logan*. Pl. 88#1-2 *H. comma harpalus=yosemite*; 6 *H. comma colorado*; 15-16 *H. comma near idaho=mojavensis*; 17 *Polites rhesus*; 18 *Polites carus*; 19-20 *H. comma idaho*. Pl. 89#30-31 *Polites carus*; 32-33 *Polites rhesus*. Pl. 90#1-2 *P. peckius surllano*; 31-32 *H. comma idaho*; 38-39 *H. leonardus pawnee*. Pl. 91#16 not from Colo. (probably Ariz.); 30-31 *N. neamathla julia*; 36 *Monca crispinus=tyrtaeus*. Pl. 92#3 *mejicanus*; 19 *P. scriptura*; 25 *Codatractus valeriana*; 32 *Piruna penaea*; 33-34 *P. albescens*. Pl. 93#32 *E. propertius?*; 41-42 *E. propertius meridianus*. Pl. 94#11-12 *T. mexicana*; 13 looks like *T. mexicana*. Pl. 95#13 *A. hianna hianna*; 15 *A. alector hopfferi=gilberti*. Pl. 97#1-2 *Hesperopsis libya libya*; 3 *Okoboji*; 5 *Hesperopsis alpheus oricus*; 6 *P. scudderii scudderii*; 10-11 *Hesperopsis alpheus graciela*; 18 *L. helloides (Aug. 28 is 2nd gen.)*; 19 *L. phlaeas alpestris*; 20-21 *C. scudderii harroweri*; 22 *L. dorcas dorcas* (odd with so much orange, but July 24 is *dorcas flight*; *dorcas & helloides* both occur at Whiteshell Park); 23 *C. occidentalis krauthii*; 25 yellow variant; 26 *P. savigny savigny=manueli*; 27 *P. occidentalis nelsoni*.

SYSTEMATICS OF WESTERN NORTH AMERICAN BUTTERFLIES.

Thomas C. Emmel, ed. Dec. 18, 1998. Mariposa Press, Gainesville, Florida. 878 p.

This book contains a lot of information concerning western U.S. butterflies, especially the descriptions of numerous new subspecies. Many authors wrote the 73 papers (chapters) in the book, especially John F. Emmel, Thomas C. Emmel, Sterling O. Mattoon, and George T. Austin. The papers vary in quality and content from substantial (esp. discovery of the *cassiope & ricei=calneva blues* and *sthenele=hypoleuca satyrs*), to unsubstantial. This review mainly is limited to corrections, and to my opinion of the status of the new subspecies. The ssp. named in this vol. that I think are synonyms are noted below; the other named ssp. not synonymized below I think are valid.

Since this book is mostly concerned with naming new subspecies, a discussion of the ssp. concept is necessary. A valid ssp. must have differences that can be seen by the ordinary person in my opinion, at least after the location of those differences are pointed out. This definition is necessary because butterflies have always been and will always be studied mostly by amateurs (the little flush of money available for university researchers toward the end of the last century will not return, as future money will go to gene jockeys), so a technical definition of ssp. will not work. (Some famous geneticists even deny the existence of ssp., for example among humans, because of so much variation of most traits in the species, but of course using this argument one could deny even the existence of species, as chimpanzees and humans are both variable in most traits and have the same variation in ABO genes and baldness etc., yet are still distinct species even while sharing ~98% of their genes. And according to this argument Neanderthals and ancient humans were not even subspecies, even though most scientists now believe Neanderthals were a separate species.) A really weak ssp. may be different from another one year, but the next year it may look a little different due to random sampling of genes for the next generation and different weather that might affect the phenotype a bit; or maybe the weak ssp. was named because the sample was small and wasn't typical over many years (*Hesperia pahaska martini* and several *Papilio indra* ssp. are examples).

A valid ssp. must also have distinctive trait(s) of its own, and cannot be an intermediate between other named ssp. with no distinctive feature of its own, in my opinion; thus a transition area or cline should have only two names (one for each end), and all names for mere intermediate populations are automatically invalid; without this rule, the transition area could end up with three, four, or a dozen silly names based on the % of one or two characters.

The subspecies concept is valid, because evolution is a forking process in which one DNA line forks into several, so if we measure the distance between tines of the fork we get a continuous gradation of distances depending on how close we are to the forking point and how rapidly they have diverged, so a species concept that has the most categories (species, subspecies, infrasubspecies, microsubspecies, etc.) best matches the different degrees of difference found in nature. We only recognize subspecies and not infrasubspecies because the ICZN tells us to without any evolutionary justification, just because of a history of abuse of naming aberrations etc., and because it simplifies the bookkeeping problem (the ICZN Code is a set of laws and latinization rules, obsessed with latin grammar, that has nothing to do with biology and applies only to names). The fact that some “subspecies” are later raised to be species, and some “species” are later demoted to ssp., proves that the ssp. concept is valid for uncertain taxa. Here’s a suggestion, used in my paper with N. Kondla & S. Spomer on *Speyeria atlantis/hesperis*: different degrees of difference can be shown in a checklist by indenting subspecies to the right: the farther they are indented, the less difference they show. Or, more simply, > can be used for one level of indentation, >> for two levels, etc.

My judgement of some of the ssp. may be wrong, because in many cases I have seen few or no actual specimens of them, and it would be necessary to see large series to determine their validity. Some of them may be weak ssp. rather than synonyms. However, the more distinctive a ssp. is, the fewer specimens you must examine to see that it is distinct, so I don’t think I have missed too many this way, given my lumpier definition of ssp. And frankly I—and I think most lepidopterists—don’t want to recognize ssp. that require a huge series to notice the difference. Obviously all the ssp. were named in this book because the authors thought they were valid, so my saying that one of them is a synonym usually means that my concept of ssp. is not split fine enough to include that one. Some splitting is inevitable as the bugs become better known (my 1986 book lumped a lot of ssp. that I now consider valid, some of which I would have considered valid had I seen specimens; and my lumpiest treatment of ssp.—the 3 “supersubspecies” into which I crammed all the *Euphydryas editha* ssp.—failed to acknowledge that *rubicunda* is quite distinctive also and should have been included). And it is true that there has been an inexorable splitting trend from Linnaeus to the present in genera and all other categories. Maybe the U.S. is doomed to repeat the European orgy of splitting that culminated in 200 ssp. of *Parnassius apollo*. But my goal is to recognize only the ssp. distinguishable by the ordinary person.

The goal is also to avoid bias in determining the status of ssp. and sp. BAD TAXONOMIST’S CREDO: “If I named it, it is a species; if you named it, it is a subspecies; if you named it and I don’t like you, it is a synonym”. I try to avoid that, but it’s definitely easy to dismiss what appears to be mindless splitting when one has not studied the taxa personally. In Colorado maybe I am known as a bit of a splitter, because I have named a new species of *Celastrina* (which actually may turn out to be a ssp. of the Cherry Gall thing), raised a “form” of one *Polygonia* into a second species (a ssp. of *oreas*), split one *Speyeria* species into two, split a common *Oeneis* in two, named a Neominois “second brood” which some authors have even considered a separate species, and have published studies splitting both *Anthocharis sara* & *Euphydryas anicia* into two species. So if I had lived for decades in those other places, I would have split some of them myself.

The book actually is justified, even though its mission is basically to name new subspecies, because the process of naming new ssp. in the regular journals is too slow and expensive; it would take \$50,000. of vanity-press charges (page charges, reprint charges, etc.) from the regular journals, as well as endless aggravation over 20 years, to get this complete coverage of the fauna. Of course, the ideal taxonomic paper naming a ssp. is a complete study of geographic variation of a whole species, listing thousands of specimens examined, but that takes a lot of time and money, which is generally not available. Sure, the piecemeal naming of ssp. and designating neotypes/lectotypes will inevitably result in some errors, but when we notice that about three-quarters of the names in checklists are synonyms anyway, a few more won’t sink the ship. A sudden flush of obvious synonyms should actually make it easier to get rid of dubious names from the past that were like vampire zombies that kept coming back after repeated whackings. (Ironically, the naming of hundreds of new ssp. makes it impossible for popular books to include any ssp., and makes it more difficult for conservation agencies to cover any of them.) And the naming of a ssp. automatically activates other lepidopterists, who collect and study the new ssp. and relatives, and they produce progress and correct errors. Basically, a new ssp.—good or bad—activates an army of lepidopterists who eventually do the work a full study would have done. And let’s face it: the ideal taxonomic study will work only on ideal bugs, and many of our bugs are the opposite of ideal; they are stencho-species (bugs that do not fit the biological species concept, bugs that have hybridized or introgressed or converged in a manner that appears random or incomprehensible, rassenkreis, or bugs that are inadequately studied because of haphazard choice of study sites or poor research, etc. Examples of stencho-species are *Papilio machaon/polyxenes/zelicaon*, *Colias alexandra/occidentalis-christina*, *Pieris “napi”*, *Cercyonis sthenele/meadii*, *Euphydryas chalcedona/colon/anicia/bernadetta*, *Chlosyne palla/[sterope] acastus/neumoegeni*, *Phyciodes tharos/cocyt/batesii*, *Limenitis arthemis/lorquini/weidemeyerii*, *Apodemia “mormo”*, *Callophrys affinis/perplexa/apama*, *C. sheridanii/viridis “dumetorum”*, *Plebejus anna/atrapraetextus/scudderii/melissa*, *Plebejus acmon/alupini*, *Euphilotes battoides*, *E. enoptes*, etc. Another generation of lepidopterists and several more decades will be required to make sense of stencho-species like *battoides/enoptes*.

All of the new ssp. are illustrated, mostly b/w. Unfortunately many of the photos in the book were printed poorly (too dark). Some of the few color photos are too red.

Several first-reviser actions are done below (capitalized), for latalinea=subaridum, ricei=calneva, davenporti=pratti, albihalos=austinorum.

Chapter 1 is an introduction to the book.

Chapter 2 fixes Boisduval's types. Later, conclusive proof showed that the labels on the Boisduval specimens in the Smithsonian that contain the wording "a/c Hofer", were written by Foster Hendrickson Benjamin while he was curator of the William Barnes collection (specifically in 1925 for most or all of the labels). And the symbols "a/c" mean "according to". And the person Hofer was Carl Höfer (with the umlaut). Scott explained all this in *Papilio* (New Series) #13. P. 9, Holland's lectotype (in his 1931 *Butterfly Book*) is the *sylvinus* female. P. 10, my *saepium* specimens from Tehachapi Mts. "fulvescens" and mine from San Diego Co. "chlorophora" don't look different from *saepium*. Most of the type locality restrictions in the book have logical justification, so are valid. The restriction of *Pyrgus ruralis* TL to the Feather River Canyon is justified by Lorquin's travels, correcting Scott's 1981 restriction to Tuolumne Co. Calif. However, the change of Scott's *caespitatis* TL (Marin Co. Calif.) also to Feather R. Can. is invalid, because their change has no justification (the text states that both Coast Range and Sierra Nevada have specimens matching the syntype, and their designation of the Boisduval TLs of *heteronea*, *saepiolus*, and *agricola* to Marin Co. proves that Scott's designation of *caespitatis* TL to Marin Co. is perfectly justified; therefore **I hereby correct the caespitatis TL to Marin Co.**; authors cannot just change TLs on a whim willy nilly without justification, therefore the first TL restriction that is justified must be accepted as the only valid TL. Their change of the Scott's 1981 TL of *pratincola* (Broderick, Yolo. Co., just W of Sacramento) to Feather R. Canyon is also unjustifiable and incorrect: they state the specimen could have come from the Central Valley, San Francisco Bay area, or Sierra Nevada, and their designation of the Boisduval TLs of *xanthoides* to San Francisco and *scriptura* etc. to Sacramento is consistent with Scott's TL; **I hereby correct the pratincola TL back to Broderick**; prior justifiable work done in a revisionary study as Scott's was, cannot be flippantly overturned for no reason; unjustifiable changes are invalid. P. 22 has several errors: Scott did not think that *pratincola* was *O. agricola*, and if Tilden's *pratincola* were to prove to be a distinct species (it isn't), the name *pratincola* would be perfectly valid for it. P. 22 and Miller/Brown checklist should have mentioned that Brown, Miller, & Clench (*Trans. Amer. Ent. Soc.* 106:77, 1980) designated a lectotype of *uricola* in the Carnegie Museum (the pale *Euphyes vestris* in Holland 1931 plate LXXIII fig. 12), which would mean that *vestris* is a syn. of *uricola*; however the O.D. is totally unlike *vestris* so that lectotype has to be invalid. P. 28 the descriptions of *saepiolus* & *rufescens* do not well match the photos; more later on *saepiolus* ssp. P. 30, Paul Opler told me that the Norden phenotype of *fuliginosa* does match Boisduval's description, thus this new neotype is invalid. P. 31, *callina* is now resolved, see the corrections on the Pelham Catalogue in this *Papilio* (New Series) #19. P. 33, the type of *irene* obviously belongs to *S. "atlantis" hesperis* (note the cream median unh spots, etc.), and not *S. egleis* as another lepidopterist has surmised (see the corrections of Butt. *Cascadia* below).

Chapter 5. Clyde Gillette told me that neither *A. mormo* nor *Eriogonum* occur at the designated TL Davis Creek Park, W of Washoe Lake, although p. 89 states that it is a "known locality"; perhaps inaccurate labeling of the "known" specimens? The citation of "Brown, 1965" at end of p. 100 should be "Three letters from Boisduval to Edwards, and the true identity of *Melitaea pola* Bdv. and *Melitaea callina* Bdv. *J. Lepid. Soc.* 19:197-211" in *Lit. Cited.* P. 102 *calchas* TL Mono Lake is low-altitude, suggesting it might be *minnehaha*, but the neotype looks like ssp. *shasta*, and a series from the mts. off Virginia Lakes Rd. in Mono. Co. looks like *shasta*, so I will treat *calchas* as a syn. of *shasta*. It is spelled *calcha* on p. 841, *calchas* p. 834 (apparently not another victim of the sex-trollers who troll the checklists and attempt to assign a sex or do a sex-change on every taxon).

Chapter 7. P. 115 professes a reluctance to redesignate TLs, which should have been applied to *caespitatis* & *pratincola*.

Chapter 8, 3rd paragraph describes the uns.

Chapter 11. *Tehachapina* is a syn. of *menapia*. *Sequoia* appears to be a syn. of *venosa* or very similar, as most *venosa* females are slightly yellowish also. *Mojavensis* isn't different enough from *pima* (and is slightly intermediate *cethura*X*pima* thus an invalid syn. anyway). *Desertolimbus* is described as intermediate *lanceolata*X*australis* (nearer *australis*) thus is invalid even if it isn't identical to *australis*.

Chapter 12. In the Calif. Coast Range *C. palla eremita* has females mostly cream, and in lower Sierras (TL of *palla*) females are supposed to be most often fulvous but are actually often intermediate or cream, then in high Sierra ssp. *altasierra* has females orange (rarely cream/orange), thus the proportion of cream females in *palla* is intermediate between *eremita* and *altasierra* and there are no other usable characters (except the male ups which may be less dark in foothill and high Sierra), so this is a cline and one of the three names must be synonymized. Nitpickingly it would be *altasierra* since it is younger than *palla* and both have most females orange and male ups may be slightly paler; however cream females are frequent in the Sierra foothills so I think it's best to call ssp. *palla* the whole mess from lowland Sierras to Coast Ranges, realizing that the percentage of orange females varies geographically somewhat; then *altasierra* is the paler orange high-Sierra thing and from Crater Lake to Mt. Jefferson. Ken Davenport notes that *altasierra* on E side of Sierra from Bridgeport to Donner Pass looks different from W side. P. 139, *sylvius* is a useful name for the female form with unspotted uns such as fig. 4 (females of all *P. faunus* ssp. vary from unspotted, to moderately spotted almost as much as males). Sierra Nevada *rusticus* is not "whitish gray" as stated, it has brownish-gray uns also (it is *hylas* & *cenveray* from Rockies that have gray uns). And the *fulvescens* were nearly all reared, which often distorts butterfly coloration, so it is doubtfully distinct,

evidently a syn. of *rusticus*. My series of *australomontana* from Big Pine Mdw. has yellower unh but the orangish ups looks like it has been hybridized with *acastus-neumoegeni*, as Davenport notes (in his Kern/Tulare Co. paper) also from Kelso Valley region, evidently because of introgression of *neumoegeni* genes into *palla*. He has a few from the summit of Nine Mile Can. on Inyo-Tulare Co. line that look like *neumoegeni*, as do some of mine from Big Pine Mdw. He wonders if the Kennedy Meadows region has a *palla australomontana* and an *acastus* that looks similar. *Australomontana* and *neumoegeni* are separate species, as in the Kelso Valley region both fly in many of the same places, with no intermediates, *neumoegeni* flying in washes and patrolling roads in L Mar-Apr., *australomontana* L Apr-May and occasionally overlapping flights, and they are sympatric on Pinyon Mtn. In the S Sierra ssp. *palla* and *australomontana* and *neumoegeni* (in Kelso V.) all feed on *Chrysothamnus*, so rearing should be done; maybe J. Emmel has found larval traits that may refer *australomontana* to *palla*. So *australomontana* is evidently heavily introgressed from *acastus/neumoegeni*, and based on phenotype it is similar to *C. acastus vallismortis* which also has yellowish unh because *acastus vallismortis* has evidently gotten genes from *palla* too (actually the proper name is *C. sterope vallismortis* since *sterope* is older than *acastus* and Jonathan Pelham has shown that *sterope* belongs to *acastus* and not to *palla*). These are *stenchospecies* (for instance, at Frazier Park the *palla* sometimes resemble *gabbii*). *Waucoba* is a syn. of *acastus* (evidently hybridized a bit also). *Eileenae* is a syn. of *pugetensis*. *Obsidiana* is basically a syn. of *mormonia*, maybe the unh is a bit darker on average?, my 6 specimens don't look different enough. *Tehama* is a syn. of *hydaspe*, *shasta* is invalid as it seems to be just an intergrade of *rupestris* X *liliana/elaine*, *hagemani* is a syn. of *rupestris*, *adiasteoides* is just the S end of *shasta* and thus is also invalid as an intergrade, *hanseni* is a syn. of *irene*.

Chapter 13. *Subaridum* is distinctive but is a syn. of *latalinea* (p. 617, J. Scott **first reviser action**). The photo of *caligulosum* holotype is bad (solid black), but Davenport & Mike Fisher told me the ssp. is extremely distinct (the uns evidently two-toned like the allotype). Most of the named ssp. of *saepium* are synonyms (my Pine Valley San Diego Co. *chlorophora*? series looks like my series of *chalcis/fulvescens* from Tehachapi Mts. and Colo. *provo* and S Ore. *saepium*, and photos of Wash. & BC *okanagana* are the same); evidently *chlorophora* if valid is only on "immediate coast" as p. 10 claims? *Rubrotenebrosus* looks like it might also be a syn. of *saepium*. *Hidakupa* looks like a syn. of *windi*. *Marinensis* seems to be *bayensis*X*windi* with no distinctive features of its own thus is invalid. The species assignment of the *stenchospecies* in *Callophrys* (*Callophrys*) are controversial as usual. *Superperplexa* larvae are said to be intermediate between *comstocki* and *perplexa*, though the adult spot pattern resembles *apama* and *comstocki*, and adults are said to be in a cline with *perplexa*. Ken Davenport finds pops. with prominent unh spots on W slope from Tulare to Madera Cos. (maybe having *superperplexa* genes?). *Pseudodumetorum* was assigned to species *C. viridis* by Gorelick (J. Lepid. Soc. 25[*suppl.* 2] table 3, larvae from Del Puerto Can. in Stanislaus Co.), but here it is mistakenly switched to *C. perplexa*, supposedly on the basis of larvae (which Gorelick placed near *viridis*), even though adults are similar to *viridis* "dumetorum" and *lemberti*, and according to old letters from J. Emmel to Scott, larvae have subdorsal ridges like *C. viridis* "dumetorum" and *lemberti* and larvae and pupae resemble *lemberti*; the leaf-feeding habit and larvae have led recent lepidopterists to suggest that *pseudodumetorum* (and *viridis* "dumetorum") belong to the same species as *C. sheridanii* *lemberti* (see A. Warren's Butt. Ore.), thus *sheridanii* becomes a ssp. of *C. "dumetorum"* (but the name *dumetorum* should and probably will be suppressed by ICZN plenary powers as I have so petitioned, so *dumetorum*=*viridis* will become ssp. of *C. sheridanii*). (Wash.-Ore. collectors insist that *affinis* and *perplexa* are distinct species based on decades of field study, because they occur in close proximity [at different altitudes, at Satus Pass area in Wash.], but close is not sympatry; *affinis* also occurs in close proximity with *homoperplexa* in S Utah [*affinis* in sage hills, *homoperplexa* in lower gulches] yet Gorelick [J. Lepid. Soc. 59:181-199, 2005] treats them as one species because they intergrade in SE Wyo.-Neb., [though the "intergrade" on fig. 5 looks nearer *homoperplexa* to Scott]. P. 165 says *pseudodumetorum* also occurs in close proximity to *perplexa*, which would suggest they are not conspecific, as Warren's Butt. Ore. noted. And the Univ. Colo. museum has 1f *perplexa* and 2m *pseudodumetorum* from Del Puerto Can. Rd. 2 mi. E County line, Stanislaus Co., Apr. 11, 1993, Reed A. Watkins, another sympatry that shows that *pseudodumetorum* is not conspecific with *perplexa*. So *pseudodumetorum* seems to belong to *C. sheridanii-viridis* rather than *C. perplexa* (I have petitioned ICZN to protect *sheridanii* from the older name *viridis*). Certainly, these are *stenchospecies*.

Chapter 14. P. 179, *insulanus* from BC is supposed to have tiny spots, but actually *C. Guppy* has found that adults from Saratoga Beach in N Vancouver I. are distinct with tiny spots, but the Victoria TL of *insulanus* has adults that resemble *amica* type to me, so I think *insulanus* is a syn. of *amica*. *Sabulosa* seems to be a syn. of *australis* (intermediate to *incognitus*, uns like *australis* but more tan, ups female brown like *incognitus*). *Excubita* is evidently a syn. of *daunia* (as are *toxeuma* & *nevada*)(someone should be investigating why the well-marked *sagittigera* flies so close to the washed-out *piasus* in S Calif., are they separate species?). *Gabrielina* seems to be a syn. of *sagittigera*. *Aureolus* is an extreme of *hilda* thus a syn. of *hilda*. *Littoralis* is evidently a weak ssp. similar to *saepiolus*. The *dedeckera* description fits *texanus* (photos horribly black), so I thought it must be a syn. of *texanus*, but Ken Davenport suggests *dedeckera* is very distinct; it "presumably" eats *Dedeckera*, a little bush with hundreds of tiny yellowish flowers in the *Eriogonum* family (*Polygonaceae*); so I'll leave it as a ssp. with *texanus* for now. *Argentata* is described as pale-silvery-blue, and Ken Davenport states that it is a syn. of *chlorina* based on research of J. Emmel, P. Opler, & Davenport (and my series from Big Pine Mdw. on *E. umbellatum* do look like the shade of blue of presumed *chlorina* from Lebec, Hungry Valley, & Frazier

Park, though none of these have more than a slight tint of green, whereas the several dozen beauties in Noel LaDue's series from upper Tehachapis had a slight greenish tint; San Bernardino Mts. *monticola* on *E. fasciculatum* are bluer. Current thought is *chlorina* eats *E. umbellatum* and *monticola* eats *E. fasciculatum* and they form a separate species *P. chlorina*. *Alpicola* would seem to be a syn. of *texanus*, as noted by C. Goodpasture. A lot of new ssp. of *P. icarioides* are named, which former authors such as John Downey and Larry Orsak felt were not worth naming. Scott dealt with these ssp. in *Papilio* (New Series) #12. *Argusmontana* is intermediate *fulla* X *albihalos* thus is invalid. *Panamintina* is a valid ssp. with large unh black pupils it seems (I have not seen specimens). *Albihalos* and *austinorum* look about the same as *panamintina* in the photos (pupils fairly large), but in my long series the unh pupils are smaller in *austinorum* and a little smaller still in *albihalos*, but these two are similar enough that *austinorum* should evidently be treated as a syn. of *albihalos* (since *albihalos* is farthest from *panamintina*). Those desert mtn. ssp. all tend to have a weak set of brown spots on uph margin as in ssp. *icarioides*, sometimes with weak orangish caps, but nothing like the orange margin in ssp. *evius* females; in the desert ssp. except *argusmontana* the unf postmedian black spots tend to be large (& wide in cell CuA_1). *Inyo* is *evius* X *albihalos* so is invalid, *eosierra* is evidently an intergrade of *icarioides* and *fulla* and *albihalos* so is a syn. *Santana* is a syn. of *evius*, *parapheres* is a syn. of *pheres* (many bad photos here)(note that after 20 years the ups blue changes "fades" toward violet on *Plebejus* and *Glaucopygma* etc. in the Cal. Acad. Sci. etc. collection, so ignore the many published statements about the blue being darker on extinct taxa of blues), *atascadero* is a syn. of *pardalis* (sl. intermediate to *evius*).

Chapter 15, *aestivalis* is syn. of *tecumseh*; a little different, but not enough for me. Ken Davenport says that most people who see a full series of *joaquina* are impressed by how it differs, so let's keep it as a ssp., but I have female *libya* from San Bernardino & Inyo Cos. Cal. & Lake Co. Ore. that are that cream on unh, so it is evidently weak. Davenport notes that Inyo Co. pop. is much larger, but it and Nev. pops. are not *lena*. The TL of *lena* is Montana, and the unh is solid gray-white in males and solid cream in females (both with a central whiter dash) on the *lena* type and in Colo. (*lena* range probably Mont.-Wyo.-Colo.-E Utah, and Scott's *confertiblanca* is evidently a syn. of *lena*); *lena* is whiter than "joaquina" and whiter than C Nev. pops., which I assign to ssp. *libya*.

Chapter 16 on *Euphilotes* has problems. It was written in 1988! and not revised since. The tables (tables 1-32) are a mess: p. 207 says that vein lengths were measured for Rs and CuA_2 veins on fw and hw, but vein Rs only occurs on the hindwing, and those measurements are not found on the tables. The tables do not contain uph aurora width measurements as p. 207 claims. Evidently every column of the tables refers to uns except the last Border M column. The amount of blue on upf is not on tables as P. 208 claims, which is just as well because the statement "this score is adjusted by adding or subtracting, respectively four times that proportion to or from four" (p. 208) is gobbledygook. And what do those 59.5...54.4 measurements mean on Table 1a??, they are not mm, are they tenths?? The tables are supposed to represent lengths of veins, spot widths, amount of unf blackish suffusion, distance from black dots to wing edge, and (last col. of the -b tables) the uph border width of males and the uph orange width of females and the amount of uph blue on females (these female traits are scored in some kind of indecipherable fashion). Unfortunately a lot of the characters in the almost-indecipherable tables are highly correlated with each other and thus are redundant, and some important characters are not represented on the tables. But it does not matter, as the tables are used little in the text anyway. Details are often disorganized, so one has to round up tidbits of data here and there throughout the paper and move them to their proper place. P. 209 claims without justification that host races or seasonal races are a step above subspecies, but don't some ssp. switch hosts in parts of their range? Most sympatric allochronic host races are claimed to be separate species in the paper, yet *E. enoptes cryptorufes* (the new ssp. on p. 209 top right) and *E. e. dammersi* are sympatric on Pyramid Peak. *E. bernardino* is obviously a distinct species reproductively isolated from *E. battoides*, but the same can't be said for *centralis* & *ellisii* & *baueri* (*baueri* flies at the hadrochilus site, but more than a month earlier). I am sure that a computer simulation of interbreeding of two sympatric but slightly allochronic populations eating different hosts would show that despite no inviability or infertility of hybrids (no prezygotic or postzygotic isolation), hybrids would have problems synchronizing with the hostplant they were or even weren't adapted to and so would produce fewer offspring, thus the two host races would persist there, even in the absence of reproductive isolation. In other words, two bugs that eat different *Eriogonum* species and are synchronized to the flowering period of those *Eriogonum*, are going to be highly adapted to those species and those flowering times, so hybrids between the butterflies are going to have a lot of difficulty producing offspring because the hybrids may choose a lousier host or emerge at the wrong time, so even if there is no courtship/mating isolation and no embryonic/larval/pupal inviability (thus no reproductive isolation) gene flow between hostplant ecotypes is going to be small, thus hostplant ecotypes may continue to exist sympatrically even if there is no reproductive isolation (and thus are just one species if reproductive isolation is your requirement for separate species). Thus the logic of separate species status for sympatric allochronic taxa is bankrupt, and *centralis* & *ellisii* & *baueri* are probably not even bookkeeping species. (The same problem will occur in the zone where a one-generation-adapted population meets a two-generation-adapted population, as Charles Oliver noted 30 years ago, as the hybrids often have fewer offspring because they are not synchronized as well.) Ken Davenport suggests that a species *E. glaucon* probably includes both *intermedia* [which now includes ssp. *comstocki*] and *centralis*, in other words most of what was once in *E. battoides*. Ken Davenport notes that ssp. *battoides* and ssp. *glaucon* co-occur at Warren Creek E of Tioga Pass, suggesting they are different sp. (*battoides* occurs at 11,000' on top of Glass Mtn. E of the Sierra totally surrounded by *glaucon*, though if they don't overlap altitudinally this is

no test of reproductive isolation). Note that Miller/Brown lists the wrong years of publication of *tildeni*, *bayensis*, *smithi*, *columbiae*, & *martini*. P. 210, it would seem that *bayensis* is a syn. of *tildeni*, and *smithi* isn't much different either. *Arenacolor* is also a syn. of *tildeni* (a weak ssp., Oakley Shields informed me), and its larvae (3rd paragraph of page) are closer to *tildeni*/*bayensis*. P. 211, *langstoni* is "probably" closer to ssp. *enoptes* and *tildeni* than *mojave*, implying uncertainty (the only real evidence that *langstoni* and *mojave* are separate species comes from a 1988 letter from Pratt stating that *langstoni* flies higher on its host *E. nudum* than *mojave* flies on its low-flying host *E. pusillum* in Jawbone Can. area, where larvae are also distinguishable but may show some gene flow). P. 211 bottom, one wonders if spring females are bluer due to environment, as in *P. acmon*. P. 226 adds *E. plumatella* to the *dammersi* hosts. *Cryptorufes* is placed in *enoptes* but p. 213 says it may be a ssp. of *mojave*; p. 213 says it does not overlap with *mojave* but p. 211 says it may overlap with it geographically near Pinyon Flat. Mattoni (J. Res. Lep. 27:175) says the female genitalia of *mojave* differ. P. 213 says that *E. ancilla* may be an artificial conglomeration, and p. 215 says the wings of the ssp. resemble sympatric *battoides* more than *enoptes*-complex taxa. *Columbiae* could be considered a syn. of *ancilla*, but A. Warren's Butt. Ore. retains both because of phenotypic differences and different hosts. P. 216, in the *battoides* group, *mazourka* & *panamintensis* fly after spring in mid July. P. 225 lists more hosts for ssp. *battoides* to add to those of p. 216. P. 217 notes a gap between *australoglaucan* and typical *glaucan*, but actually *glaucan* flies at Lookout Mtn. 8350' filling the gap (specimens sent to me from Derham Giuliani). *Mazourka* and *australoglaucan* and *panamintensis* seem to be synonyms of *glaucan*. P. 220 *intermedia* is said to be "sympatric" with *battoides* (*intermedia* at lower elevations on *E. lobbii* & *marifolium* according to p. 214). P. 221, Pratt (e-mail to Ken Davenport) now suggests that *vernalis* belongs to *E. baueri* because it is sympatric & synchronic with *battoides* in Coso Mts. on *E. nudum*; though how could those pops. so far apart be the same ssp.?, and Pratt states the *vernalis* allozymes are closer to *glaucan*. This paper claims that *allyni* belongs to *E. battoides*, then claims that other "dark coastal" populations belong to *E. bernardino*, based on unpublished allozyme work etc. Pratt (e-mail) states that there is little difference in wing pattern between *allyni* on *E. parvifolium* and nearby "near *bernardino*" pops. on *E. cinereum* but claims allozyme differences "in sympatry". Allozyme work is cited in many places, but that work has not been published or properly analyzed, and allozymes have proven to be problematic data that mostly are not relevant to the characters distinguishing the wings and bodies of the butterflies or their hostplants or behavior (most are neutral alleles); and allozyme studies are virtually never verified by other authors and thus amount to blind faith; and based on the recent total failure of mtDNA to help in *Phyciodes* and many other taxa (mtDNA is very variable and overlaps greatly between species, Wahlberg Oliveira & Scott, Syst. Ent. 28:257-273; and a little bit of introgression can result in the mtDNA of one bug completely taking over in the other bug, as has happened in *Papilio glaucus appalachiensis* [see Butt. W.Va. & their Caterpillars below] & *Phyciodes diminutor incognitus* & *Polygonia progne*, etc.), I do not think allozymes should be given much weight. Mattoni published a paper on *bernardino* (J. Res. Lep. 27:176, 1988) in which he carefully and cogently argues that *allyni* is part of *bernardino*, with *garthi*, based on wing pattern and 4th stage morphology, size & hosts, and on the wing pattern of *bernardino* 1 km S of Malaga Cove that overlaps wing pattern of *allyni*. P. 223 emphasizes *allyni*'s fusion of M₁₋₂₋₃ macules with spots bordering orange aurora, but *garthi* from Cedros I. has these 3 macules nearly always fused & has larger spots. What to make of this disagreement? Mattoni's treatment would seem preferable. P. 222, *centralis* does not eat *E. umbellatum* or *flavum*, which are errors. P. 222, *hadrochilus* surely belongs to *battoides* (actually to *glaucan*, since *battoides* & *glaucan* are different sp.) and not to *centralis*, based on having the same host as *mazourka*/*panamintensis*, flight time like *mazourka*/*panamintensis*, wing pattern like *glaucan*, etc. (and look at the map, it is ludicrous to have this one pop. from Westgard Pass belong to a species 400 mi. to the east). *Inyomontana* is misspelled –us on map (another victim of the sex-trollers?); it seems to be a syn. of *bernardino* intermediate toward *martini*, and Oakley Shields suggests it is closer to *martini* (female ups bluer etc.), thus would seem to be invalid as an intermediate (Ken Davenport notes some difference but says it's hard to place exactly where *bernardino* & *inyomontanus* separate because of individual variation within populations—and surely a ssp. isn't worth keeping if the variation isn't much greater than individual variation). P. 226 *euromojavensis* host in Spring Range is actually spelled *heermannii* var. *sulcatum*. P. 226 the W-C Nev. *heermannii* pop. was named *ellisii basinensis* on p. 551. P. 227 holotype *avawatz* evidently coll. Aug. not July; it looks like a syn. of *euromojavensis* to me but Oakley Shields suggests it is distinct.

The maps of *enoptes* & *battoides* are a complicated mess, and benefit from a lot of pencil work. On the *enoptes* map, the "giulianii" symbol near Lake Tahoe/Carson City must be an error for ssp. *enoptes*? A *tildeni* dot can be added for Coso Range. *Ancilla* occurs in S Alta. and S Sask. The Marina Beach dots are *enoptes arenacola*. On the *battoides* map, both symbols from east-central Nev. (the "inyomontana" dot and the "battoides on *Eriogonum shockleyi*" dot) are actually the same bug, *bernardino minuta* on Baking Powder Flat, named by Austin on p. 549-50. The Coxey Meadow dot is *battoides vernalis*, the "centralis" *panamintensis* is *battoides* (really *glaucan*) *panam.* on p. 219; *inyomontana* is misspelled. The island pop. W of Baja Calif. is mapped as dark coastal *bernardino*, but was named *garthi* by Mattoni, a pop. even more heavily spotted than *allyni*. P. 813 lists "near *garthi*" from Pt. Loma San Diego Co.

One disturbing fact: I have two specimens, which have one valva like *enoptes*, the other like *battoides*. Does this have devastating implications for the taxonomy of *Euphilotes*? Is there valval polymorphism due to dominant/recessive genes, such that *enoptes* & *battoides* are really the same species, or several species that each are polymorphic in valva form?

Chapter 17. *Confusa* is intermediate *pallescens*X*elvira*, thus invalid. There are many such intermediate populations. Davenport notes that Mono Co. pops. are extremely variable, some at S end of county may be *elvira* while others cannot be placed into ssp. The separation of *E. pallescens* from *E. rita* is still dubious. P. 272 states that hostplant is sometimes not a good clue to taxonomic relationship in *Euphilotes*, as the bugs switched to an annual.

Chapter 18, *calneva* is a good ssp., however it is intermediate toward the even more extreme *ricei* on p. 815, therefore *calneva* is invalid and as a **first reviser action** I declare *ricei* valid and *calneva* a syn.

Chapter 19. *Purissima* I thought might be a syn. of *bohartorum* which is supposed to have a wide upf border, though Oakley Shields and Ken Davenport say it is quite distinct (smaller spots)(but only half a dozen *bohartorum* have been caught, so how do we know what it looks like?). There are evidently a number of local odd populations of this tiny hard-to-see bug, including *P. speciosa leona* in Ore.

Chapter 20. *Cassiope* is an outstanding find of a new biological creature (eating the odd *Cassiope mertensiana*, a spreading juniper-leaved bush <30 cm tall with little white bell flowers), with less upf blue near upf apex and unh postbasal discal cell spot vestigial, the upf border wider. Obviously *podarce* is a distinct species from *A. glandon*, with its conical unf submarginal spots and different flared-out white larval dashes and pupal abdomen etc. *Cassiope* would seem to be a ssp. of *A. glandon* related to *megalos*, and the larvae and pupae (figs. 1-3) are similar (and variable, comparing 1b with 8); however the recent claim by Steve Kohler that he has found *cassiope* in western Montana means—if true—that *cassiope* is a separate species too if it occurs with *megalos* there. *Kelsoni* appears to be a weak ssp. with more linear unf submarginal spots (a little more similar to *megalos* than is *cassiope*). *Cilla* unh looks a little darker than *podarce* on the photos but this trait isn't mentioned in text, only the larger unh spots are noted but these are stated to have "considerable variation", so perhaps *cilla* is a syn. *Klamathensis* is very different in Davenport's series so is a ssp. (my one male from Scott Camp Crk. has wide upf border and does not look different).

Chapter 21. *Nesiotes* is a syn. of *amyntula*.

Chapter 22. *Pseudoxerces* from Santa Rosa/Santa Cruz Is. isn't much different from *australis*, but unh black pupils are a bit smaller thus the white spots are more apparent; it can be left as a ssp. as it isn't an intermediate to *xerces* (black spots much larger on unf, whereas in *xerces* black spots are about the same size on unh as unf), and it isn't *xerces* as several people say, though maybe some of the same genes may be active. P. 309, *xerces* are darker more violet blue now because of fading of the blue.

Chapter 23. *Catalina* is intermediate *santacruz* (closer)X*sylvanoides* thus invalid.

Chapter 24. *Channelensis* is a syn. of *sabuleti*?, or maybe a little darker (most book photos are dark).

Chapter 25. P. 322, ice-age lowering of sea-level by 300 feet brought biota to Channel Islands that could not move northward as mainland biota did in interglacials (in glacial maxima the Californian biota moved to Baja Cal., in interglacials including the "Recent" the biota moved back north). *Hypoleuca* is an intermediate *sthenele*X*behrii* without any different characters of its own, near *sthenele* but the unh white marks not quite as strong as the San Francisco *sthenele* fig. in Chapter 2 fig. 115 and in Comstock's Butt. Calif. & Holland's Butt. Book, thus *hypoleuca* is invalid, and is a syn. of *sthenele*. Another fantastic discovery!, which can be called near-*sthenele*.

Chapter 26, figs. show basal upf red streaks, and *elegans* is a syn. of *wrighti*.

Chapter 27, p. 333 repeats the folly split of *fulvia* from *leanira*, which was superficially based only on adult traits, despite the intermediacy of *pariaensis* (which has intermediate wing color plus 2 traits of *alma* & 2 of *fulvia*), and despite the fact that the ssp. change gradually (step clines) from *leanira* to the intermediate "cerrita" to *alma* to *fulvia*, and the different characters (older larval body color, width of black larval bands, number of white dots in the bands, larval head color, adult color & pattern) change in different places, all explained in Scott 1992 *Papilio* (New Series) #6 and Scott 2006 *Papilio* (New Series) #12. *Obsoleta* is an interesting extinct ssp., correctly a subspecies and not an aberration. P. 344 the ~25 glaciations in the last 2.5 mya surely affected the Colorado Plateau greatly also. *Oregonensis* & *nebularum* are syns. of *leanira*, *elegans* is a syn. of *wrighti*, *austrima* is a weak ssp. with a bit more ups orange than *wrighti*, *basinensis* & *flavodorsalis* are syns. of *alma*.

Chapter 28. I think *C. chinatiensis* is still a distinct species, because the "intermediates" shown on figs. may fall in the range of variation of ssp. *bolli* (which looks intermediate even in NE Mex.)(thus 175-177 can be interpreted as *chinatiensis*, 169-174 & 178-183 as *bolli* [though 169 & 170 do look somewhat intermediate], and note that 175-177 are 1988 and 178-183 are 1987 from same site, as if the climate favored one sp. one year, the other the next), and one dot of *chinat.* is in Durango Mex. right next to a *thekla* dot, and the Stanford/Opler county record maps show both *chinatiensis* and *theona* from all over W Tex. and Eddy Co. NM (plus Hidalgo Co. NM & Cochise Co. Ariz. though, where *chinatiensis* is dubious), and J. Glassberg (Butt. through Binoculars The West., p. 162) found both *theona* & *chinatiensis* in Chisos Basin. More collecting in Tex. is needed. *Ezra* is prob. a ssp. of *theona*, and is somewhat intermediate *theona*X*perlula* but not intermed. enough to be invalid; the "lectotype" of *ezra* would seem to be the holotype. *Perlula* is prob. a ssp. of *theona*. *Minimus* is a syn. of *theona*.

Chapter 29, *ehrichi* is similar to *editha* but may have more extensive cream bands thus is valid (don't confuse it with *augusta* [Edw.] which is a syn. of *quino*), *karinae* a syn. of *editha*, *bingi* a syn. of *nubigena* (and as transitional to *aurilacus* is an invalid intermediate), *trinitina* a syn. of *sierra*, and *varicolor* a variable intergrade population thus not a ssp.

Variicolor from TL Sonora Pass was first described by Scott ("1978", J. Res. Lep. 17:245-252), whose 5 specimens from above Sonora Pass had valva prong type frequency of 0,0,0,1,2,2 (short to long), thus the genitalia is closest to *anicia* as p. 401 admits, so why call it *E. chalcedona variicolor*?? (the checklist on p. 828 plops it into *E. anicia*), evidently it actually should be called *wheeleri-sierra-macglashanii*? (corrections to Scott 1978: the Hallelujah Jct. Pop. should be called *macglashanii-wheeleri*, the King Can. site *macglashanii*, the Sonora-Ebbets Pass sites *wheeleri-sierra-macglash.*, the Carson Pass site *sierra-macglash.*, the "quino" sites *hennei*, the Satus Pass site *colon-paradoxa*?) and anyway is an intergrade pop. so is invalid as a ssp. The Carson Pass alpine pop. is also very variable from black to orange, yet has very variable prong length frequency (0,16,19,5,1,0, see Scott p. 251), thus is closer to *E. chalcedona*. These two alpine sites show intermediacy between *E. chalcedona* and *anicia*. Why did they name the Sonora Pass pop. and not the Carson Pass pop.??

Chapter 30, *koreti* is a syn. of *lehmani*, *tahoensis* & *bingi* are syns. of *nubigena*, *monoensis* a syn. of *edithana* (same ovip. on *Collinsia parviflora* too), *owyheensis* and *mattoonii* are syns. of *edithana*.

Chapter 31. This is a nice paper by Austin & D. Murphy, clearly showing blending between different phenotypes with photos of numerous specimens. (I wish many other chapters of this book had more good mass photos and less text.) P. 420 *klotsi* is not redder than *corralensis* & *kingstonensis*. P. 421 Comstock (1927 pl. 35) figures *wheeleri*—not *olancha*—from Mono Lake itself. P. 422 the wings of *wallacensis* don't look more pointed than *veazieae* on the photos. The leftmost *wheeleri* from Pine Valley Mts. in Fig. 7 looks like *hermosa*. Ssp. *klotsi* from C Ariz. is like *hermosa* but darker on ups, the yellow spots more prevalent than red ones. *Kingstonensis* in wing pattern is a syn. of *hermosa* (distinguished only by a slightly greater frequency of yellower specimens) yet they have different valva prong so are valid ssp. that have been placed into *chalc.* and *anicia* (surely they look similar because of gene flow). Ssp. *wheeleri* is a little redder than those. Ssp. *morandi* is variable but has dark basal half of uph, fewer yellow spots than *wheeleri*, ups often tawnier overall. Ssp. *olancha* (higher alt. in Mono Co. than the lake) is distinctively white-spotted. Ssp. *sierra* is the distinctive reddish higher Sierra bug. Ssp. *macglashanii* is a mostly-whitish mess with some redder ones, and looks like *veazieae-macyi* in wing pattern, except the valva prong places them in different species (their similar wing pattern surely proves gene flow doesn't it?); Scott's *macglashanii-wheeleri* series from N Hallelujah Jct. Lassen Co. (J Res. Lep. 17:150) showed a lot of variation with some red adults and some long prongs. *Macyi* is like *veazieae* but a bit redder, and one wonders if they are separable over their whole ranges from Nev. to Ore. & Wash. From the photos *nevadensis* is a syn. of *wallacensis*. *Alena* is similar to *wheeleri* (the photos in fig. 7 show strong contrast between uph central yellow & outer red, but photos in other books [Butt. Colo., Butt. Rocky Mts., Howe] and my specimens do not show this). *Maria* is evidently distinctive and perhaps has introgressed from *colon* as its larva is somewhat white-hairy like *colon*.

Recently Austin et al. (J. Lep. Soc. 57:176-92) changed his mind and split the Nevada bugs into two sp. *E. chalcedona* and *E. anicia*, mostly because of a change of species concept to the "phylogenetic species" (a pompous name for bookkeeping species, a portion of a stenchospecies that must be broken off and treated as a separate species after the areas of interbreeding between them are ignored, which allows each taxon to be treated as a separate species where they do not interbreed); *E. chalcedona wallacensis* and *E. anicia veazieae* are sympatric without interbreeding (though their flight periods only partially overlap) in NW Nev. (suggesting two species), while in NE Nev. *E. chalc.* "nevadensis" interbreeds freely with *E. anicia wheeleri* in the small altitudinal and temporal overlap between them (suggesting one species). (In *Papilio* (New Series) #12, I discussed the whole stenchospecies in W N.A., and now it appears that *E. colon* may be a valid bookkeeping species.)

Chapter 32. As noted in *Papilio* (New Series) #12, *carolae* is an obvious syn. of *eurytion* whose TL is identical to the *eurytion* TL of "probably South Park Colo." (note 521 in Miller/Brown). P. 433 NW not northeastern. Host must be *Cast. integra* (*chromosa* only occurs in SW Colo.) if they eat *Castilleja* there, although S. Spomer found no feeding damage on that plant at TL, and Frank Stermitz reported *Besseyia plantaginea* in S Park also (*Besseyia alpina* is a known host for ssp. *brucei* in Colo.).

Chapter 33, *yolaboli* is intermed. to *egleis* thus invalid, and should be treated as a syn. of *oweni*. My Warner Mts. series is near-*egleis*.

Chapter 34 on *carolae* displays warped logic: if *carolae* is intermediate thus lower in status than distinct species, why is it raised higher in status to a species??? The holocentric chromosomes in Lepidoptera mean chromosome # does not matter, because even if broken the pieces go to the proper end of the dividing cell. In table, fw shape of female *zerene* is rounded or square. Fig. of *carolae* male fw shape resembles *malcolmi*. Size is larger in *coronis* & *carolae*, smaller in *zerene*. Most traits of *carolae* in my long series fit *zerene*, except the first two in table (fw shape male & female) which are weak; the good traits of overscaled unh submarginal band and triangular unh marginal spots and spots outside unh postmedian spots are similar to *zerene*, thus *carolae* is actually a ssp. of *zerene* closest to *malcolmi*. S. *coronis semiramis* should have been figured too.

Chapter 35, *puntareyes* is syn. of *myrtleae* (photos too light [overexposed]).

Chapter 36, *sonomensis* is syn. of *myrtleae* (& photos look darker not lighter).

Chapter 37, *giulianii* looks like a syn. of *lasus* slightly intermed. to *macswaini* on the photos, but Davenport and J. Brock think it is much different (evidently because of "very bright fulvous ups" which isn't apparent on the photos).

Mattoonorum is spectacular-looking (but note that only some have tiny unh spots as shown on photos, most in my long TL

series have larger unh spots), so is a ssp. if it isn't just a local intergrade; my only question is, how widespread and stable is this phenotype?, because MacNeill described many odd populations of *dodgei*, so are there other *dodgei* pops. this spectacular also? Shapiro (J. Res. Lep. 29:35) showed the variation of *H. comma* at Asbestos Gulch Trinity Co. (some individuals look like 3 ssp. incl. *mattoonorum*, though that site is a blend zone near ssp. *idaho*).

Chapter 38 is not good and has many problems. *Ssp. lindseyi* is not defined, as the *lindseyi* TL is not mentioned ["Nellie & Ukiah" in Miller/Brown, Napa Co. Calif. on label on type], the locality of the ssp. *lindseyi* photos is not given, and the paper does not state how *macneilli* differs from *lindseyi*. Using my crystal ball, I deduce that evidently the TL of *lindseyi* is supposed to be Inner Coast Range, *macneilli* Outer Coast Range. P. 478 claims *septentrionalis* is lighter but photos are darker. P. 476 text says *lindseyi* ups is darker and unh spots less contrasting but photos show the opposite. At any rate, all 3 (*macneilli*, *eldorado*, *septentrionalis*) are syns. of *lindseyi*. If they were distinctive, MacNeill would have named them in his *Hesperia* book. *Septentrionalis* might be a valid weak ssp. with larger unh spots, but the spots on the ssp. *lindseyi* photos are just as large. K. Davenport notes that S Calif. *lindseyi* look different from Monterey & Fresno Co. *lindseyi*, none of which were covered in this incomplete paper, which is so disappointing that I don't want to consider these weak names seriously.

Chapter 39, "harpalus" is now *idaho* (Scott, *Papilio* [New Series] # 11). *Mojavensis* is intermediate *idaho*X*susanae* (its unh spots bigger than *susanae* but like *idaho*, unh paler than *susanae* most often like *idaho*), thus is invalid and best lumped into the synonymy of *idaho*. The same phenotype is on N Rim Coconino Plateau in Ariz.

Chapter 40, *sierra* seems to be a syn. of *nevada*, with "greenish-olive" unh, the unh spot size on photos is the same size as in Colo. Paper doesn't mention my 1992 paper (*Papilio* [N.S. #6] noting that Front Range Colo. *nevada* have sl. browner unh than sl. greener South Park (TL) *nevada*. "Nr." in fig. 1 legend means NV.

Chapter 41, *gentryi* has wing pattern like *freemani* but unh spots weaker, so is a ssp. of *A. aryxna*. Again, chromosome # does not matter in holocentric chromosomes (*baueri*'s # of 15 is similar to *gentryi*'s 13). Paiute is described as "orange-yellow" but mine are yellowish, so maybe this isn't a valid ssp. Lots of "species" have been described in *Agathymus* but insufficient attention has been paid to relationships (for instance, *valverdiensis* & *estelleae* are obviously same sp. as *A. remingtoni* when you carefully consider ranges/wings/anatomy/host/ larvae; my 1986 book was **first reviser** in treating the former as syns. of *remingtoni*.) It's easy to argue that even *A. alliae* could be considered a ssp. of allopatric *aryxna*.

Chapter 42. *Indistinctus* is a syn. of *pylades*, *eureka* a syn. of *phyleus*, *flavus* a syn. of *wrightii* (if you accept *wrightii* as valid--*wrightii* isn't much different from *eunus*, just a little paler, so you may want to treat *wrightii* as a syn. too), *obscurus* a syn. (a bit darker) of *eunus* (there are too many names for grades of pallidity in *P. eunus*; *alinea* is the only really distinct ssp.). P. 503 right says *e. eunus* is smaller in spring, but stats show *e. "flavus"* is bigger in spring--?? G. Austin told me "obscurus" is extinct at TL but still flies near Pyramid L. P. 504 spelled *Dechambeau*?. *Tenebricosus* is a syn. of *campestris*; photos are too dark. P. 505 top, both darker and much yellower? *Transmontana* is a syn. of *coloradensis*; Davenport suggests it is consistently larger & lighter than *coloradensis*, but the photos don't look any different than the gazillions I have caught in Colo. P. 507, # of spots varies in *titus*. *Occidentalis* is a syn. of *Satyrium titus watsoni*. *Purpurascens* is a syn. of *sheltonensis*. *Clenchi* is a syn. of *Strymon istapa istapa*. *Deserticola* is a weak ssp., the same as *oro* but unf spots a bit bigger. *Albomontanus* is a syn. of *gertschi* (see *Papilio* [New Series] #12). Horrible problems in writeup of *L. carinenta streckeri*, which is actually a syn. of *larvata*. First, *streckeri* was named by Field in 1938 (Univ. Kansas Biol. Ser. 39:132) as an infrasubspecific "dimorphic form", therefore the subspecies must take authorship & date of *L. car. streckeri* Austin & J. Emmel 1998. (Note that Scott & Wright [J. Res. Lep. 30:258 bottom] warned readers about a grossly misleading example at end of 1985 Art. 45[g]. We wrote to inform ICZN that this example was rotten, yet in the 4th ed. 2000 rules the mistake is turned into code as the similar Polinski example at end of 45.6.4.1. shows. Tolerating bad examples; turning errors into rules; promoting instability by changing sexed suffixes; giving half the space to obscure details of latin grammar; allowing the principle of priority to force taxonomists to waste thousands of man-years studying rotten old names and types; that is the unchanging legacy of the anachronistic authors of the ICZN Code. Evidently it will take decades, and the death of the current officers, for the ICZN to become a valued part of biological science.) The whole point here is that the *larvata* TL is San Antonio, so *larvata* is whatever population flies at San Antonio, not what the syntypes look like (the pop. may vary somewhat so the consensus is what matters), and I understand that at San Antonio the pop. is more like the Mexican than the E U.S. ssp. (my one San Antonio specimen is like the Nev. specimen figs. 47-48, and San Antonio is in the range of the Mexican thing in Butt. Tex.). The TL of *streckeri* is Donna in Hidalgo Co. Tex., so it is the Mexican thing like *larvata*. Figs. 49-50 is actually ssp. *bachmannii* from Dallas (coll. by Kemner, not Klemmer)! They should have shown photo of *larvata* holotype. They misinterpreted the biology and geographic variation. The N Mexican ssp. *larvata*=*streckeri* is not a local phenotype, it is widespread from S Tex. to Ariz. and northern Mex., and these bugs are MIGRATORY so the traits of two syntypes may not matter (migrants into Colo. are mostly *larvata* but some are *bachmannii*). The costal white spot is widely fused to the outside-cell spot in *bachmannii* and offset in *larvata*, and they fail to note that important character on *larvata* syntypes (their statement that the spot on costa and that at end of cell and that in cell M_3 - CuA_1 are in line on *larvata* suggests that spot in costa is not fused to that at end of cell as it is in *bachmannii*, thus the *larvata* syntypes must be more similar in this critical character to Mex. ones)! The orange on rear of fw is near discal cell in many *larvata*. (By the way Timothy Friedlander examined numerous U.S. *Libythea* and published his conclusion that

carinenta and bachmanii are distinct sp. based on genitalia, and then seemingly everyone thought they were two species. I sent him a series from Acatlan, Puebla, in which wing pattern is all larvata yet genitalia varies from bachmanii to carinenta, and then I wondered why suddenly everyone accepted that they were one species, maybe because Friedlander told others?) Grisea is very weak, basically a syn. of coenia. Powellii was named as an infrasubspecific “transition form” by Field in 1936, so the subspecies name takes authorship & date of L. lorq. powelli Austin & J. Emmel 1998 (the description and stated location of holotype and figure all seem to make the name available). P. 515, the posterior fw ocellus is much smaller in all ssp. Paulus isn't a Great Basin endemic as masoni from Colo. looks the same to me (however Davenport & J. Emmel state that E Mojave Desert specimens are closer to masoni than to paulus, and Spring Mts. sthenele look like Colo. masoni; Davenport states that paulus is lighter brown on ups and has more whitish unh scaling, masoni more umber brown on uns; but I am not sure I can see any difference between paulus & masoni in my drawer, so it's not a very good ssp. Davenport notes that fresh masoni often have a strong rusty flush on unf apex, which is introgression from rufous Cerc. meadii which interbreeds with C. sthenele in N Ariz.-S Utah.).

Chapter 43. Profugus is a syn. of huachuca. Apertorum is a syn. of scriptura (fails to note that spring adults have larger white spots, which Scott 1986 called form pseudoxanthus). Flavaventris seems to be a weak ssp. similar to sonora, with slightly yellower uns. Longinqua is a syn. of sonora (intermediate sonoraXflavaventris, thus invalid). Omnigena is a syn. of sylvanoides (K. Davenport notices the difference, but I don't want to recognize a ssp. with this small a difference), and is sylvanoides (very near)Xbonnevillea thus an invalid intermediate anyway. P. 527, napa is a syn. of sylvanoides also. Sacramentorum is a syn. of yuma. Lutea photos look like a syn. to me from the photos, but A. Warren insists it is different, with yellower unh (the shinier unh areas in photos evidently). P. 528, Mt. St. Helena is surely mislabeled.

Chapter 44. Magnamenapia is a syn. of menapia (noticeably larger, but not different enough for me). Pseudothoosa is similar to ssp. sara and ssp. thoosa, closer to sara. It is a little different from both, but in characters and range it is intermediate saraXthoosa thus would seem to be an invalid syn. The paper should have discussed the relationship with ssp. sara. (Anthocharis is discussed in Papilio [New Series] #18.)

Chapter 45. Incana is a syn. of rubidus. P. 540 “olive-gray” is dubious, as it surely is not green (I hate the color “olive”, is it a green olive or a brown olive or a black olive??). Rava is a syn. of heteronea. Rutila is a syn. of heteronea plus a homonym of European L. dispar rutila/rutilus. Praetexta is a syn. of browni, lavender on only outer 1/3 of unh like browni (nivalis is lavender on outer 1/2). Intermedia is a syn. of citima (citima often has larger orange spots, they aren't always small). Obscurafacies is evidently a weak ssp. (uns a bit darker the dots a little larger than californica), & Warren's Butt. Ore. implies it is a syn. Megapallidum lacks tails like dryope, but is larger and uns is whiter so most lepidopterists consider it valid. Obscurofuscum may be a syn. of saepium (the bad photos are too dark; the photos are truly obscurofuscum). Concava is a syn. of augustinus and has nothing to do with iroides (the unh base is dark on augustinus, pale on iroides, and the median unh band averages more of a “concave” nook in augustinus [because the posterior portion of the band has a bit more of an anterior jaunt that forms the rear edge of the nook], a little less of a nook in iroides [because the posterior portion of the band averages straighter]); augustinus and croesioides seem identical also (the only different ssp. in this augustinus group is the new chocolate ssp. in N BC), and iroides and annetteae seem identical. Some people think the polyphagous iroides is a separate sp. from the Arctostaphylos uva-ursi-eating augustinus, but where's the proof? Mojavensis may be a ssp., but why is TL only a couple dozen miles from the designated TL of fotis? My series from Tooele Co. lacks ups orangish. The range would make sense if mojavensis occurred only N and fotis S of Colorado R., but paper states that fotis also occurs N in Virgin Mts. Pallescens is syn. of eryphon. Montanorum looks like Colorado amyntula valeriae, but is evidently distinctive (maybe the postmedian spots are more-distal); it not a syn. of amyntula which occupies the Great Basin. Minuta is evidently a valid ssp., that even looks like allyni in border & some unh spots. Basinensis is a ssp., but it may belong to E. bernardino where Shields placed it, as the wing pattern & flight period fits bernardino not ellisii, and this isolated pop. doesn't fit with the range of ellisii 300 mi. away on the map, though p. 226 says larva is more similar to euomojavensis. Fusimaculata is a syn. of panamintensis and glaucon. Shieldsi is a syn. of purpura (even Oakley Shields agrees). Gilvatunica may be a syn. of ancilla (& the uns spots are large as they are at ancilla TL), but there is controversy about how spotted toptype ancilla are (see Papilio [New Series] #18). Opacapulla is a syn. of dammersi. Aridorum is a syn. of enoptes. Primavera is a syn. of langstoni. Arenamontana is a syn. of pallescens (& uns spots are not smaller than pallescens, see p. 282). Septentrionalis is a syn. of speciosa. Minipunctum is a ssp. or just a var. of deserticola with sl. smaller spots. Alateres is a syn. of melissa. Pallidissima is a syn. of minnehaha (Ferris' [1976] plates don't show any difference either). Goodpasturei is a syn. of alupini. Autumnalis & parva are syns. of mormo.

Chapter 46. Carsonensis is a syn. of apacheana; even 658 types can't make this name valid. Reidi is a syn. of utahensis (as is toyabe). Expedita is a syn. of arachne. P. 576, monache has much more unh white. Robusta is a syn. of vallismortis (my 2 actually have upf median posterior black patch half the size of that shown for “robusta”, and my vallismortis have the large black patch shown, and the ups orangish and unh yellowish doesn't seem to differ either). (The checklist p. 842 lists dorothyi from Nev., which is very dubious.) Arenacolor is actually a valid local ssp. distinguished by large uph postmedian dots, melanic female upf, whiter female unh (but the comparison C Utah sample could be either P. cocyta selenis or P. batesii anasazi). Pullum is a syn. of milberti. Alkalorum is the extreme of--thus a syn. of--pallescens whose TL is nearby, as a cline oetus-pallescens-alkalorum can have only two—not three--names.

Chapter 47 is a nice paper with lots of photos of the variation of *Nevada tullia*, incl. intergrades *elko*X*pseudobrenda* in NE Nev. and introgression *ampelos*X*pseudobrenda* in C Nev. P. 587 the color olive again, the wing is NOT olive-brown! (in Ventimiglia Italy I sampled about 20 kinds of olives at a market, which were all colors from green to brown, all sizes, yet all were so horribly bitter they could pucker a rhino [cheap U.S. olives are green or black]). They show that *pseudobrenda* is a valid replacement of what everyone wrongly called *brenda* which is a syn. of *california*. *Mono* is a ssp. like *ampelos* with unh postbasal marks. *Elko* seems to be a syn. of *ampelos* (postmedian bands don't seem narrower either). *Eryngii* is a syn. of *california* (fewer unh ocelli but intermed. to *ampelos* in this, thus invalid).

Chapter 48 *latalinea* is a nice ssp. (*subaridum* on p. 159-160 & 169 is a syn. of *latalinea*, J. Scott **first reviser action**).

Chapter 49. It's totally obvious from the photos (and from my series) that *interrupta* is really *lemberti* X *comstocki*, and thus is a syn. as it is intermediate without any different characters of its own, thus is invalid. More intermediates between "interrupta" and *comstocki* are illustrated, which may have a 2nd gen. like *comstocki*. P. 622 top mistakenly claims *lemberti* is limited to high elev.; actually it ranges quite low on the E edge of the Sierras (where Mike Smith found it at Hot Creek, Davenport found it at Martis Creek, R. Wells and O. Shields & I found it at Fredericksburg, etc.--and Ken Davenport found *perplexa*? resembling *lemberti* flying under 2000' on W side at Tule River Tulare Co.). And note that *Papilio* (New Series) #18 discusses intermediates between *sheridani*X*comstocki*" paradoxa.

Chapter 50. Based on the types, Scott (1998, *Papilio* [N.S.] #11) restricted the *harpalus* TL to "Sierra Nevada Mts., west of Carson City", in other words way west of Carson City, way into the Sierra Nevada where the unspotted ssp. occurs that was formerly called *yosemite*, not at low altitude in Nevada where the spotted green-unh ssp. *idaho* occurs. Scott restricted the TL of *cabelus* to that same way-west locality; they are all synonyms of *harpalus* (*oregonia* evidently properly refers to *idaho*X intergrades from N Calif, despite #11.), while *idaho* is the lowland Great Basin ssp. (By the way, Scott 1998 used the name *H. colorado* for these taxa, but I now think that reports of sympatry in Cypress Hills Alta. etc. were bogus, and the N. Amer. entities should be called *H. comma* again.) Austin is correct in redesignating the TL of *cabelus*, however he meant to say "western slope of the Sierra Nevada", not "west of the Sierra Nevada" (which would be in the central valley such as Sacramento), so I hereby correct Austin's TL to "western slope of Sierra Nevada". C. Gillette couldn't find *mormo* at Davis Creek Park.

Chapter 51. This is a bad paper. The table fails to give the unh ground color! even though that is an important character, while numerous tiny-near-worthless characters are given in exhaustive detail (table 1 is humongous, wow!). There is considerable variation in these bugs, and I don't have really long series, but all these names *magnus*, *mandan*, *mesapano*, & *skada* look poorly-distinguished to me. P. 647 states that *mesapano* has a "poorly defined distribution", and says *skada* has "expanded dark and orange patterning" (how can both dark and orange be expanded?). I was informed that Wyo. ones are large like *magnus* but mine isn't and my largest female is from Alta.; and I was informed that W U.S. pops. are not *skada* or *magnus*. The BC book says that *magnus* ranges north to BC. Cris Guppy states that *magnus* (forest, unh russet brown) flies together with *mandan* (prairie parkland, tan unh with a few russet areas) at 3 sites in Peace R. distr. in BC, and finds different DNA, thus he thinks they are different species. Localities are missing on figs.

Chapter 52 has some problems. Its interpretation of the name *cupreus* and its TL is wrong, as Scott fully explained in *Papilio* (New Series) #12: the *cupreus* type is identical to *lapidicola*, and was collected in the Sierra Nevada, and was not collected in Ore. by Gabb. Thus, *lapidicola* is a syn. of *cupreus*, the TL of *cupreus* is Tioga Pass, and *artemisia* is a different ssp. Some minor mistakes: *lapidicola* does not have two gen. as p. 664 surmises is possible, and *snowi* is brassier colored dorsally.

Alpestris clearly differs from E U.S. *hypophlaeas* (which replaced the name *americana* in Chapter 2), and *alpestris* is similar to *arctic feildeni* but has a narrower orange band. *Bichroma* is a ssp. which is usually quite yellow on unh, and the lavender occupies 1/2 the wing; in *browni* the uns is often quite yellow, and the lavender only occupies the outer 1/3 of unh, whereas in *nivalis* the uns is seldom very yellow and the lavender occupies the outer 1/2 of unh. Actually the unh color varies greatly from bright yellow to yellow-tan in the Cal. Sierras (where I have very bright yellow females) & Colo. & BC, variation that is puzzling; *browni* females actually vary incredibly from brown to tawny on ups. Note the convergence of larger uns spots on both Warner Mts. *warnermontana* & *jacquelineae*. *Dorothea* looks too much like *jacquelineae* to me so is a syn. (the male & female fig. by Dornfeld have smaller spots and orange female upf! & the f is not melanic). Delete last parag. on p. 667, a duplicate from p. 669 right middle. *Micropunctata* looks like a weak ssp. of *gorgon*, as SF Bay Area *gorgon* have uns spots about as small; Davenport notes a blend zone with ssp. *gorgon* in Kern R. Valley, eastern Greenhorn Mts. and upper Kern Can. *Gravenotata* TL is spelled Plainview (near Coal Creek, Jeff. Co. CO), not Plain View. The old specimens of *heteronea* in USNM from Sonoma & Marin Cos. are no doubt violetly because of old age (same color change as in old *icarioides* & *xerces*). *Submaculata* is intermediate *heteronea*X*gravenotata* on uns, but the female ups is said to be more tannish-orange so it is evidently a ssp. (and A. Warren notes that it is different). *Northi* looks like a syn. of *klotsi* (which is best treated as a syn. of *heteronea*) on uns (the syntype on p. 66 has blue upf, but p. 671 says some have brown upf!), but is a ssp. if female upf is usually bluer. P. 671 end, *Yolla* not *Yola*. P. 672 says that *submaculata* and *klotsi* occur only 4 km away in N Calif. and differ in host and larvae (so why are similar *Euphilotes* raised to species status but these are kept on one species?) *Obsolescens* looks like a ssp. if most of them are weakly-spotted, though there are a lot of individuals from S Calif. that look like that (even some from Point Richmond near SF). Davenport notes that a pop. in the

Inner Coast Range in Monterey Co. is consistently chalk white below with very tiny spots, which makes *obsolescens* look obsolete!

This chapter's treatment of *Lycaena xanthoides* is very bad. Actually, *L. dione* is a distinct species (differing grossly in unh orange and small solid black round spots etc.) and *dione* is sympatric & synchronic with *vurali*, while *vurali* and *editha* may yet be ssp. of *L. xanthoides* (all having the same short narrow unh orange band and pale-centered brown unh spots etc.) that differ only in wingspan and relative size of unh spots. Scott fully discussed this in *Papilio* (New Series) #14 p.35-36. *Pseudonexa* is an intermediate between *editha* & *xanthoides* without any distinctive traits of its own, thus is invalid. A published paper (Pratt et al. 1993) that claimed that "*pseudonexa*" is just *L. editha* and not intermediates, was highly flawed, and made the ludicrous claim (p. 188) that Scott mis-measured! the unh spots in his original paper on "*pseudonexa*". Ssp. *editha* and *xanthoides* intergrade in the Dunsmuir area to W slope Ball Mtn., at Mather on W side Sierras, & Silver Can. & Sherwin Summit on E side Sierras, & in SW Ore. *Pseudonexa* is quite similar to *nigromaculata*.

Chapter 53. P. Opler (pers. comm.) told me his 1970 Norden neotype fits the O.D. better than the new neotype, and the ICZN Code seems to preclude the redesignation of a neotype and requires cases of improper designation of neotype to be referred to the ICZN Commission, so this is evidently another inappropriate redesignation, which is invalid. *Tildeni* appears to be a syn. of *fuliginosa* though is said to be darker, and *albolineatum* appears to have larger unh spots than both but those spots are not mentioned in text and only the "whiter" unh is mentioned, which doesn't look much different on the photos, so maybe it is a syn. too. *Maculadistinctum* looks distinct by having larger unf spots, and is said to be darker, but my series has smaller spots and looks like *semiluna* so maybe this isn't distinct either. At any rate, only *semiluna* is certainly a valid ssp., and Warren's Butt. Ore. gives a much better analysis of ssp.: some tiny-stigma pops. range north along W slope of Sierras nearly to S. Wash.; maybe *S. fuliginosa* is a different species from *S. semiluna*.

Chapter 54 is not good. The TL of *multicaudata*, "Mexique" on syntype, should have been restricted. *Grandiosus* is a good ssp. from SE Mex.-Guat. with oranger ups, but *pusillus* TL NE Nev. is a syn. (at best a very weak ssp.) as the variation is described as a cline from S to N, and the male holotype is just an abnormal odd spring form (note that photos in books Butt. BC & Butt. Ore. & Butt. Cascadia are not like holotype). In Colo. I have caught tiny spring adults the size of *Argynnis* (*Speyeria*), and giant Aug. females with orangish uph, variation much greater than the supposed geog. variation. Ken Davenport accepts the ssp. because summer forms of it are much smaller than Ariz. mult. But are those still smaller due to environment, and is size alone enough? I don't accept it in Colo.

Chapter 55, *shastensis* is a syn. of *indra*; S. Spomer states that most specimens from TL are not as narrow-banded as these photos.

Chapter 56, *calcolica* is a syn. as it is intermed. *martini* & *fordi* thus invalid. S. Spomer states that reared *panamintensis* adults do not resemble orig. photos and instead resemble *calcolica*, thus *calcolica* is a syn. of *panamintensis* which is invalid for the same reason. Davenport notes that one of his four males from Westgard Pass looks like *panamintensis*.

Chapter 57, *pygmaeus* is another syn. as S. Spomer told me his adults from there look like *panamintensis* & small size is not consistent (& photos look like *martini*, thus it is a syn. anyway).

Chapter 58 is very annoying. P. 717 claims alpine Colo. *P. phoebus hermodur* do not differ from low alt. *P. p. smintheus*=*sayi*; actually *hermodur* is smaller and much darker, and my reared *hermodur* emerged just as distinctive as in nature (Cris Guppy J. Lep. Soc. 43:148-51 proved that altitudinal melanism is mostly genetic). The dark C Mont. ssp. is nicely corrected to *maximus*. The separation of *phoebus* into four sp. (European *sacerdos*, Asian-Yukon *phoebus*, Yukon-Colo. *smintheus*, Cal. *behrii*) is unsupported splitting. The paper justifies the splitting because of micropyle pattern, but if you use a magnifying glass on fig. 11 *smintheus* you see that it is actually rather similar to *phoebus* on fig. 11, and *sacerdos* is said to have micropyle like *smintheus*. P. 718 cites *Hesperia micropyle* as being a great species-level trait, but Scott (J. Lep. Soc. 29:156-61) showed clinal variation of micropyle between *Hesperia comma colorado* & *H. c. oroplata*. P. 719 says *phoebus* eats *Sedum rosea*, but in Colo. "*smintheus*" ssp. *pseudorotgeri* also eats *S. rosea*, and flies in mesic habitats typical of *phoebus*. *Yukonensis*, *magnus*, *xanthus*, & *sayii* are all syns. of *P. phoebus smintheus*.

Chapter 59 again relies unfortunately on only one character. Thus the wider tip of a small rod on the valva is supposed to prove that *B. titania* of Europe is a distinct species from *B. "chariclea" grandis*, even though multiple characters of wing pattern surely created that remarkable wing pattern similarity, which is extremely-doubtfully due to "convergence" as guessed on p. 728. "Worship the god of genitalia" strikes again (& the fig. of *ranieri* has a sl. wider rod tip). Many people in America and Europe think that it is the arctic *chariclea* that is the distinct species from *grandis/titania*. All the American bugs are *B. titania* to me. (It's interesting that hypocritical dogma considers *B. astarte* & *distincta* conspecific because of similar genitalia when numerous traits of wing shape and pattern differ, yet *magdalena* and *mckinleyensis* are usually treated as separate sp. despite being different in only one trait [red fw flush] that is intermediate in Yukon.)

Chapter 60 is a puzzle because of the differences between their ssp. and those of Troubridge & Wood. The 2nd paragraph is a wishful dream of the actual history in which Scott (1986) was the first person to raise *natazhati* to species rank, based on Troubridge & Philip's research, not Shepard's unknown unpublished work. The O.D. way back in 1920—not Shepard--recognized 2 specimens from Bernard Harbour as *natazhati*. *Bankslandia* is a syn. of *natazhati* (maybe frons & palp differ a bit, but 2 of my 6 have orange—not black—antenna). Why name it *Bankslandia* when it occurs on Victoria I., NOT Banks I.??, that's very confusing (nobody nowadays cares that people 200 years ago used the word *Banksland*).

Why are Coppermine adults in my slides like natazhati & paler than Bernard Harbour adults? Finally, surely tarquinius from Canadian arctic (Baffin I. etc., see Butt Canada fig.) is also conspecific with natazhati, not B. freija (this paper's assignment of it to freija based only on "granitic substrate" drags one-character taxonomy to a new low), thus natazhati is a ssp. of B. tarquinius; a thorough study of CNC tarquinius would be gneiss. (By the way Scott's Pleistocene Fritillary is a perfectly fine appropriate name because it occupied Alaska etc. in the Ice Ages, so its replacement by "Beringia Fritillary" in Butt. Canada is very deplorable, esp. because Beringia implies a Siberian distribution that does not exist—and now Brock & Kaufman have muddled it further with their bad name "Cryptic Fritillary".)

Chapter 61 is a useful study of intergradation of P. pulchella. Inornatus resembles pulchella (unf little black, nudum black) but has ups postmedian spots yellow like camillus, the uns is paler orange tending a little toward camillus, and ups of photos looks a little oranger suggesting influx from montana. Inornatus is an intergrade of those ssp. thus would seem to be invalid with my ssp. definition; however, since the intergrading involves three rather than two parent ssp., maybe my intermediate-is-invalid rule does not apply to this case, and inornatus can be considered valid; so maybe? I should compromise here, and say that popular usage should decide: if you like it, use it. Vallis is a syn. of montana, as it looks like the inornatus stuff that has interbred more with montana thus is oranger; yes it is oranger, but when I caught them long ago I called them intergrades; vallis seems to be just montana. Inornatus, thus a simple intermediate thus invalid. Davenport notes orange P. pulchella from E of Carson Pass, Alpine Co., which resembles inornatus, not montanus or vallis as map predicts. Nudum is black in all of Nev., thus all these names are not related to ssp. owimba. C and N Nev. have intergrade populations showing influx from camillus (also, Shapiro found pulchella-montana intergrades in Plumas-Sierra Cos., and Ken Davenport & I find considerable influx of camillus into the pulchella on E side of the S Sierras). Davenport states that "California is a horrible mix of pulchellus, as S Sierra Nevada has a mix of ssp., and ssp. deltarufa appears to be a blend zone between pulchellus and montanus thus invalid" (though I will not sink deltarufa as a syn. at this time because I wonder if the unf discal cell bar blends at all between those, and because my Mather series & Davenport's Yosemite Butt. paper demonstrate that blacker variable pulchella in the low-altitude Sierras separates San Joaquin Valley deltarufa from high-altitude montana).

Chapter 62. More publicity for Ackery & Vane-Wright's thersippus=strigosus.

Chapter 63. Idaho is a syn. of lahontani (unh not pale enough, and anyway intermed. toward archippus thus invalid). Todd Stout notes that Utah bugs are also variable and intermediate between archippus & lahontani. Colors are much too red on most photos incl. fig. 8; actual specimens show that uns of idaho & archippus are very similar.

Chapter 64. Harbisoni is yet another syn. of coloradensis.

Chapter 65. P. 771 Festuca "ovina" is basically a European sp., as its "vars." are now mostly considered to be species (this mistake has caused idiot government workers to seed alien European F. glauca widely in Colo. mts. etc.). Klamathensis is a syn. of mardon.

Chapter 66. Reeseorum & grandiosa are syns. of lasus; some people who have seen lots of these think they are valid, but to me it looks like there are too many names here. Fulvapalla is valid but terraclivosa is intermediate macswainiXfulvapalla thus invalid. Photos are too red.

Chapter 67. Pratti is a syn. of davenporti named in next chapter (**first reviser action**, J. Scott; davenporti has a far greater range so should be used). Nigrescens is a syn. or weak ssp. of virgulti (described as limited to one colony SW of San Bernardino with dark unh margin, but my virgulti from San Diego Co. & S have darker unh margin than virgulti from L.A. to Santa Barbara, so are those San Diego pops. nigrescens that ranges from TL to Baja?, didn't they notice that difference?).

Chapter 68. Davenporti is the same ssp. as pratti (see above **first reviser action**). Dialeucoides is a syn. of dialeuca; it has the same appearance and host and flight period and mountain habitat as dialeuca, so why can't they be monophyletic? Arenaria is a syn. of virgulti (even though allozymes are said to differ). Obviously there is more than one species of A. "mormo", but the group is a mess. Ken Davenport has found deserti sympatric & synchronic with davenporti, so they must be separate species. He has found tuolumnensis flying with his 2-gen. Kern Can.-Greenhorn Mts. bug (#128b in his 2003 Butt. Kern & Tulare Cos. paper) which based on a series he sent I assign to a new ssp. that looks somewhat near deserti with some virgulti influx (128b is the same as 131c-d-e I think, I have a small series of 131c), so evidently belongs to A. virgulti, thus tuolumnensis is a separate sp. from apparent virgulti. Pratt & Ballmer (J. Lep. Soc. 45:46-57) classified the S Cal. taxa biologically thus (names brought up to date by me): type 1) A. virgulti, multivoltine, young larva hibernates, larva may aestivate, egg duration short, larval duration short, this species evidently includes ssp. deserti, virgulti, & now evidently nigrescens, arenaria, mejicanus [extensive orange upf, uph, unh], duryi [same as mejicanus but unh browner], maxima (deserti & duryi sometimes use Krameria as host, perhaps a clue they are same species); type 2) A. dialeuca, univoltine mostly spring-flying, young-1/2-grown larva hibernates, larval duration much longer, this species now includes davenporti=pratti, mojavelimbus, dialeuca=dialeucoides, evidently peninsularis; type 3) A. mormo, univoltine fall-flying, eggs (or sometimes young larvae) diapause & take longer to hatch, eggs larger than types 1-2, includes langei, pueblo, ssp. mormo=parva=autumnalis=mormonia. Now we add type 4) A. cythera-tuolumnensis, univoltine fall-flying, traits presumably like those of A. mormo, as a separate species because Ken Davenport's 2003 Kern & Tulare Co. paper (Gillette Mus., Colo. State Univ.) reports it sympatric with ssp. mormo along Hwy. 58 in E San Luis Obispo Cos., etc. The Pratt &

Ballmer paper was jumbled and incomplete (data on diapause stage & egg size is very incomplete), but seemed reasonable. But it seems to have been discarded by the new papers. However, the new “1998” papers were probably written in 1988 as was the Euphilotes paper, so are we to believe the J. Lep. Soc. paper?? Those papers left the mormo group (as with Euphilotes) in chaos. There are also problems. A lot of populations are hard to assign to “species” because there aren’t enough wing/host characters to distinguish the ssp. and sp. and there is very skimpy reporting of immatures. Butt. Baja Calif. states that occasional specimens from middle altitudes in W Sierra San Pedro Martir are intermediate *virgulti*X*dialeuca* (a problem that would disappear if that Baja “*virgulti*” was misidentified *peninsularis* and *peninsularis* is a ssp. of *dialeuca*, but they describe 2 gen. for their *virgulti* and say it loses the uph orange southward). *Dialeucoides* is supposed to have just one spring gen., but I have specimens that look like it from Aug. 16-Sep. 24 (some from the same site where they flew June 4-27—if this is what the end of p. 804 describes, then their “univoltine *A. mormo* ssp.” is just the 2nd gen. of *dialeucoides*). *Peninsularis* is said to be distinct from *virgulti* (flying <1 km away with allozymes well-differentiated), and its flight time & host resemble *dialeuca*, but adults look more like *tuolumensis* & *virgulti*; could it be a link between *A. dialeuca* & *A. mormo*?, which are not known to be reprod. isolated, thus *A. dialeuca* may just be a ssp. of *A. mormo* (sympatry & allochrony is NOT a test of reproductive isolation so the two can be just one species).

Chapter 69. *Maritima* & *obscura* are syns. of *polios*. Tawnier uph margin isn’t common on *polios* or *obscura* either; photos don’t match some features of description; fringe is checkered on photos. The B.C. book couldn’t separate BC material from E *polios*.

Chapter 70. *Ricei* is a fantastic new find. *Calneva* is a syn. of it (**first reviser action** noted above, J. Scott)(uns spots are smaller on *calneva* photos on p. 281). I have several *ricei* adults with nearly all uns spots absent.

Chapter 71. *P. saepiolus* is a nest of worms in identifying ssp., like *P. icarioides* (for *P. saepiolus* see also p. 14, 65, 176-179, 197-8). I surveyed this mess and offered my accepted ssp. in *Papilio* (New Series) #12 (here I correct *maculosus*). The ranges of *aehaja* & *rufescens* are wrong. The photos of *gertschi* & *albomontanus* aren’t good (male ups isn’t green and uns is tan-gray not brownish). Ssp. *littoralis* (coastal Del Norte Co. to coastal Ore., a few are similar inland in Siskiyou Co.) is a var. of *saepiolus* (perhaps a weak ssp.) said to have smaller unh spots with white rings. Ssp. *maculosus* (C-E Nev., TL Snake Range) I thought might be a good ssp. with larger spots as described, but as noted in *Papilio* (N.S.) #18, in Univ. Colo. museum I found good series from Snake Range and Monitor Range which have ordinary-sized spots (some even smaller in Snake Range), thus *maculosus* is the same as the bug in NW Colo., and evidently isn’t significantly different from ssp. *saepiolus* based on my few males of *saepiolus* and A. Shapiro’s Field Guide to San Fran. Bay & Sacramento Valley, which shows a female with no blue and says “Sexual dimorphism is extreme in our [Marin/Sonoma/Napa Cos.] populations, but on the far north coast some females have a fair bit of blue above”, thus ssp. *saepiolus* TL Marin Co. evidently has little blue and extends to NW Colo. *Albomontanus* is a syn. of *gertschi* which ranges from Cal. to S Colo.

Chapter 72 is checklist of Calif. butt. It repeats the popular error that *Nastra neamathla* occurs in Cal.-Ariz.-Baja; it actually is limited to SE U.S. (Fla. to SE Tex.), and Butt. Houston & SE Tex. says it intergrades with *julia*, thus they are ssp. *Bairdii* is a ssp. of *machaon*, & *hollandii* is a form not a ssp. It’s *Ascia josephina howarthi*. *C. philodice eriphyle* is actually just a local BC ssp. with some orange on ups of many maybe due to historical introgression with currently-allopatric *C. eurytheme*. *Ariane* & *baroni* are syns. of *nephele*, *carsonensis* is a syn. of *gabbii*, & *walkerensis* is a syn. of *nephele* (partly intermediate toward *gabbii* thus invalid). In Calif. I would use *P. tharos tharos*, *P. cocyta selenis* (mislabelled from Warner Mts. so is not in Calif. anyway), and *P. phaon jalapeno*. *Spey. hesperis* occurs in Cal., not S. *atlantis*. *Xerces* is an obvious ssp. of *lygdamus*. *Fridayi* is the alpine Sierra bug with wings like *lotis* but bigger blue spots & *gnathos* intermediate but nearer *melissa*, which I assigned to *P. atrapraetextus fridayi* in *Papilio* (N.S.) #12. *Hesp. libya lena* doesn’t occur in Calif. or Nev.

Chapter 73 is a checklist of Nev. butt. *P. eunus alinea* doesn’t occur in Nev. does it? Surely *acastus dorothyi* cannot occur in Nev., just variants like it. More *S. hesperis*. *P. tharos* in Nev. might be ssp. *riocolorado*?, or ssp. *tharos*.

Index. Brenda, add p. 589. *Catalina*, Och. sylv., add pp. 313-316. *Editha bayensis* entry and *Euph. ed. bayensis* entry, move p. 210-211 to *enoptes bayensis*. *Editha edithana*, photo of lectotype p. 126. *Flaviventris*, p. 525 not 526. *Phyciodes hondana*, remove 359. *Ivallda*, change 827 to 828. *Karinae*, add p. 399. *Obscurafacies*, add p. 543. *Phyciodes hondana*, remove 359. *Ricei*, *Euphilotes*, move 176 to *ricei*, *Lycaeides*. *Sineocellata*, delete 321, 322. *Tehama*, change 147 to 146. *Thessalia perlulu*, change to *perlula* & remove 359.

FIELD GUIDE TO BUTTERFLIES OF THE SAN FRANCISCO BAY AND SACRAMENTO VALLEY REGIONS. Arthur M. Shapiro, artist Timothy D. Manolis. 2007.

Univ. Calif. Press, Berkeley. 346 p.

At last, a butterfly book full of expert information on hostplants and butterfly biology! Shapiro has studied butterflies in the area for 35 years. Overall, the book text is outstanding in the biological research, and offers numerous interesting insights that can’t be found in most books. He gives us summaries of his research, such as interesting accounts of geographic variation in voltinism and hostplant tolerance of *Papilio zelicaon* across the area. The treatment of local hostplants is excellent, as there is good local info on many species. Shapiro writes about many butterflies that were common 30 years ago on the Calif. lowlands that have now disappeared there (esp. in the Sacramento Valley), such as

Euchloe ausonides & *Lycaena helloides* & *Satyrrium sylvinus* & *Celastrina* & *Glaucoopsyche* ecotype at Suisun Marsh & the *Phyciodes pulchella deltarufa* TL & *Limenitis lorquini* & *Epargyreus* & some *Pyrgus scriptura* sites & *Pholisora catullus*, some of which would seem to Scott to document global warming, though maybe other factors like introduced parasitoids might have wiped out some species. The color plates are disappointing; they are paintings (Manolis' sole contribution to the book), and the butterflies are often positioned annoyingly with curled wings in spread-winged downbeat position, often with forewings spread way forward in a position that butterflies only achieve on a lepidopterist's spreading board (*Nathalis iole* are inexplicably in museum-mounted position)(only Superman with his X-ray vision could freeze butterflies in flight to see them this way), and some such as *mossii* & *muii* & *mylitta* & *orseis* & *juba* uns don't look right, the faunus female is too dark, etc.; butterfly photography has exploded, and great photos are available for every species treated. All the illustrated butterflies seem to be identified correctly. In matters of taxonomy, the book isn't great or is often out-of-date, the worst being the *Callophrys* "dumetorum complex", which gives them numerous names and doesn't bother to choose among them, and gives a perfunctory account of two of them and then misses "pseudodumetorum" which flies in Del Puerto Can. and Trinity Co. Art Shapiro basically does systematics research, but concentrates on environmental causes of seasonal forms, and electrophoresis-then-DNA studies of various populations, transect studies of butterflies and their hosts, etc., while he disdains actually naming any species or subspecies and never completes any taxonomic "revisions" and does not study morphology, thus his work never achieves the kind of robust conclusions that make taxonomically-inclined butterflyers happy. (Though he once did a nice taxonomic study of *Poanes viator* and named a new ssp.; why not do it again?) But no matter: he gives us nice research on the topics he likes, and we gladly accept his gifts. The description of mate-locating behavior in the book is very annoying, as Shapiro casually uses all the anthropomorphic clichés & deer phrases: some are "territorial", or they "fly a beat", or they "lek", or "lek & mate on the hostplant", they are "highly territorial", they are "very active territorially in late afternoon, perching on the host plant", "lek around the tops of pines" (for *Neophasia*, which really patrol the tree tops to seek females), "territorial perchers", etc. In describing mate-locating behavior, Shapiro is out of control! (Scott has recently—in *Papilio* [New Series] #14, and *News of Lepid. Soc.*—proposed four new words to describe mate-locating behavior, which hopefully will become widely used and will remove anthropomorphism and misapplied deer behavior from the description of bug behavior.) The book uses "puddling", another word I don't like as the only butterfly I know that actually lands on puddles is *Apyrothrix araxes*, which lands with wings flat on the water surface! to drink, no other butterflies land on puddles; I used "sip mud" in my book, which is still not great but is better ("suck salty water from moist bare ground" is precise but too long). P. 75 has this gem: "Since subspecies qualify for legal protection, the [Endangered Species] law affords safeguards to things that may not be "real"."

There aren't many mistakes in this book. P. 42 says there aren't any hosts of *Nathalis iole* in Calif., but its selection of *Compositae* hosts is pretty wide in Colo., so there are palatable ones in Calif. P. 42 and 127-8 claims that *Habrodais grunus* adults do not feed and that it has no close relatives in N. Amer., and p. 128 writes that he is mystified at its "odd dance—presumably a lek—around the host deep into the gathering evening gloom". Just read Scott's 1974 paper (*Amer. Midl Nat.* 91:383-90) on the close relative *Hypaurotis crysalus* (yes, that is a 2nd N. Amer. *Theclini* sp.—while "poodiae" will be a separate sp. from *grunus* when poodles grow wings & fly) and you will learn that they use their equally short proboscis to suck mud and the sap oozing from oak twigs, and you will learn that males patrol about the host canopy (fleeking behavior in Scott's new terminology) in late afternoon-dusk using the violet-ultraviolet wing color to find females and court and mate with them. Shapiro notes that *grunus* aestivate for 3.5 months+, fascinating! P. 55 says *H. grunus* perch on the host trees and sally forth to investigate passing insects (raiting behavior in Scott's new terminology, perching behavior in the old), but p. 127 says "they come out in late afternoon to fly around the host plant"; Scott's observations of *grunus* suggest the latter, they fleek around the host to find females as does *Hypaurotis*. (While we're on mate-locating behavior, p. 56 says *Papilio eurymedon* patrols a beat to locate females; yes it does in Colo., but there Scott finds that it patrols only in little clearings among ridge/hilltop trees, back and forth in the small area—flaiting behavior in the new terminology, meaning patrolling in a small area where females fly to mate) P. 52, the world does not appear to butterflies as a mosaic of dots, they have neural network circuits beneath the ommatidia and in the brain which create an interpreted picture of the scene in their brain, just as humans have. P. 56 claims that *Erynnis propretius* patrols, but I found that it rait (perches) in clearings on the sunny edge of a ridgetop/hilltop (sometimes on gulch banks) to await females. P. 31 bottom, the terminal egg inside the female is often fertile, and I have gotten fertilized eggs by dissecting females many times, as has Richard Heitzman who first told me of this technique. P. 60 middle, males can't mate very often (3-5X at most usually), because their spermatophores are large and greatly deplete their energy reserves. P. 74 says that *Pontia occidentalis* looks identical to *P. callidice calora* from Tibet (Scott still calls it *P. callidice occidentalis*). Interesting that *Battus philenor* larvae living in full hot sun become redder! *Papilio indra* males in my experience rait on rocky cliff-like spots just below a hilltop/ridgetop. *Colias eurymedon* might overwinter poorly in alfalfa because larvae do not diapause so die after mowing (but Jack Harry found they do survive 0°F in Utah). P. 135 the May-June "Inland Bramble Hairstreak" records may refer to *pseudodumetorum*, which was missed in the book and is conspecific with *C. sheridanii-viridis*, not *perplexa*. And *viridis* & *perplexa* do co-occur at Marina Beach Dunes near Monterey according to John Emmel's research, and both once flew in San Francisco before *perplexa* was paved over there (SF *perplexa* in CIS coll.). The *Callophrys* name is mistakenly used in parentheses as a subgenus on p. 135-141; the book should have used *Mitoura* "Callophrys" *spinetorum*, etc. P. 145 my

Cupido comyntas from Davis Calif. have orange spots & uns just like E U.S. ones. P. 146 Cupido amyntula evidently does not live inside pods in spring; and Ballmer & Pratt (J. Res. Lepid. 27:50&66) write that the larval tubercles are noneversible in some pops. esp. in S. Cal. where larvae live inside Astragalus pods, while the external feeders have eversible tubercles in N Calif.; and in Colo. the main host is Astragalus flexuosus whose pods are only 3-4 mm wide, probably too small for a mature larva to live inside. P. 149 there never was even a bit of evidence that xerces was a distinct species, its hosts and immatures were the same as lygdamus and the distinctively-white-spotted adults were merely polymorphs, and the lavender color of museum specimens is just due to fading of the blue that fades the same way in Plebejus icarioides! P. 155, synonymize tildeni to bayensis!; taxonomy isn't hard, if it's not right, just change it! Echo would seem to be a ssp. of Celastrina neglecta, which evidently goes across Tex. to Ariz. "cinerea" and Calif. with little change; these multivoltine polyphagous bugs look very similar. Missionensis is just an intermediate pheresX pardalis, therefore is invalid, & the San Bruno Mtn. pop. is also intermediate, but is nearer pardalis. Lots of synonyms in the ssp. of Argynnis (Speyeria) zerene, all 3 "myrtleae" ssp. are synonyms of ssp. hippolyta. P. pulchella pulchella occurs at Suisun Marsh, & ssp. deltarufa in the Sacramento Valley. My Papilio (New Series) #13 gave good characters of female uph median band etc. to distinguish P. orseis. All the Phyciodes hibernate as unfed 4th-stage larvae. P. 181 fails to mention the distinctive extinct C. leanira obsoleta in Marin Co. Variable Checkerspot was the name for the lump of E. chalcedona/anicia, but that stenchospecies has been resplit into bookkeeping species chalcedona & anicia. Why didn't Shapiro name the Sierra Nevada chalcedona a new ssp. with vastly different larva?, that's not hard either. The ssp. taxonomy used for Euphydryas editha is decades outmoded: bayensis is now considered a syn. of editha, luestherae is another syn., and baroni is a very distinct ssp. with shockingly-red uns limited to the vicinity of Mendocino. There are seasonal trends in adult forms of Polygonia satyrus, but the individual variation in each season is almost as great. Zephyrus is a ssp. of Polygonia gracilis, while oreas is not a ssp. of P. progne; it's best to use goats just to make their delicious cheese, and rather than sneer and whine about butterfly names, just change them!, it's not hard to do. The story of Nymphalis californica's migration N & E to higher altitude to produce a generation that then flies to the high Sierras to aestivate seems convincing (Shapiro has published on this bug several times before); so why haven't we found it at high altitude in Colo., where it seems to be a non-migratory native limited to the foothills? (no migration because the Colo. host Ceanothus fendleri is edible all summer?). Aglais milberti males rait on cliffs or clearings just below a hilltop (just like Papilio indra), and sometimes rait on gulch banks. If Vanessa cardui fly SE to NW from the U.S./Mexico deserts to get to Davis, and Scott's Colo. cardui fly to NNE all day from the same deserts, why do they not interbreed in winter in NW Mex. and lose their orientation? Our U.S. V. atalanta rubria differs from European atalanta, therefore ours is native. Interesting that after months of adult estivation, Coenonympha tullia californica lay eggs and the larvae fast until the grasses grow. Why not use Scott's "Scrub Wood Nymph" name for C. sthenele?; hypoleuca is similar to thus a syn. of ssp. sthenele to Scott. Ralph Wells found O. nevadensis in Eldorado Co. P. 223 the left & right Erynnis valvae work differently during mating, and the species fit in 3 groups: left-handed, right-handed, and ambidextrous (see Papilio [N.S.] #14). Salix records for E. persius in E U.S. are based on E. icelus. Pyrgus "albescens" was mapped NE of Stockton by Burns J. Lepid. Soc. 54:58. Maybe albescens isn't a distinct species: Shapiro notes that higher Sierra Nevada Pyrgus "communis" have the "molecular data" of P. "albescens"; & in Latin Amer. the costal-fold-less "adepta" replaces albescens at higher altitude yet some individuals have the costal fold & valva of the other; & the way albescens is neatly replacing communis in Fla. suggests that a dominant-recessive gene for valva shape is linked to temperature tolerance and they are conspecific and one gene is just replacing another; rearing studies are needed. Maybe Colo. communis differ too, they do not "perch well off the ground, up to waist height" as this book describes in Calif., instead they mostly flait (patrol a small site) in low weedy spots and land low to the ground. Hesperia juba does NOT eat bunchgrasses: in Colo. Scott's more than 100 host records (Papilio [N.S.] #6) show it eats a low mat grass Bouteloua gracilis, plus several winter annual grasses Poa secunda & Bromus tectorum which are green from fall to spring. And juba adults have two gen. & do NOT overwinter, as Scott in Colo. (Papilio [New Series] #6 p. 109, and Lepid. News 1982#1 p. 5) found that Sept. females ovip. on winter annual grasses etc. and the larvae have fast development so they mature in fall and spring in time for a new May generation; and A. Warren (Butt. Ore., p. 36-37) also disproved adult overwintering & actually saw those spring generation adults migrate just above ground level upslope to higher altitude! (Shapiro claimed that adults overwinter because spring adults were found with Chrysothamnus pollen from fall flowers, but that logic is wrong—the entire science of palynology is based on pollen wafting for miles and settling in peat and fossil deposits and serving as good indicators of the age of those deposits, and if pollen can survive for millions of years that way, some surely survives the winter on the bushes and attaches to a spring juba landing thereon). (In general, teeny things like pollen, bacteria, & fungus spores spread everywhere.) P. 240 H. columbia is not greenish. Mysteries remain in H. lindseyi: I found it common at a trashy weedy spot where winter annual grasses looked like the only possible host. Get rid of the name "Dogstar" Skipper!, yck! Scott has numerous Ochloides yuma "sacramentorum", which does not differ one iota from ssp. yuma and is a worthless synonym. (Some "conservationist" morons in Kern Co. removed all the Phragmites from one site thinking it was the invasive weed rather than the native strain, and exterminated the yuma pop. there.) The ssp. of Euphyes vestris are very dubious. Amb. vialis eats 5+ sp. of hay (taller wide-leaved) grasses (Scott [N.S.] #6). P. 270 Oeneis & Neominois pupate "underground" in loose soil or litter, and the turf skippers such as Hylephila & Atalopedes & Polites peckius-sonora etc. evidently pupate in the larval nest in loose soil/litter/base of grass clumps. P. 274, Shapiro is

optimistic that amateurs will take over the study of natural history, but Scott sees collectors disappearing, and butterfly watchers failing to provide usable information on hostplants or any other scientific info. It takes a lot of butterfly collecting and endless hours of staring at rigid specimens to develop identification skills and find suitable localities where the bugs are common enough to get results in observing natural history, and it takes a lot of plant collecting and microscope study to learn the plants, in order to do that research, which butterfly watchers just are not able or willing to do. But it won't matter anyway, as our bugs are going to fry with global warming, and when the oil runs out in two decades, we will starve.

BUTTERFLIES OF CANADA, by R. Layberry, P. Hall, and J. Donald Lafontaine.

1998. University of Toronto Press, Toronto, Ontario. 280 p.

This is a good book, with mostly-accurate text, nice color plates and maps, a gazeteer of localities, a professional discussion of generic controversies, etc.

There are some misidentified photos on the fine color plates: Plate 1 fig. 8 is brizo male, pl. 5f4 is hybridized with canadensis (the unf submarginal band is a stripe instead of spots), 7f24 is the spring form vernalis=ariadne, 8f1-2 are from Colo. not Alta., 10f5-6 are *L. rubidus rubidus*, 10f13 is *L. e. michiganensis*, 10f31 *S. sylvinus nootka*, 10f53-54-55-56 are *C. gryneus rosneri*, 11f3 ups is same mossii female ups as 2, 13f16 is *snyderi*, 11f23 evidently *Cupido amyntula maritima*, 11f24 *C. a. amyntula*, 11f33-35 *Plebejus atrap. atrapraetextus*, 11f34 *P. atrap. alaskensis*, 11f40-41 *E. glaucon* near *glaucon*, 12f14 *S. callippe chilcotinensis*, 13f2 *S. zerene picta*, 13f6 *S. hesperis dennisi*, 13f8 *S. h. minor*, 13f9 *S. h. rhodope*, 13f13 is *S. hesperis dennisi* female, 13f14 *S. zerene picta*, 13f16 *S. coronis simaetha*, 14f15-16 *B. tarquinius natazhati*, 15f5 *C. whitneyi altalus*, 15f11 *P. cocyta cocyta*, 15f12 *P. cocyta selenis*, 15f13-14 *P. pulchella owimba*, 15f23-24 *E. a. helvia*, 16-8 *P. faunus* ssp. near female *f. silvius*, 17f10 *L. weid. weidemeyeriiXrubrofasciata* (oranger unh), 18f12-13 later named *C. sthenele sineocellata*, 18f14-15 *C. oetus oetus*, 18f21 is *magdalenaXmckinleyensis*, 19f119 was later made holotype of *O. alberta ojobwe*, 20f8 *O. polixenes luteus*, 20f9 *O. polix. subhyalina*, 20f10 *O. p. yukonensis*, 22f11 may be *P. pallida barnesi* as it has white patches on top, 28f1 is *P. alupini lutzi*, 28f5 is *newcomeri* (*neoperplexa* isn't in Canada), 29f2 *N. menapia tau*, 32f7 is *B. titania grandis*, 32f8 *O. polix. luteus*.

There are no life zones used in the book. Southampton Is. in Hudson Bay looks like a distribution dot on most maps but isn't. P. 12, Queen Charlotte Is. had 100 m lower sea level during glacial advances and had butterflies. *Boloria alberta* & *B. improba* nunatak probably survived on Alta. nunataks also. P. 13, alvar not defined in glossary (=a place with bedrock exposed at surface, often limestone) P. 22, upright egg means that the line from micropyle to middle of egg base is perpendicular to attachment substrate (parallel in flat egg).

Detailed notes (the review for Butt. BC adds other notes): *Erynnis pacuvius* was reported on *Ceanothus cordulatus* for ssp. *lilius*, not in Canada. Yukon-W BC *afraunus* seem dubious. Poplar & willows are errors for *E. persius*. P. 47, it's actually not hard to find *Hesperinae* larval nests in grasses because they bunch the leaves and there is often a "loop" of a grass blade whose tip is silked into the nest but grows faster from the base than the other nest leaves so loops upward next to the nest (*Satyrinae* larvae are hard to find). *Hesperia juba* adults do not hibernate. I wrote once that *Hesperia colorado* and *H. comma* are different species based on Canada research, but now it seems that *Hesperia comma* and *colorado* are conspecific, as Jon Shepard has altitudinal series that intergrade between *H. comma idaho*=*harpalus* and high-mountain *H. comma manitoba* in B.C., N. Kondla says the *manitoba* and *assiniboia* are nowhere sympatric in Alta., Paul Klassen says *borealis* and *assiniboia* blend at The Pas Man., and the "colorado" and *manitoba* from Cypress Hills on the Butt. Canada maps are errors and are *assiniboia* (*colorado* from Cypress Hills of Sask. on the Butt. Canada map is an obvious misidentification of *assiniboia*). There are three kinds *manitoba* *assiniboia* & *colorado*, but they are not reprod. isolated (see *Papilio* [N.S.] #12). *H. nevada* has just one gen. (even in lab early stages take 100 days). *Poa pratensis* is an error for *Polites sabuleti*. *Polites draco* immatures descr. in *Papilio* (N.S.) #6. *Polites origenes* hibernates as 3rd-4th-stage larva. *Polites mystic*, host is *Echinochloa crus-galli* not repens. *Festuca idahoensis* is dubious for *Polites sonora*. *Euphyes vestris*: *Tridens flavus* actually came from A. Shapiro's Butt. of Delaware Valley. *Amblyscirtes osleri* eats *Bouteloua curtipendula* (*B. gracilis* was a "probably" guess error by Opler & Malikel). Norbert Kondla told me the dot from Copper Mtn. in W-C BC on *Parnassius clodius* and *P. icarioides* etc. maps is really Copper Mtn. in the Kootenay Mts., so those dots should be in SE B.C. not in WC B.C. *Smintheus* is a ssp. of *P. phoebus* (micropyle very similar with magnifying glass, *sacerdos* has *smitheus* micropyle, upf submarginal band wide in *pseudorotgeri*, etc.). *P. brevicauda* may be a ssp. of *P. machaon*. *Dodi* occurs on S prairies & is a syn. of *brucei*; ssp. *bairdii* is NOT in Canada or N U.S. (similar specimens must be *kahli* or *nitra*). P. 86, all *Papilioninae* hover while nectaring. *Rutulus* & *canadensis* are ssp. of *P. glaucus*. *P. multicaudata* does not eat *Salix* & *Populus*. The *Pontia protodice* dots from W Alta.-N BC are dubious according to N. Kondla. *Pieris "napi"* in Colo. are quite variable, on uns at least, and Adam Porter found that there is very little difference between *bryoniae* & *napi* in Europe where people once insisted they were very different species, so I am mystified at the splitting within "napi". *Pieris angelika* is evidently a ssp. of *Pieris hulda*. The Yukon line isn't a natural boundary, so the *angelika/marginalis* dots look artificial there. Geiger & Shapiro's paper was tentative with small sample sizes, one of their *angelika* pops. was treated as *meckyae* by Eitschberger, etc. There are *Euchloe hyantis/lotta* intermediates on the Kern Plateau in Calif. *Papilio* (N.S.) #18 fixes *Anthocharis "sara"*. *Colias eurytheme*, the northern dots from BC to Ont. are dubious. The *Colias occidentalis/alexandra/christina* treatments are an incorrect mess; see *Papilio* (N.S.) #12; *Colias christina/krauthi* is the

same species as *C. occidentalis*; Butt. Alta. has the correct map for alexandra range. Jack Harry has since reared *C. meadii johanseni* larvae/pupa (in Taxonomic Report, TILS), which are identical to Scott's Colo. meadii. The treatment of *Colias tyche* is very good. *Colias pelidne*, the NW BC dot is error (for *Colias interior* also), and probably all the NW Canada dots are misidentifications. Scott's book had an X for *Phoebis philea* in N.B., not N.S. P. 122, *Lycena xanthoides* has pale-centered unh spots. Canada has *L. rubidus rubidus*. *L. heteronea*, move W-C BC dot to Copper Mtn. in SE BC. *L. epixanthe* is not on pl. 14. The dorcas map is a mixture of *L. dorcas* & *L. florus*; *L. florus arcticus* & *L. f. dospassosi* are ssp. *L. mariposa*, *Polygonum* is error. *S. acadica* "montanensis" from Waterton Lakes are *S. sylvinus nootka*, which occupies the BC range also. *S. californica brashor* was later named in BC, not putnami. *S. liparops aliparops* evidently occurs only in S Alta. *Okanaganum* is a syn. of *Satyrium saepium*. Canada evidently has *Callophrys sheridanii newcomeri*. *C. gryneus rosneri/barryi* writeups are wrong, the Alexis Crk. & Williams Lake dots are barryi (lots of Juniperus), the Vancouver I. "barryi" dots are misid. rosneri, the plate photos don't look properly identified, and A. Warren's Butt. Ore. corrects barryi TL and says rosneri =barryi =plicataria =byrnei. Call. lanoraicensis is not in Conn. R. Robbins states that *C. niphon* & *eryphon* do hybridize. *Strymon melinus*, only ssp. melinus & atrofasciata are valid. Alce is a syn. of *Echinargus isola*, which I have never seen migrate. *Cupido amyntula maritima* evidently occurs in most of Canada, and ssp. amyntula in BC, though there are forms like albrighti (grayer uns) & immaculata in BC according to Kondla. *Nigrescens* would seem to belong to *Celastrina neglecta* (with echo), while *Celastrina argentata* evidently includes the Cherry Gall feeder (now known as *serotina*) & *humulus*. *E. battoides* is evidently *E. glaucon* near-glaucon assoc. with *Eriog. heracleoides* based on Warren's Butt. Ore. *Glaucopsyche piasus* larvae eat only flower buds. *Glaucopsyche lyg. kournakovi* is the alpine Alta.-Yukon bug with silvery ups, and ssp. *afra* occurs in the plains with tiny unh dots. *Lycaeides idas* is a mess of 3 sp. (see *Papilio* N.S.#12). *Samuelis* is a separate sp. or ssp. of *P. atrapraetextus*. Move the W-C BC Copper Mtn. dot of *Plebejus icarioides* to SE BC. *Plebejus alupini lutzi* occurs in Canada, with ssp. *spangelatus* in alpine Alta. Argynnids were lumped into *Heliconiini* by Scott & David M. Wright 1990 (Chapter 2 Butterfly Phylogeny & Fossils, in Otakar Kudrna's Butt. Europe). *Plantago* is a dubious host of *Euptoieta*. W.H. Edwards described *Argynnis* (*Speyeria*) *hesperis* immatures. The *Speyeria* & *Boloria* were discussed in the Butt. BC review. Labrador *B. eunomia* lack silver spots, and Churchill *eunomia* differ on uns too. Why is *B. frigga* absent in most of BC, did glaciers scour the willow fens? Early stages are known for *B. improba* in Colo. & Wyo. Based on appearance & range, *natazhati* is a ssp. of *B. tarquinius*. Beringian Fritillary is ridiculous, as it is not in Siberia; Scott's Pleistocene Fritillary should be used. Evidently *B. t. natazhati* occurs on Mt. Natazhah, Mentasta Mts. Alaska, Montana Mtn. Yukon, Coppermine NWT, & Victoria I. NWT, while *B. t. nabokovi* irrationally occurs on Sentinel & Stone Ranges BC, White Mts. Yukon, Mackenzie Mts. NWT, & Bernard Harbour NWT. The *natazhati* is said to ovip. & assoc. with *Dryas integrifolia*. NE BC *B. alberta* dots are errors according to N. Kondla. Half-grown larvae of *Chlosyne gorgone* hibernate. *Sterope* doesn't occur in BC and is same sp. as *acastus*. *C. palla* doesn't eat *Castilleja*. *C. "damoetas"* is *C. whitneyi altalus*. Canada has *Phyciodes tharos tharos* in S Ont., *P. t. orantain* in Alta.-Man., two-gen. *P. diminutor* in S Ont., single gen. *P. cocyta cocyta* in Nfld.-NS, *P. cocyta selenis* in most of Canada (which does not have pinkish spines). Canada has *P. pulchella* (best identified by the orange [no black] subapical costal unf patch) ssp. *owimba* & ssp. *tutchone*; *camillus* stops 300 miles from Canada. *P. pallida barnesi* has blackish larvae. The *Phyciodes mylitta* larva photo has white patches near top, which is odd. *E. anicia* are errors in Man., the "chalcedona" treatment is a mess; *E. bernadetta* occurs in Cypress Hills along with *E. anicia*. The Klotassin R. Yukon is surely an error for *Polygonia satyrus* & *oreas*. Narrow-banded "oberfoelli" are just individual variants in *Limenitis weidemeyerii*. P. 214, in contrast, *Lethe eurydice fumosus* eats the narrow-leaf *Carex aquatilis* in S Minn. and narrow-leaf *C. emoryi* in Colo. *Nipisiquit* is a ssp. of *C. tullia* (they fly within a few feet, but the nipisiquit flight period is different than immigrant *inornata* so they can't interbreed, and they look similar enough that hybrids would not be detectable anyway, as in adult appearance it isn't even a valid ssp.), despite distinctive mtDNA (mtDNA fails here again, as it failed in *Phyciodes* & *Polygonia progne/P. oreas* & *Papilio glaucus appalachiensis*, etc.); it has different larval overwintering stage. Move the *Cercyonis pegala* dot from Copper Mtn. in W-C BC to SE BC; *ino* & *boopis* are syns. of *nephele*. The 3 *C. sthenele* dots in SE BC are errors. Canada has *C. oetus oetus*=*phocus* (*silvestris* is Calif. ssp. of *C. sthenele*). Yukon bugs are intermediate *Erebia magdalena*X*mckinleyensis*. I use the names *E. stubbendorffii pawloskii*, *E. s. canadensis*, *E. dabanensis youngi*, & *E. kozhantshikovi lafontainei*. *E. episodea* has no ssp. *Neominois* has only 1 gen. *Papilio* (N.S.) #12 divided *O. chryxus* into 2 sp. *O. uhleri* from km154 & mile 80-97 of Dempster Hwy. Yukon are dark brown (nahanni?). SE Man. *O. alberta* were later named ssp. *ojibwe*; the E BC dot is error. P. 28 states that Kurentzov's *Oeneis* names apply in N.A., & I agree that *O. alpina* is in N.A. as the O.D. illustrates holotype male and valva which is like *excubitor*. But N. Kondla has photos of both syntypes of *rosovi* and neither look like *O. philipi*, and he states that *beringianus* does not apply in N.A. either. Kondla found major mistakes in the range of *yukonensis*, and Alta. *O. polixenes* has males near *brucei* & females near *subhyalina*. NE BC *Danaus plexippus* are errors. P. 258 *patrolling* is not flying in a specific territory! Bibliography lacks Ferris (1977)(p. 127) & (1987)(p. 142) & K. Johnson (1976)(p. 140).

BUTTERFLIES OF BRITISH COLUMBIA, Crispin S. Guppy & Jon H. Shepard.

2001. University of British Columbia Press, Vancouver, BC. 415 p.

This book is beautiful, with good photos of adults, nice-looking maps, and lots of great photos of eggs larvae and pupae (often borrowed from California or Washington or elsewhere, but still great). It has lots of good information. The authors tried to write a definitive treatise, and named various subspecies to place names on more of the diversity of BC butterflies. This attempt is very commendable, compared to some skimpy Canadian province & U.S. state books that merely provide a picture and a pathetically-starved writeup for each species. The book has lessons about butterfly conservation too. One lesson is that clear-cut logging opens up habitat to benefit various butterflies (idas, icarioides, etc.), land clearing creates grass habitat for others (*C. tullia*), and a christmas tree farm creates habitat for others (*E. editha*, which went extinct when the tree farm closed and weeds invaded). These lessons are contrary to the tree-hugging environmental nuts who plague most conservation organizations today and like to scare the public with clear-cutting scares really designed to raise money from ignorant idiots (and reminds us of Ken Davenport's discovery that conservation workers removed all the *Phragmites communis* at a Kern Co. site thinking it was an introduced weed, resulting in the extermination of that *Ochloides yuma* colony). Another case blames pesticide spraying for extirpating *Limenitis archippus*. Various species (*B. bellona*, *S. atlantis*) have odd ranges that miss SE BC, which is interesting, although N. Kondla suggests that *B. freija*, with a similar range, would have more records there if it flew later into summer collecting season. And C. Guppy notes that the book lumped a lot of *S. atlantis* records with *hesperis*, and Guppy can now distinguish them easily and finds that *S. atlantis* is much more common in C and N BC than the maps indicate. The book wastes a lot of space trying to determine the origin of scientific names (such as the dubious attempt for *Limenitis*); this etymology annoys me because it is irrelevant to butterfly behavior/ecology/taxonomy etc., and because so many scientific names are grossly inappropriate or misleading.

The book does have a large number of mistakes however, arising from a large fauna with not enough lepidopterists or time to study them. Too often, the subspecies are listed and mapped but no distinguishing features are described (for those species written by Jon Shepard), leading readers to question their validity or believe that they were determined by locality rather than actual characteristics. Hostplant information is quite weak, with hundreds of hosts missed that were published by Scott in particular (in *Papilio* [New Series] #6).

Below are detailed comments and corrections I have noticed, with numerous others added by Norbert Kondla. P. 13, the records of *V. atalanta* from Queen Charlotte Is. and Atlin are in both the Canada book and the BC book, so are correct according to Guppy (pers. comm.), who states that it is a migrant to BC, far northward in some years.

P. 89 the right *Erynnis valva* is also useful for identification, nearly all BC *Erynnis* can be identified from wing pattern alone; and the tibial tuft character is useful (missing on some species such as *pacuvius*). P. 92 *E. afranius* records are surprising, the two specimens from New Aiyansh were part of a local collection acquired by Shepard, so should be from that locale. Mislabeling does occur (A. B. Klots' collection of *S. zerene* and *S. hydaspe* in N New Mex. was mislabeled for instance, the latter even named "conquista"), and misidentifications of the *valva* occur. However C. Guppy notes that those "afranius" records were definitely collected from that site, but is doubtful they are really *afranius* and may be an undescribed sp. in the *afranius-lucilius* group (I note that there are strange records of genitalic "lucilius" from N Man. & "afranius" from Yukon etc., and a report of *afranius*-genitalia bug from Pink Mtn. BC, hinting at a boreal Canadian creature now unnamed?). *E. persius* records for *Salix* & *Populus* are errors for *E. icelus*. Scott 1992 *Papilio* #6 found that *Colo. persius* eat several legume genera. The white uph base spots described on *P. centaureae* are actually missing on this species and on the photos. *P. ruralis* is easily identified on upf. *Carterocephalus* is in subfamily Heteropterinae; most of the ssp. of *palaemon* are dubious to me because individual variation is great, although N. Kondla suggests that the the boreal forest/aspen parkland *mandan* has a tan unh with a few russet areas, and the cordilleran *magnus* has a russet-brown unh as in photo. The ups of *O. garita* is not greenish, and Scott in *Papilio* #6 gave half-grown larva as overwintering stage. *H. juba* adults do not overwinter: Scott Pap. #6 proved that *H. juba* has two generations, fall females lay eggs that hatch and larvae feed and grow very fast compared to other *Hesperia* (allowing two generations), no adults occur in March-M May when real overwintering adult butterflies fly, and obviously Shapiro's May adult sat on an inflorescence that survived the winter and picked up stray pollen (and A. Warren recently observed *juba* migrate to higher altitude). The female "juba" photo is *H. comma idaho*. The live male photo of *H. comma "oregonia"* is *hulbirti* as the book defines and maps it. *Oregonia* is an intergrade of *idaho*X*hulbirti*Xetc. in N Calif., so *hulbirti* is the correct name for *oregonia* in BC, and *idaho* is the correct name for BC "harpalus". *H. nevada* adults are easy to identify (though Guppy says some BC *H. comma* are similar), and eggs do not turn pale green. *P. peckius* half-grown larvae hibernate (*Papilio* #6). In *P. sabuleti*, *Eragrostis trichodes* is a misidentification error (see *Papilio* #6). *Papilio* #6 gave many hosts for *P. draco*. The photos of *P. mystic* do not look like an undescribed ssp. *Atalopedes unh* is not tweedy green. Pap. #6 described immatures and numerous hosts of *O. sylvanoides*, and the "phenotypic response" theory of ssp. is wrong. *Papilio* #6 gave numerous hosts for *E. vestris* (*Cyperus* is lab only) and *A. vialis*.

The *P. clodius* ssp. are worthless to me, though Guppy says he can distinguish them; the gray chevrons are present on *claudianus* photos, and all the pops. from W N.A. in Scott's coll. have weak chevrons except a Stevens Pass pop. E of Seattle with strong chevrons, which this book labels as *claudianus* characterized by weak chevrons! *P. phoebus smintheus* seldom oviposits on the host (Shepard observed *phoebus* in Yukon ovip. on the host, but this may be only occasional; Scott

found that they are laid about 10 cm away on average). *Smintheus* belongs to *P. phoebus* (Shepard & Manley's 1998 fig. 11 *smintheus* micropyle looks a lot like *phoebus* micropyle when you use a magnifying glass!, and other characters used to justify the split such as the moist habitat and *Sedum rosea* host of *phoebus* also occur in SW Colo. *pseudorotgeri*; and Evgeny Zakharov has shown that *bremeri* in Asia has the same mtDNA as *phoebus* and intergrades with it, and is therefore another ssp. of *phoebus*, along with *sternitzkyi* and *behrii*. (Also, the number of micropyle spines differ in Colo. *H. comma colorado*, but intergrades clinally to *H. c. oroplata*, J. Lep. Soc. 29:156-161). About its ssp., *yukonensis* females have large red spots approaching those of *phoebus*; and *magnus* & *olympiannus* are surely syn. of *smintheus*. The male *phoebus* photo has rather small spots, contrary to text. *Bairdii* is a ssp. of *P. machaon* as shown by F. Sperling, and *dodi* is a syn. of *brucei* as it has no distinguishing differences; actually *hudsonianus* hybridizes with *zelicaon* only in some areas, and *pikei* is intermediate between *hudsonianus* and *oregonia* (the eyespots like *hudsonianus*). *P. zelicaon* form *nitra* should have been mentioned, although Guppy states that *nitra* does not occur in BC; *zelicaon* could have two gen. near the coast but seldom three (Guppy told me a third gen. is partial and overlaps with late 2nd-gen. adults). *P. indra* larvae take much longer than 18 days in nature. *P. glaucus* is conspecific with *canadensis*, *rutulus*, and *alexiares*, recent splitting here violates the biological species concept. *P. g. canadensis* intergrades with *rutulus* in SE BC (Scott & Shepard Pan-Pac. Ent. 52:23-28) where numerous intermediates abound, and the map shows intermediates. *P. g. alexiares* wings and valva look like *rutulus*, yet females are often black like *glaucus*, its hybrids with *glaucus* had high fertility and 1:1 sex ratios (Scriber & Lederhouse JRL 27:222-232), and its mitochondria are very similar to *glaucus*. Also, *arcticus* is a *rutulus*-like form in Alaska etc., & *canadensis* intergrades with *glaucus*. *Pusillus* is a syn. of *multicaudata*. *P. eurymedon* males flait (patrol a small area among trees on ridgetops to await females), in Colorado also.

Tau is a good ssp. near the TL that has a larger black fw patch (not a less-red unh); most of the map has ssp. *menapia*. *Beckerii* surely has only 2 or possibly 3 gen. in BC, not 4, though Guppy assures me that a few adults are the 4th gen. Some *P. sisymbrii elivata* females are yellow, so I wonder if *flavitincta* is a weak ssp. Hybridization expts. with Siberian *P. callidice* (rather than with the one or two European adults as Shapiro did), which resemble *nelsoni*, would probably show compatibility with *occidentalis*. The *Pieris napi* group needs lots more work, as everyone agrees. It seems to me to be highly muddled in the book, so I offer some possibilities below, while admitting that these possibilities are speculative as I have little knowledge of these bugs: Guppy and Kondla claim to find *marginalis* and *oleracea* sympatric in numerous locations. But based on the maps, *P. oleracea* and *marginalis* could be the same species (see the isolated central BC cluster of *oleracea* that fits nicely into the missing area of the *marginalis* map), and the book's *marginalis* species looks phony to me because *guppyi* and *tremblayi* are surely the same species as *angelika* based on very similar wing pattern (the photos all have narrowed unh veins near margin, etc.), and the graph beside map shows that *guppyi/tremblayi* have only one generation (Guppy told me that he has collected 2nd gen. but it is obscured because of its late season when few collectors are about and by the wide altitude gradient). But the 4 photos of "angelika" are *tremblayi* according to N. Kondla, and C. Guppy notes that *angelika* is closer to *oleracea* than to *hulda*, so maybe *angelika* is a ssp. of *P. marginalis*. The wing pattern melanism sequence is: unspotted summer forms > *oleracea* > *marginalis* > *reicheli* > *tremblayi* > *guppyi*. *P. oleracea* may have recently dispersed along Alcan Hwy. to Yukon just as *P. rapae* did (though Guppy told me that *oleracea* W of Rockies are an unnamed ssp. hence cannot have dispersed recently). The graph shows mostly one gen. (sometimes two) for *oleracea*, whereas *angelika* has one gen., so *angelika* might occur at higher altitude than *oleracea*?, but Guppy told me they occur at the same altitudes. The immature traits seem weak, for instance *tremblayi* larva has spiracles in cream band & magnifier shows cream around spiracles (Guppy told me that cream is absent around spiracles and that the "cream band" is not pigmentation, it is shiny reflection from hairs, as noted in text for *reicheli* larvae; though I doubt that hairs aimed toward the camera would reflect at all--the pale lateral line next to black spiracles of *tremblayi* larva photo surely is not tracheae?). *P. angelika* is said to have lower pupa projections, but that was perhaps based on pupae that hatched into *tremblayi*? The adult traits are so variable as to be questionable also, and the phenotype changes drastically in seasonal forms. At any rate, I have no answers here, just these speculations, but the scheme presented here and by Eitschberger etc. seems to have many questions so a lot more work is needed. Chew & Watt (Biol. J. Linn. Soc. 88:413-35) did mtDNA of a few arctic specimens, but one wonders about their identifications, particularly the Racing River specimen in NE BC. (More: the Alaskan *passosi* according to Guppy is a mixture of *shapiro* & *oleracea* syntypes thus is currently useless; the *pseudobryoniae* TL was stated to be Finnmark Scandinavia in Miller/Brown while Pelham's Catalogue lists it as *Nulato* Alaska.) Looks like *Pieris rapae* spread along the Alaska hwy. *Anthocharinae* is not a valid subfamily (Scott & Wright, vol. 2 Butterflies of Europe, etc.). (American *ausonides* looks rather like European *E. simplonia*, which until recently was called *ausonia*, because both names have been transposed in Europe). Pavel Gorbunov's new Russia book suggests that *E. naina* is a ssp. of *E. ausonia*. If *ogilvia* really has darker uns, perhaps it is conspecific with *naina*, as Gorbunov reports great variation in Siberia in *E. ausonia naina*; *ogilvia* and *naina* are sympatric in Yukon, where they may be separate species, or forms? (a few adults identical to *naina* occur in high-altitude Colo. *ausonides*). *E. naina* photos do not show rounded fw. *E. hyantis lotta* is suggested by intermediates reported from Kern Plateau Calif. The book distinguishes "A. sara" from "A. stella" well, though the book states that females become yellower inland, and the bar of *stella* becomes like *sara* westward, implying that they are the same species. Guppy told me that "stella and sara both fly on the west side of the Coast Range, near Vancouver. They maintain separate phenotypes, and in the Skagit valley apparently use different habitats." The real

ssp. *stella* has yellow-tinted males and is limited to the Sierra Nevada of Calif., so the “*stella*” of this book is a related ssp. *columbia*. The black marginal spots and yellow veins described do not match the *Anthocharis* photos. More work needed. (See *Papilio* [New Series] #18 for some clarification of *Anthocharis*.) *C. philodice* is very often paler yellow on ups, though the photos look too pale (Guppy states that many eriphyle are this pale); *Astragalus* “*carodurpus*” is misspelled (for *caryocarpus*?) as is *leucanthus*. Note that eriphyle (based on photos sent by Guppy) is a good ssp. limited to BC which has a lot of orangish-tinted adults, which I think must be due to introgression with *eurytheme* in some past eon such as the altithermal period. Colo. *C. eurytheme* adults survive fall frosts quite well, and I have never seen real migrating *eurytheme* adults, so I assume it is native in Colo. as J. Harry finds in Utah. *C. eurytheme* surely overwinter and do not migrate in BC (though Guppy did see a migration in BC), as is true of *eurytheme* and *philodice* in Colo. Guppy states that it cannot be native in BC because the phenology histogram shows very few spring records, but could the few records be caused merely by its scarcity in the province (“their populations never get large”)? (Colo. *eurytheme* and *philodice* are not common in spring). S. Ae (1958) did write that his larvae on the Yale roof did not diapause in fall and died in the winter while *philodice* diapaused 3rd stage (maybe because the diapause inducing mechanism was not yet adapted to that latitude?). But why don’t I see migrating adults in Colo.?? *C. alexandra* could still have been named after Princess Alexandra, as a princess is not exactly obscure; and W. H. Edwards lived in West Virginia so he may have liked the south. Queen Alexandra’s Sulfur was first used in 1956 Colorado Butterflies. But true *C. alexandra* actually does not occur in B.C. *Papilio* (New Series) #12 updates the treatments of *Colias* “*alexandra*” & *C. christina*. Recent collections in Grant Co. Oregon, by C. Ferris and A. Warren etc., and the MS thesis of Clyde Gillette that concluded *wasatchia* was a ssp. of *occidentalis*, have shown that *C. occidentalis occidentalis* intergrades with *C. occidentalis wasatchia*=*pseudochristina*, so are one species; “*astraea*” [now called *sacajawea*] and *christina* and *krauthi* are obviously the same species as *wasatchia*, so all these are ssp. of *occidentalis*. The holotype *pseudocolumbiensis* photo in book is oranger than the actual specimen which has only a slight orangish flush. The female *pseudocolumbiensis* in the book is *Colias edwardsii* according to N. Kondla, who currently considers them to both occur in S BC. The book maps the range of *pseudochristina* as a ssp. of *occidentalis* (and misses the Ida. range of *pseudochristina*), which is correct, but was not correct for the limited concept of *occidentalis* that the book espoused, because *pseudochristina* was named from Utah and is related to *christina*. The ssp. status of *kluanensis* is doubtful (is it a syn. of *christina* at the TL?), because series from west of the TL in Yukon are distinct with grayer unh than *krauthi* and *christina*, but specimens from north of TL have unh yellow-green like *christina* and *krauthi* (though the unh at both sites has a little less orange than *krauthi*), and Butterflies of Canada states that there is a cline in S Yukon from *kluanensis* to *christina*. Ssp. *occidentalis* does not occur in NWT, this page should mention that C. Ferris Bull. Allyn Mus. 116:17 designated male as lectotype from Gulf of Georgia=Strait of Georgia beside Vancouver I., while the females from “Fort Simpson on Mackenzie R.” are other species such as *C. p. vitabunda*, which the book does state. *C. meadii* white females are very rare in Colo. and Wyo., where *meadii* flies in both subalpine and alpine. (By the way the Colo. *meadii* larva is identical to *C. meadii johanseni*.) *C. hecla* females are usually orange, and *C. canadensis* females usually white, though some are yellow; Scott 1986 first separated them as distinct species (Ferris described it as a ssp. of *hecla*). No dots are shown for *canadensis* along the Haines Hwy. where text claims sympatry. *C. nastes* varies from almost blackish to pale yellow, with considerable individual variation, so some of the N.A. names are not valid ssp., but the two ssp. in BC evidently differ and have different fw shape, though they don’t look much different on the photos (Guppy notes that he has since coll. *streckeri* in SE Yukon). Gorbunov’s book does not list *chippewa* from Russia, and Tuzov’s findings do not show sympatry of *palaeno* and *chippewa* so may have limited relevance. *C. interior* has a satellite spot (sometimes a small smudge, note photos), and N. Kondla states that the unh in N BC (Pink Mtn.) is more greenish (and Yukon area records of “*pelidne*” are not that; *C. interior* needs more study). *Skinneri* is a syn. of *C. pelidne minisni*. Alaska *gigantea* adults are actually quite large, so *mayi* is distinguished by very slightly less blackish on unh base and slightly less unh green. *Eurema* doubtfully comes from the word *heurema*, and *iole* is very doubtfully from the word *iov*, as there is never any purple flush on N. *iole* (the unh may be dark green). N. *iole* is not delicate and is somewhat migratory. *E. nicippe* photos should be orange, and N. *iole* should be yellower, as the yellows and oranges of *Coliadinae* did not reproduce well (their colors are bad in Scott’s book too).

E U.S. *phlaeas americana* is native, not introduced from Europe (where non-arctic pops. have brownish uns and tails, unlike *americana*); *phlaeas* also occurs in S Utah and Wind River Mts. Wyo. *L. phlaeas* larvae hibernate half-grown. Ssp. *henryae* is a syn. of *snowi*, which does have a bipartite range, separated by ssp. *artemisia* in the sagebrush zone in Wyo.-Mont.-Utah; the larva photo is Calif. ssp. *cupreus*=*lapidicola*; the unh is gray on photo. *L. thoe* is the proper name, as the figure of *hyllus* type is a female *L. thersamon* (A. Kopač 1983 *Priamus* 3:3-5, 1984 *Priamus* 3:95) or *alciphron* etc. *L. dispar* is not the sister species of *L. dione*. *L. dione* is a distinct species which is sympatric with *editha* in Laramie Mts. Wyo. without interbreeding; but *editha* (p. 358) is a ssp. of *L. xanthoides* and intergrades with it in N Calif. and the lower Sierra Nevada. *L. dione* is rare at lakes in Colo., where *thoe* and *helooides* fly by the million at the *Polygonum coccineum* zone around lakes and *dione* is absent, evidently because its eggs cannot withstand submergence (David Wright found that *dione* egg cells are quite wide and shallow compared to subgenus *Epidemia* eggs, evidently suggesting that *dione* eggs cannot sustain underwater plastron respiration for a long period); so BC folks should look in weedy meadows and along

creeks for dione, not around lakes (2008 update: N. Kondla has found dione in quite a few weedy sites & roadside ditches assoc. with *Rumex* within a limited range in SE BC). “Historically correct” does not apply to common names such as Dione Copper, because priority does not and should not apply to them; common names should be appropriate. Guppy told me that *Polygonum amphibium* is an error for dione as a host in BC or is rare. *L. rubidus* glue their eggs on just like other *Lycaena*. The larva photo of *L. heteronea* is ssp. *clara*, and the pupa is Calif. “submaculata” (=heteronea X gravenotata). *L. dorcas* and *helloides* are a close group, and *L. florus* is now considered distinct (see *Papilio* [N.S.] #12 & #18). Real *dorcas* is sympatric with real *helloides* on the flatlands of Alta. & NE BC; Yukon material is *L. florus arcticus* & *L. dorcas*; *L. florus* no doubt occupies much of BC (the single-generation populations), while real multi-generation *helloides* near-*helloides* occurs in S BC. C. Guppy has found real *L. dorcas* at Quesnel BC, assoc. with *Potentilla palustris* in a wet bog/fen. Fig. 64b looks more like *helloides* or *L. florus* (not *dorcas*) based on unh band and fw shape and spots etc., the live photo of “*helloides*” on p. 187 looks like *helloides* or *florus*. The photo of male “*dorcas*” from Valemont BC cranberry bog looks like *L. florus* or *helloides*, & the cranberry bog assoc. suggests *L. florus* to me (though Guppy insists they are not *florus* or *helloides* & are *dorcas*, possibly a new ssp.), and the female “*dorcas*” from Yukon looks like *L. florus arcticus* (*florus* is very variable, ranging from adults that are very dark to adults that are oranger). The “*helloides*” photos look like *L. florus*, not ssp. *helloides*, but they are from a two-gen. pop. in Vancouver I., so evidently they are an unnamed darker S B.C. ssp. of *helloides*. True *L. dorcas* is very different from all these pictures (darker, less orange, stubbier fw, black dots relatively closer to base, small). *L. nivalis* has all the unh spots of other *Lycaena*; the larva is ssp. *warnermontana*. Only very young *L. mariposa* males have a slight purplish tint (common to most very young male *Lycaena*, where they reflect ultraviolet even when older). *Charlottensis* actually differs by having a dark median unh band, not by yellow color; *penrosae* is a syn. of *mariposa*.

Benjamin & Ziegler provided the modern definition of *Satyrium*, prior to Clench. Ssp. *immaculosus* has smaller unh dots in most of BC, ssp. *titus* has larger spots but occurs only in SE U.S., so the ssp. in Peace River is evidently *immaculosus*X*watsoni* (the *titus* uns photo is very dark, surely an aberrant specimen, unlike Butt. Alta. photos). *S. b. columbia* looks identical to *behrii* so is a syn. *S. fuliginosa* does overwinter as eggs; the larva/pupa photos are “*maculadistinctum*”, a syn. of *semiluna*; the Waterton Lake dot is missing from map. Wash.-BC *S. californica* have less ups orange, later named ssp. *brashor* Kondla & Scott; the Waterton Lakes record is missing from map. The blue spot has orange cap in *californica* (but very reduced in many BC specimens according to Guppy), none in *sylvinus*. The map of *S. acadica* on p. 358 shows no dots in Idaho, meaning that the book treats *coolinensis* (TL Coolin, N Ida.) as *S. sylvinus*, implying that *nootka* is a syn. of *coolinensis*, because the name *coolinensis* is older; if Coolin adults are *sylvinus* and not *acadica*, then *coolinensis* is the proper name for *nootka*. However the *coolinensis* types are definitely ordinary *S. acadica*, and thus were mislabeled from Coolin. Ssp. *liparops* males rait (perch) on gulch trees to locate mates, perhaps explaining their presence on *Amelanchier*, though it could be a host. The “*affinis*” photo unh has many white spots unlike most *affinis* (Guppy says BC adults are variable for this trait, with at least a trace of white usually present, uns is lighter green than *sheridanii*); real *affinis* males rait=perch on hilltops to await females, which Guppy states is frequently true in BC. In nature the *sheridanii* pupae survive freezes and there is no prolonged emergence (Hiruma’s 200 days were a lab artifact). P. 210, *C. johnsoni* can be orange on ups. *Barryi* and *rosneri* are ssp. of *C. gryneus*, and the maps interdigitate nicely as ssp. should, and the photos do not show the diagnostic traits claimed (the *rosneri* photos do not show less contrast, the photo of mounted *barryi* is not pink, the live photo of *rosneri* is pink), there are no areas of sympatry (they interdigitate in S interior BC but are in different habitats because hosts occur in different habitats according to Guppy), and identification problems are severe (I thought most identifications for the maps have probably been made based on the presumed hostplant trees, not the butterflies, though Guppy insists he can separate them by wing phenotype). Warren’s Butt. Ore. recently corrected the *barryi* TL & suggested that *barryi* & *plicataria* & *byrnei* are all syns. of *rosneri*. The *augustinus* and *iroides* pops. are not closely adjacent on maps., and are surely one species. The “*augustinus*” photos are oddly dark chocolate on uns (Colo.-Ont. Calif. adults are reddish-brown on uns, not chocolate), so this is a new undescribed ssp., but is not ssp. *augustinus*; N. Kondla and now C. Guppy both think that these “*augustinus*” photos are really *iroides* (by the way, the only difference I can see between real *augustinus* and *iroides* is the unh base averages less dark--oranger--in the latter), and Guppy now thinks *augustinus* occurs in BC in western foothills of the Rockies. Distinguishing features of *C. mossii* are not listed; actually ssp. *mossii* from Vancouver I. is distinguished mainly by longer russet marginal unh spots, and the amount of white differs little, as ssp. *mossii* may have slightly less whitish overscaling on unh beyond white line (although the live ssp. *mossii* photo from Victoria I. has a lot of white like *schryveri*) and E Ore. “*schryveri*” and the Creston BC “*schryveri*” photo is intermediate between *schryveri* and *mossii* in the amount of unh white, thus BC lacks pure *schryveri*. *Callophrys polios* ssp. are rather worthless. The unf cell bars are rather worthless for separating *niphon* from *eryphon*. *C. eryphon sheltonensis* has a more purple uns; though N. Kondla has found that purplish adults are found occasionally elsewhere in BC. *S. melinus atrofasciatus* may be a valid darker ssp. with less red on unh postmedian line, with *setonia* a syn. of it. The mounted *atrofasciata* photos are dark because they are the spring (April) form; the other photos of *atrofasciata* and *setonia* are paler because they are the summer (July) form.

E. comyntas from Ore.-BC are later-named ssp. *pacnowe* Scott & Kondla with blacker rounder unf postmedian spots (and with slightly smaller orange spots on average). If *E. comyntas* was brought in by “nursery stock”, they must use

herbaceous legumes as nursery stock there (actually *E. comyntas pacnowe* is native from Ore. to BC). The female D and female V are given separate localities on 372 despite obviously (note antenna position etc.) being the same specimen. BC *amyntula* could be ssp. *amyntula* but the photos look like ssp. *albrighti*, and N. Kondla notes ~3 varieties in BC. Scott actually published that the *Jamesia* feeder is a darker-uns critter more like BC *ladon* (the whitish Colo. taxon is *C. humulus*), Scott did not mention BC, and Scott would use *C. neglecta echo* and *C. lucia* for several BC species. The map dots are almost entirely allopatric though, suggesting that the two species have been confused in BC and a more careful assessment of uns color and unh patch and ups color and voltinism (*lucia* has much more silvery ups blue, at least in Yukon & Edmonton) would greatly change the maps (and *C. Guppy* informed me that he now knows of 6 locations where *lucia* and *echo* are sympatric in BC, and they are easily separated by phenotype). *Guppy* now suggests that there may be three species in BC, *C. echo*, *C. nigrescens* (bluer ups, in SE BC), and *C. lucia*. Because the “echo” has only one gen. in BC (except for the one female), it might be thought that there is just one sp. in BC, but *Guppy* and *Kondla* assure me that there are several species, showing sympatry. In E U.S. the southernmost *C. lucia* has darker bluer ups, so maybe *nigrescens* is the darker-blue ssp. of *C. lucia*. *Kondla* now suggests that at Genelle BC 95% of adults can be separated into “*bakeri*” earlier, and “*nigrescens*”, which overlap in flight. The larva of *E. battoides* is “ssp.” *glaucan*, while the pupa from Riverside Co. is surely *E. bernardino*, a separate species, as Riverside Co. is south of the known range of *E. battoides*. The larva of *G. piasus* is ssp. *sagittigera*. The *G. lygdamus* larva is described as light green but the photo is pink. The female *G. lygdamus* “*couperi*” photo is *columbia* according to N. Kondla. Ssp. *couperi* is evidently limited to eastern Canada (where lowland Kananaskis Alta.-Man.-Ont.-N.B. males have darker-blue ups), because the Yukon-NW BC-Alta. mts. males are a distinct ssp. with very pale silvery ups that *Kondla* states is the Siberian ssp. *kurnakovi* (both “*couperi*” figs.). Scott (*Papilio* [N.S.] #12) improved the *Plebeius idas* mess. BC evidently has 1) *Plebejus atrapraetextus* which eats legumes (*Astragalus*, *Lupinus* in Colo. etc.)(*Guppy* [whose larva was amazingly multicolored] & *C. Schmidt* found that *alaskensis* eats *Lupinus* in Yukon, so is evidently *P. atra. alaskensis*), 2) *P. scudleri* in C BC to eastern Canada etc. which eats *Ericaceae*, and 3) the whitish-uns *P. anna* (with ssp. *vancouverensis*) in S BC. The book states on p. 234 & maps that *P. idas* is allopatric to *anna* in BC, but the ranges actually overlap (see *Papilio* [N.S.] #12). Scott prefers *Anna Blue* and *Melissa Blue* to the possessive case (these butterflies were never owned by those people). The *melissa* larva is Calif. ssp. *paradoxa*. *Kondla* notes that *melissa* in BC is evidently an undescribed darker ssp., which does not eat alfalfa. P. 238, the *saepiolus* uns wing base is never green, and *Kondla* finds that some BC males are a bit green on ups. I think *insulanus* from the TL is a syn. of *amica*, but *Saratoga Beach* adults from N Victoria I. have smaller spots, an undescribed distinct ssp. The larva photo is S Cal. ssp. *hilda*. No characters are listed to distinguish *pembina* from *montis*, the blue colors are bad on the photos (note the background colors), and these are no doubt synonyms, though *blackmorei* is distinct. *P. acmon lutzi* is actually *P. alupini lutzi*; real *P. acmon* ranges north to *Chelan Wash*. Ssp. *spangelatus* also occurs in the alpine zone in Alta. (& a similar ssp. *cotundra* in Colo.). *Peace R. glandon* is NOT *lacustris*, which is very different & heavily spotted beneath (TL woods north of *Lake Winnipeg*, not prairie); prairie pops. are an undescribed ssp. somewhat intermediate *rusticaXmegalo* based on the photos of prairie adults in this BC book & my specimens (*Chris Schmidt* will describe it). The “*mormo*” photo on p. 246 is *A. virgulti deserti* female.

P. satyrus uns varies from golden tan to dark brown; syn. *marsyas* is misspelled; the larva rests on the bottom of the leaf which is curled downward around it. The weakly-spotted-female-uns form of *P. faunus* is form *silvius*. Ssp. *rusticus* is the browner-uns Calif. ssp. (Scott, *J. Res. Lepid.* 23:209) which I doubt extends northward, though *Butt. Cascadia* suggests it occurs northward so Ore.-Wash. adults should be compared to real Calif. *rusticus*; BC adults are the grayer-uns recent ssp. *cenveray* from *Ida.-Mont*. There is no valid reason for separating *zephyrus* from *gracilis* (the male ups *zephyrus* looks more typical of *gracilis* with darker uph margin but the uns shows it is *zephyrus*, whereas both “*gracilis*” photos are actually *zephyrus*); *zephyrus* is distinguished by less-contrasting uns base-median vs. postmedian. Perhaps *Sugden’s* willow host of *gracilis* was *P. faunus*, although if it eats alder it may eat willow also (*faunus hylas* eats both willow and *Ribes*, so perhaps *P. gracilis* can also). The p. 254 photo of *threatfuli* male holotype may be female. *P. oreas threatfuli* is basically an intermediate between *silenus* & *oreas*, which I (in *Papilio* [N.S.] #12) considered an invalid intermediate. Further study may show *threatfuli* grading to *silenus* toward the coast; the rarity of specimens makes this difficult to study. Concerning *Roddia*, *Polygonia* have middorsal scoli on larvae also; *Polygonia* and *Roddia* have silver spots on pupa, absent on *Nymphalis*; the larval horns are small on *Roddia*; *Roddia* pupal shape is like *Nymphalis*; and *Roddia* evidently lays more eggs than *Polygonia* but less than *Nymphalis*. Thus *Roddia* seems valid. N. *Wahlberg* & *Soren Nylin* later plopped it back into *Nymphalis*, but I call it *Roddia*. Someone should look into the *Denis* & *Schiffermueller* names, if there are many and overturning them would cause much instability, then they do need to be retained (ICZN is considering them according to their website). *N. californica* is rare in Colo. too, where it is evidently a native. The idea that *herri* is due to “maturation in mountain environment” is doubtful, because altitudinal “forms” have proven to be genetically distinct when reared, and *herri* does not really differ from *californica* (the book does not give distinguishing characters). *A. Shapiro* evidently would say that the Apr.-June hill in the histogram are bugs migrating N into BC, and their offspring are the July-Sept. peak, which then migrate S. In *N. antiopa*, weeping willow is *S. babylonica*. *Aglais milberti* larvae are creamy on bottom half; only adults hibernate; willow and *Helianthella* are erroneous hosts without actual published data. *V. cardui* is orange, not pink, though *Guppy* states that fresh adults have pink tones that fade; at any rate the common name is frivolous nonsense like

most butterfly names. Overwintering of *Vanessa* seems difficult to study. It might be thought that *V. carye*=*annabella* does overwinter in BC as the graph shows two generations including many in spring and the map shows many dots, however there is only one record before mid May and adult hibernators first appear 4-6 weeks earlier than that. *V. atalanta* hibernates as adults, and actually any *Vanessa* could emerge in Feb. in Wash. and fly into BC the next day, but this must be rare or there would be more early-spring records; larvae fold the leaf upward and live on top of the leaf (the opposite of *P. satyrus*).

P. 271, most *Argynnis* (*Speyeria*) males have concave fw margin. *E. claudia* surely attempts to breed in BC and surely succeeds at times; there is no OFF-SEX-EGGS button pushed by customs at the border; it is rare so rarely breeds in BC. *S. cybele pseudocarpenteri* is dimorphic, just less so; S BC populations are nearer *pugetensis* than *leto* according to N. Kondla; *cybele* occurs in many habitats. Kondla (*News Lep. Soc.* 43:100) showed that *whitehousei* was named for A. C. Whitehouse, a Forest Service worker stationed in Jaffray. The tan circle trait on unh (between CuA_1 and CuA_2 , not beside M_3) of *aphrodite* is an interesting character, which does work most of the time, but often does not work (only 14 of 33 Elko specimens have the halo according to Kondla), and occurs occasionally on some other *Speyeria* (*cybele hesperis zerene callippe egleis* etc.); a better trait for *aphrodite* is the pale color along unh Rs vein across the disc, visible on all three photos. No distinguishing features are cited for *columbia* and *whitehousei*; *columbia* is a little darker on ups bases, while *ethne* is a syn. of *whitehousei*. *S. zerene sitka* seems to be similar to *hippolyta* (the unh submarginal band is not wide on photos as text states), which ranges along the coast from N of San Francisco to the Ore.-Wash. coast and the NNE Olympic Mts. (see Butt. *Cascadia*) and Alaska panhandle. *S. zerene garretti* is a syn. of *picta*; the female *garretti* photos are the *picta* phenotype, and the male *picta* photos are *garretti* phenotype, according to N. Kondla; who states that *zerene* does not retreat to high altitude in dry summers, though Guppy states this occurs (maybe the low ones died from drought). *S. zerene* larvae are not gregarious when young as eggs are laid singly; the larva/pupa photos are unfortunately too dark; the *bremneri* TL is Vancouver BC. (the locality on Holland's lectotype). The *S. callippe semivirida* disc varies from green to light brown. The *chilcotinensis* holotype is stated to be female, but the photo is male; the authors corrected this in *News Lepid. Soc.* 44: 51 by stating that the holotype of *chilcotinensis* on p. 277 is the female photo, not the male. The photos of *S. "atlantis"* are actually *S. hesperis brico*, and the confusion is amplified by P. Hammond's mixup in the text. N. Kondla (*Boreus* 22:17-24) suggested that the C BC *S. a. hollandi* dots are *S. hesperis brico*, but Guppy has found that those dots are correct, as *S. atlantis* is quite common in C BC in Black Spruce bogs and very wet open White Spruce forests, while *S. hesperis* is common in drier sites; they are almost always easy to distinguish. *S. hesperis helena* is a syn. of *dennisi* (*dennisi* is older, a mistake in the paper by Scott Kondla & Spomer, *Papilio* [N.S.] # 8); the *helena* female photos are male; *beani* larvae do have narrow yellowish heart-lines; *brico* is a perfectly valid ssp. which the book confuses with *S. atlantis*; "Northwestern Frit." name was coined by Hooper 1973; at Atlin they are not limited to the warm springs. N. Kondla has published (*Taxonomic Report* 3[1], 2001) that *S. hydaspes minor* (whose author is McDunnough) is on the coast, and *S. h. rhodope*=*sakuntala* is inland; they occur in any kind of forest. Mapping *S. m. eurynome* in Alta. and *opis* in BC would seem doubtful as bugs don't often respect artificial political boundaries, but Guppy states that they are easily distinguished, evidently by the browner disc & darker ups in *opis* (though those traits aren't apparent in my few specimens, which have a rather pale disc & are mostly unsilvered), & N. Kondla finds that *eurynome* occurs in Flathead Valley of extreme SE BC; *erinna* is usually silvered, and one wonders if *erinna* really has the odd range shown and occurs in BC, (evidently it does); *mormonia* males do not hilltop in a mate-locating sense; the TL of *washingtonia* is Mt. Rainier, and it is a distinctive ssp. (not a syn. of *opis*) with dark ups and dark-green disc. N. Kondla states that ssp. *bischoffi* is often silvered (almost 1/3 of specimens according to Guppy), and its TL was restricted to Anchorage AK by L. Grey *J. Lep. Soc.* 43:7; Guppy now notes that the lowland Yukon bugs incl. Kluane Lake are just *opis*, and *bischoffi* occurs in NW BC along the Tatshenshini River, and likely in the St. Elias Mts. in Yukon. *Polygonum* is the host of *B. napaea* in N.A.; *Vaccinium* mostly grows in woods (though *V. cespitosum* grows in meadows etc., and *B. napaea* occurs in meadows contrary to the name *napaea*), *Polygonum* in meadows (nearly all Scott's numerous and carefully-demonstrated *Boloria* hosts were ignored by this book). Gorbunov's book states that *B. napaea* and *B. alaskensis* are sympatric in the Ural Mts., so according to him N.A. has *B. alaskensis*, not *napaea*. Flight periods of *B. eunomia* reflect biology and overwintering stage, not the desire to avoid mating interference; numerous Scott hostplants of the truly polyphagous *eunomia* are missing. *B. eunomia* also has silver unh spots (p. 285). *B. bellona scoli* on T1 differ in length from those of *B. selene*; both *jenistorum* & *toddi* are synonyms of *bellona*, no distinguishing features are given; the disjunct range is very odd but evidently real. *Youngi* is an obvious syn. of *improba* (the *youngi* type was just a yellow aberration, see Holland 1931 pl. LV fig. 28); the E-C B.C. ssp. is *nunatak* Scott; N BC pops (Pink Mtn.) have some adults showing a suggestion of the traits of *nunatak*, but this pop. is ssp. *improba* also. *B. alberta* is surely the Alberta Fritillary, not Albert's; Edwards would have named it *alberti* if he had named it after Prince Albert, and *alberta* has a feminine suffix if named for a person. The "wide white area" on *B. polaris* unh is the row of white lunules like on *freiya* and *astarte*. Gorbunov treated *kurentzovi-erda* as ssp. of *B. polaris*, based on study of genitalia, habitat etc. Scott's 1986 book first invented a common name for *natazhati*, the Pleistocene Fritillary, which is perfectly appropriate; "Beringian Fritillary" invented later in Butt. Canada is misleading nonsense as it is not found in Siberia, so this inappropriate name must never be used again. There are lots of data showing that *astarte* flies every year at many Alta. sites such as Plateau Mtn. (records of Scott & Shepard, Butt. Alta.) etc. The difference between *astarte* and *distincta* is

vastly greater than the difference between *magdalena* and *mckinleyensis*, though a slight hint of *distincta* can be found in Pink Mtn. *astarte*. *Chariclea* belongs to *B. titania* (the wing pattern of *grandis* and European *titania* is very similar in numerous characters; we cannot set up species based on single weak characters, ignoring many others that point differently). (A lot of hypocrisy here: *distincta* is lumped with *astarte* despite large wing pattern/shape differences, and *myrina* etc. from NA is lumped with European *selene* which looks quite different, while *chariclea* is split from *titania* despite identical appearance in some ssp. & very minor valva rod difference, and elsewhere *mckinleyensis* is split from *magdalena* despite the only difference being the amount of red which is intermediate in Yukon.) The *ranieri* uns photo looks like *helena*. The *butleri* male photos are actually *grandis* (from Atlin BC, a range extension or mislabeled); Guppy has since found *grandis* along Top of the World Hwy. W of Dawson Yukon, with *butleri* in the alpine (many people consider that there are two species in N.A. *B. chariclea butleri* in the alpine and *B. titania grandis* lower down, except they hybridize considerably in some areas; that treatment is better than treating all American ssp. as *B. chariclea*, which is wrong as *grandis* is nearly identical to European *titania*). Scott proved that *titania helena* is polyphagous, though based on many dozen records the main host is various *Vaccinium*. BC *Phyciodes* “*tharos*” is actually *P. cocyta*; it has mostly one generation based on the graph, with very few records in May or after mid Aug. (note that *P. pulchella* has the same type graph). Porter & Mueller actually misidentified some *cocyta* as *tharos* (see *Papilio* [N.S.] #13), and thus only proved that Mich. and Ohio bugs are similar species. *P. cocyta* occurs on the plains of Alta. also (along river woods), sympatric with *tharos*; *pascoensis* is evidently a synonym until proven otherwise. The *P. batesii* crescent is usually paler than ground color, and there is no purple on any *Phyciodes* larva; young larvae rest on top of a web, not in it. Alta.-BC has *P. batesii saskatchewan*, named later (*Papilio* [N.S.] #13). All *Phyciodes* hibernate as 4th-stage larvae. Scott (*Papilio* [N.S.] #7, #10, #13) showed in nauseating detail that *P. pulchella* was the correct name, not *pratensis*, and Kondla & Guppy *J. Lepid. Soc.* 56:171 showed the same thing. BC has ssp. *tutchone* in NW BC and *owimba* in S BC (N. Kondla), both of which have orange antenna nudum, versus black in Calif. *pulchella*. Ssp. *owimba* is just as valid as most of the ssp. in this book, and a continuous distribution does not invalidate a ssp.; the book did not bother to investigate where the small neat-rowed *tutchone* changed into *owimba*. To demand here that valid ssp. require a distribution break would invalidate most of this book’s ssp. (most of the ssp. in this book are mapped as if they have continuous distributions into the next ssp., including *threatfuli* etc., although Guppy notes that there are abrupt topographic changes such as parallel mountain ranges & valleys that provide barriers between some of the book ssp. that don’t show as a gap on the map dots). *P. 303* meant to say that there are no *P. mylitta* records from the Lower Mainland Coast of BC (the map region shown on p. 10) before 1902, whereas it presumably was present all along in the Southern Interior map region (*P. mylitta* eats all thistles of several genera, so was probably native to the southern interior of S BC before Canada Thistle invaded, happily eating the same native thistles as does *P. pallida*). *C. whitneyi altalus* occurs in BC, and has pointier fw and brighter coloration than true Colo. *damoetas*; populations also occur below timberline where rockslides extend downward into the forest; Scott (*Papilio* [N.S.] #6) described *damoetas* early stages and Scott (1986) has photo of pupa. The book fails to state how *C. hoffmanni manchada* differs from *segregata* (it doesn’t). The book badly mixes up *Euphydryas chalcedona* & *ancia*. Actually the easiest way to see the valva prong is to merely brush the scales away from a specimen as Scott did; there is no need to remove the abdomen or dissect or mount or preserve it in a vial or slide at all, and Scott’s genitalia examinations did not mangle the valva using any of those procedures. (This accusation reminds me of Gordon Pratt’s ridiculous accusation [*J. Res. Lepid.* 30:175] that I mismeasured the unh spots of N Calif. *Lycaena xanthoides editha* X *xanthoides* populations, and later Pratt published photos of specimens identical to Scott’s and named those intergrades a ludicrous “subspecies” *pseudonexa*.) The BC book photos on p. 307 are stated to be “*paradoxa*” but all are the blackish coastal colon (“*perdiccas*”), they are NOT the redder *paradoxa* which should have been illustrated; *paradoxa* is a redder (look at Gunder’s original type photos) variable ssp. of *chalcedona* that originated from intergrading with *ancia*. Ssp. *colon* or near-*colon* occurs on Vancouver I and Olympic Pen. *E. anicia* isn’t completely reprod. isolated from *E. chalcedona*, as proved by Scott morphologically (and proved using electrophoresis by Peter Brussard and others)(see *Papilio* [N.S.] #12 for recent info), and p. 308 last sentence admits that true *paradoxa* can be distinguished from ssp. *hopfingeri* only by the valva. The *chalcedona* map missed the presence of *wallacensis* in NW Mont. (Missoula etc.). Page 309 “Biology” admits that blacker adults near the *paradoxa* range resemble *wallacensis* (which resembles *colon* but has a slightly longer valva prong). The unf postdiscal band character is rather worthless. The photo of “dark colour form” of *ancia* is actually an aberrant specimen (of course aberrations can be called forms, but I prefer to use the word “form” for commoner variations). The *ancia* map shows *bakeri* in SW Mont., where *bernadetta* occurs too. *P. 310*, the dorsal arm of other *Euphydryas* has small spinules too (see Figs 71a & b). The valva of *E. gillettii* is similar to *E. editha*, and DNA study by Niklas Wahlberg plops it phylogenetically into the middle of various Eurasian *Euphydryas*. The chart shows *L. arthemis* has only one gen. in BC, except for rare records. *L. archippus* may never have eaten apple as a usual host (it has failed to colonize Denver despite common apples and superabundant crabapples, yet is found along Denver creeks and all over W N.A. on *Salix exigua*); *Salix* was probably the most common host. *L. archippus idaho* is a syn. of *archippus*, it’s much less distinctive than *lahontani*, and is intermediate and close to *archippus* thus a syn. The name *burrisonii* was clearly named as a ssp. in the O.D. which did not mention the word hybrid, and is now a nomen dubium; C. Guppy suggests that *burrisonii* and *itelka*

differ and will designate a neotype. Scott would have preferred names for the two *L. lorquini* ssp. that are relevant to the butterfly, rather than using middle names of obscure people.

The separation of *C. tullia* into two species based on one weak character is rather dubious (for instance Fig. 73 shows a 2X ratio for both, despite claims of 1.5 and 2.5 in the text), and even if the drawings are poor and this genitalic difference is statistically valid (later photos by N. Kondla showed little difference), the difference does not mean they are separate species. Demorest Davenport published a revision of American *tullia* decades ago, and recent attempts to hypersplit the species have been debunked by G. Austin and others (by the way *C. tullia nipisiquit* never meets *inornata* either since they are separated by different flight periods and habitats). My series from Seward Pen. of “*viluensis*” seem synonymous with *kodiak* from Brooks Range-Denali Hwy. (so the Siberian *viluensis* does not occur in Alaska or is a syn.), and *yukonensis* does not seem distinct either. W Mont. pops. are generally considered to be *sweadneri*, which is a little paler than *inornata* (and Kondla finds *sweadneri* in SE BC, which flies in cooler moister higher areas than surrounding *ampelos*); *insulanus* and *columbiana* are syns. of *ampelos*; *benjamini* is paler than *inornata* though lumped by some. (In these *Coenonympha* and *Cercyonis* etc. some people like to split much more finely than I do.) Evidently all BC *C. pegala* are ssp. *nephele*; and the Alberta book showed no paler outer uns band. The *sthenele* text blames others’ misidentifications, and then misidentifies the male *C. oetus* “*phocus*” photos, which are really *sthenele paulus* (and *phocus* is a syn. of *oetus*; photos in Butt. Alta. are ssp. *oetus* also); Butt. Canada has a *sthenele* dot from ~Chapmans in SW BC. *C. oetus* unf lacks a brown line, not a light band; Scott’s *C. oetus* hosts are just as valid as all the other literature “hostplants” for Satyrinae (studying hosts of Satyrinae is difficult, because larvae are hard to find in nature, females often oviposit rather haphazardly, and larvae will eat numerous plants, etc.). The eyespots are not in line on the photos of *E. mancinus*, the text evidently meant “approximately” in line; one wonders whether the adults flying together on Dempster Hwy. were just picked out of a series based on one variable character without confirmation from other characters, however Butt. Canada mentions genitalic differences. *E. magdalena* and *mckinleyensis* are no doubt biennial; females oviposit on the edge of boulders near vegetation (grasses, sedges, rushes, etc.); *saxicola* is a synonym of *magdalena* unless you want to be a hairsplitter and peer minutely at a few body hairs that can’t be seen on photos; Yukon pops. are intermediate between *magdalena* and *mckinleyensis* in amount of fw orange, which proves that they are conspecific, since that is how they differ. Little sign of a bipartite range appears on the *E. discoidalis* map, though C. Guppy suggests there could be a gap in N BC. Gorbunov’s book suggests to me that *pawlovskaia* is a ssp. of *E. stubbendorfii*; and *E. theano* is limited to the Altai as the Russians treat it. The valva tip of one Siberian specimen (type of *tshuktscha*) resembles *youngi*, which with the slight difference and considerable variation in the four photos of *dabanensis* valva in B.C.S. Warren’s *Erebia* book, suggests to Scott that it is a variant of *E. dabanensis*, of which *youngi* is a ssp. The male photos of “*youngi*” actually seem to be *E. kozhantshikovi lafontainei* (note the spikes at outer edge of unh median band in mid cells, & paler uns); the unh has more of a slight reddish than pinkish tint. Some *E. epipsodea* larvae are green due only to food, and they surely have 5 rather than 3 stripes; the great variation in eyespot number is noted, yet the book persists in recognizing ssp., even though *remingtoni* is surely a synonym of *epipsodea*, and *sineocellata* no doubt is a syn. of *epipsodea* also; the Alberta book and Manitoba book photos resemble ssp. *epipsodea* also, and the Alberta book assigned prairie & Peace R. pops. to ssp. *freemani* = *sineocellata* and said there was little difference from ssp. *epipsodea*, and mentioned that the BC ssp. was *hopfingeri* (which BC book synonymizes with *epipsodea*)! It’s time to throw these ssp. names into the *epipsodea* dumpster, there are no ssp. within *epipsodea*! Scott (Papilio [N.S.] #6) showed *Poa pratensis* to be a hostplant of *epipsodea* in nature, among others. *O. chryxus* is an open-forest species in the Front Range of Colo. Still no discussion of the possibility that *macounii* is a ssp. of *O. nevadensis*, despite its allopatry and close geographical replacement and similarity in nearly all physical and behavioral traits (*macounii* lack the male stigma, have slightly darker upf veins, and on average the unh mesial band looks more distinct and unh striations are less, though the unh traits vary considerably in both); would a *nevadensis* female care if her mate had androconial scales? (obviously *macounii* females do not care, since their males lack the patch)? Why are their ranges barely allopatric, if they aren’t ssp.? (Guppy suggests the habitat changes). Papilio (N.S.) #12 divided “*chryxus*” into two species which both occupy most of BC, *O. chryxus*, and *O. calais altacordillera* & *O. c. caryi*. *Phalaris arundinacea* is doubtfully a host for *O. chryxus*, since this grass is 2 m tall and occurs in wet spots. The unh median band of *O. uhleri* is weak, not pale; *uhleri* never has two generations per year despite ancient ridiculous reports of two (C. Guppy told me he has seen two near Plateau Mtn. Alta., and I suggest that the late individuals were misguided oddities who got out of sync, as rarely happens in various butterflies); and Scott has never seen males hover and spy downslope (they merely perch in hillside swales near the host to await females, hovering and spying would require an enormous waste of energy); Scott proved that *Poa* etc. are larval hosts (oviposition, association, lab feeding, what else is needed?). The same ludicrous two-generation nonsense is repeated for *O. alberta*, will that ancient garbage never die? (two gen. was reported in error in the 1800s for Colo. *alberta*, where dozens of recent collectors have never found a 2nd); *Oeneis* are very slow larval feeders, and there is zero possibility of ever producing two generations, though a mixed-up individual may emerge in a wrong month. The photos of *O. “alberta”* from Peace River look like many *O. calais* near *altacordillera* to me, at least on ups, and the unf postmedian line doesn’t have as long a jog as most *O. alberta*; this bug flies early on dry low-altitude grasslands like other *O. alberta* in Canada, but are evidently orange so are at least a new ssp., and should be investigated further. Holland’s (Sandberg’s) description of *O. bore* larva stripes was very poor, and larvae may vary somewhat like *O. uhleri*; there is a two-year cycle in Colo. also; the

unh of my edwardsi specimens is not darker and less contrasting. The live photo of “mckinleyensis” looks like *Oeneis philipi*. *O. jutta* always has a two-year life cycle; ssp. *reducta* is oranger and is not in BC or Alta., the “*reducta*” and “*ridingiana*” photos are chermocki, and chermocki is a syn. of *alaskensis* (European *jutta* is similar but usually? has a tawny spot near upf tornus in cell CuA₂); the photos show whitish inner margin of unh median band. Nights in the alpine/arctic zone are usually too cold for *O. melissa* larvae to feed at night. Is ssp. *atlinensis* different from ssp. *melissa*? Many more dots exist in S Alta. for *O. polixenes* than shown on map; the photos of *yukonensis* are too overexposed white; *yukonensis* is similar to Colo. *brucei*, while Scott’s Alta. males are similar to Colo. *brucei* while most females are browner; *luteus* is yellower, and also occurs at Churchill Man. Gorbunov treats *philipi* as a syn. of *O. norna oeno*; his drawings of *norna* and *polixenes* genitalia suggest a resolution of *philipi*’s status, but Scott’s examination of *philipi* and *brucei* valvae indicate that they vary greatly (and average midway between Gorbunov’s pictures of both taxa, the average valva tip more square), with *philipi* valva tip perhaps averaging wider like *norna* but one individual is quite narrow only 25% of basal width; the valvae of these taxa and *O. melissa* are too variable to draw confident conclusions). N. Kondla suggests that the female “*rosovi*” is *O. polixenes*, and his photos of all the *rosovi* types (2 syntypes) show that neither resembles *O. philipi* (one is yellowish with weak unh band!). Why should the Kootenays have eastern N.A. monarchs??

Appendix 3 provides data for photos, a useful feature. Delete “*A. c. campestris*” on p. 357. P. 359, replace *dumetorum* with *perplexa*. Two *L. lorquini* ilgae live adults are listed, so which data go with which photo? *Shus?ap* is misspelled in *M. rosneri*. *P. oreas* threatfuli holotype is male. *S. behrii* is from Sherwin Summit. *S. liparops* larva has no locale (none available).

THE BUTTERFLIES OF CASCADIA. Robert Michael Pyle. 2002.

Seattle Audubon Society, Seattle, Washington. 420 p.

Most butterfly folks will like this book. The photos are nice, and the text has interesting digressions about the fauna. It covers “*Cascadia*”, an artificial area including Washington and Oregon, though the maps include some surrounding area. The book includes for each species something about identification, variation (subspecies), hostplants and number of generations, habitat and range, plus a little blotch map. This review mainly lists the things in the book I think need correction.

For easy identification, the text of a species and its photos should be on the same page, but in many cases photos of related species appear on a species’ page, which is quite confusing. Many of the photographs in this book are misidentified, especially in the tough groups such as *Argynnis* (*Speyeria*) and *Phyciodes*. To identify butterflies requires staring at them for a long time in order to develop an ability to recognize their subtleties, which is as difficult as identifying human faces (thus people can identify and name only a few people, those they have stared at the longest), and I think this requires a large personal collection that one can stare at interminably, so butterfly watchers have a disadvantage here. Another disappointing feature of the book is the wholesale inclusion of numerous recently-named subspecies, especially those creeping into the fauna from California and Nevada, which appear to be synonyms or are very weak. I now recognize many more subspecies than are in my 1986 *Butterflies of North America* book, but I still think many recent ssp. names are not distinctive enough. The map on p. 20 should have included the names of the counties to show the reader where they are. The book sometimes uses strange nonstandard english that inhibits communication, including “*afforesting*” on p. 46, “*flickering*” (*fluctuating*?) on p. 361, “*cumbrous*” p. 75, “*oystery*” on p. 60, “*individuating*” on p. 181, etc. This book, like the BC book, I think wastes a lot of space on the etymology of the scientific names, an academic exercise having nothing to do with the butterflies since the butterflies exist in nature just fine without names. Some people seem to like this exercise, but many scientific names are inappropriate, misleading, meaningless, frivolous, or stupid, so deducing their origin involves a plunge into idiocy. Contrary to p. 35, there were three strong efforts to organize common names of N.A. butterflies, the first being Scott 1986, who—most importantly—also corrected all the inappropriate misleading names (such as correcting “*Early*” *Hairstreak* *E. laeta*, which flies in July, to *Turquoise Hairstreak*, correcting the *Lupine Blue* to *icarioides* not *P. alupini*, etc.). It is quite disappointing to see so many folks (such as the untrained amateurs in the North American Butterfly Association) ignore appropriate names and return to bad misleading inappropriate names, especially when there is no Code that requires the use of older bad names. And it is disappointing to see so many lepidopterists still using priority, or counts of the frequency of use of bad names by trashy books, to choose between common names (such as the *Red Admirable* in this book). The name *Lupine Blue* should not be used for *Plebejus alupini*, when the butterfly has absolutely nothing to do with lupines (larvae do not eat lupines and adults do not nectar on lupines), and the name is totally perfect for another species *P. icarioides* which is totally restricted to lupines. (As a humorous aside, maybe we could transport these common name transgressors to the international border S of Osoyoos, where local politicians cannot agree on whether their river and jurisdiction is the Okanagan or Okanogan, and we must, using the handle of a stout butterfly net, nag on their noggins until they confess their sins and agree to fix those names.) As I stated in my critique of one compendium of common names (*J. Lepid. Soc.* 47:170-1), “The study of butterflies is young compared to the study of birds (because there are fifty times more ornithologists working on half the number of species), so there is no need to rush into mandating particular common names when so many of them are bad. The lesson...is this: common names should be appropriate descriptive names for the

common person; they should not be the most common error.” It is nice that the book does have the courage to reject some of the inappropriate common names used by NABA.

The lists of hostplants found by Pelham here and there in the book, are unfortunately not documented (ovipositions?, larvae found?, association?) and are not published elsewhere; Pelham told me that most are based on larvae found or ovipositions, but at present we have to wonder if they are all valid.

Detailed comments: Foreward: Lepid. Society is losing members, thanks to harassment of lepidopterists by the U.S. Government. P. 24, the worst conservation problems in the west are overgrown pest trees (Lodgepole Pine, Engelmann Spruce, etc.) choking out butterfly habitats due to 100 years of misguided fire suppression by the federal government etc., weeds such as *Bromus tectorum* (cheatgrass), *B. inermis*, *Centaurea diffusa* & *repens* etc. overgrowing whole landscapes (*B. tectorum* dries in June and later often catches fire and burns up native sagebrush etc.), and development and water diversion ruining lowland riparian sites (probably 50% of *Speyeria nokomis* sites were wiped out by 1900 by water diversion). P. 28 Norbert Kondla has a bigger collection than other private NW collectors, and has published much more than any other area lepidopterist. P. 45, *A. vialis* photo is a male. P. 49 what does “sub-cell” mean? P. 50 spelled *aemilia* (L. Miller J. Lep. Soc. 39:51); all butterflies flap their wings below and above the body; *mexicana* raits on hilltops. P. 51 *icelus* does not eat locust. P. 53 *pacuvius* is rather readily identified by the strong upf mottling; at high density *pacuvius* males usually patrol their hillside hostplants. P. 55 *fredericki* (a syn.); upper photo looks like a female. P. 56, *centaureae* ssp. are weak and the top photo has a very whitish unh because photo is overexposed. It occurs in both subalpine and alpine zones, and subalpine is below timberline. P. 57, *P. scriptura* also flies L Apr.-May. P. 58, Kondla finds that *Pend Oreille communis* pop. has one gen. M June-M July in shrubby open woods, and Pelham notes that there are numerous univoltine foothills pops. P. 59, *albescens/communis* may be a dominant/recessive polymorphism (A. Warren says that *Mex. P. communis adepta* [which mostly lacks costal fold]/*albescens* relationship is complex, as some males which lack costal folds have *albescens* genitalia, and some males with costal folds [*albescens*] have *adepta* genitalia, which I think implies that costal folds and genitalia are polymorphic and may be just a dominant/recessive situation, which would explain why *albescens* can take over Florida and displace *communis* (what usually happens when one species invades another’s range is one bug moves in, while the other bug remains at least for a while). Rearing studies are needed (raising whole families from a number of wild-caught females, cage matings). P. 60, the waxy bloom on Hesperiid pupae is best called “glaucous”, it is not blue. P. 62, pupa may have glaucous waxy bloom but is not purplish. P. 63 both photos are of course ssp. *libya*; and true ssp. *lena* has a solid cream unh and occurs in Colo.-Mont. and obviously does not occur in Nev. P. 65 the *oregonia* m f are near-idaho. P. 66 the ssp. are weak and difficult to assign in *C. palaemon* (Pelham). Kondla says *tan-unh mandan* occurs in boreal forest/aspen parkland in N Alta.-NE BC, *russet-unh magnus* in cordillera mts.; *C. Guppy* has new research on these. The Tilden & Mattoon paper did not even report unh color in their giant table! I do not have confidence in any *palaemon* ssp. now as adults vary a lot and everyone who looks at them comes up with a different scheme because the differences are overwhelmed by the variation. P. 70 *eureka* is a synonym; *muertovalle* is limited to Death Valley-Needles CA. P. 72 both photos of course are ssp. *uncas* (Warren reports *terraclivosa* in Ore., which looks like *fulvapallaXmacswaini* thus is invalid & has darker unh like *macswaini* thus is a syn. of *macswaini*, but based on Pyle’s description of white veins, ssp. *lasus* occurs in Ore., not *macswaini*). P. 73 *juba* adults do NOT overwinter, explaining the total lack of any specimens collected between the two Sept. and May gen. *H. juba* is not much bigger, and Scott proved (*Papilio* [N.S.] #6) that *Bromus tectorum* (a winter-annual grass) is a very common host in Colo. of the winter generation, and this bug has 2 generations; its larvae grow much faster than other *Hesperia*, permitting a 2nd gen.; the Oct-April larvae feed on winter-green grasses at lower temperatures, the May-Sept. larvae feed on a few green and maybe not-so-green grasses at higher temperatures; Shapiro’s work was quite incomplete, and his 1st-gen. males must have picked up stray pollen in May when the new adults emerged from larvae (pollen falls to the ground and lasts thousands of years in soils, so it surely can overwinter on an old flower head that a spring male rests on while doing raiting=perching behavior for mate location in some gulch). P. 74, *manitoba* TL is *LaHache* [Lac la Hache] BC; *harpalus* is not a syn. of *idaho*, it is the ssp. in Cal. Sierra Nevada (TL far west of Carson City in the Sierra Nevada of Calif. where unh spots are reduced and yellower); *oregonia* is a useless name. P. 65 J. Pelham told me that plate 2, 3rd row left and middle, and p. 75 left two photos, are all the isolated Puget Trough *H. comma* [near idaho] with unh spots flat white or even tan, pearly in some females. P. 75 right middle photo, Pelham and others think this represents montane forms of *H. comma idaho*, which occur in montane areas within the range of lowland *idaho*, and evolve independently. This is controversial, since all “altitudinal forms” that have been reared have proven to be genetically different, and darker *manitoba* genes in the ice age probably just moved upward in altitude and interbred with *idaho* moving northward in the lowlands, so the “habitat convergence” producing the similar appearance of such high-altitude bugs is probably mostly due to shared genetic ancestry. Scott believes that *H. colorado* belongs to *H. comma* after all and the purported cases of sympatry in Canada do not exist. Since *oregonia* is a useless intergrade name, *H. c. hulbirti* might be used for W Ore. bugs with greenish-yellow unh and yellowish spots also. Pl. 3, *juba* male is female. P. 80 *sierra* is a syn. of *nevada* (and the Front Range Colo. pop. has a slightly more tan unh than greener South Park pops.). P. 82 *tenebricosus* is a syn. P. 83 it was common in S Minn. in 2002 also. P. 85 Scott repeated the old stuff that *peckius* larvae and pupae both hibernate but I don’t believe it, larvae probably hibernate. P. 86 *alkaliensis* is basically a syn. of *sabuleti*, my series from Lake Co. Ore. is definitely ssp. *sabuleti*, and *aestivalis* is a syn. of *tecumseh*.

P. 87, there are usually lots of Sept. flowers. P. 88, waxy pupa is glaucous again; and *klamathensis* is basically a syn. of *mardon*. The lower right photo looks like *O. sylvanoides orecoasta* f. P. 90 *draco* is in NW BC too and Yukon. P. 92, *themistocles* seemed absent or scarce on Colo. *Poa pratensis* lawns for a long time, but in the last few years became common again (did it suffer from parasitoids brought in by the invading *P. peckius*, then adapt later to immunologically encapsulate most of the parasitoids??). P. 96 *flavaventris* is described as similar to *sonora* but uns a bit yellower perhaps, so is a weak ssp. at best, and doubtfully occurs in Ore. P. 98 *nemorum* isn't very valid either. P. 99 *omnigena* is intermediate between *sylvanoides* and *bonnevilla*, near *sylvanoides*, and lacks characters of its own, so is invalid. Pelham suggests that most NW *sylvanoides* are somewhat intermediate, but my Yakima Co. bugs are ssp. *sylvanoides*, and I have seen near-*bonnevilla* from Grant Co. to Kennewick. P. 101 upper left photo is female. P. 102 *lutea* is a syn. of *yuma*. P. 104 *kiowah* is a syn. of *vestris*. P. 106 *vialis* hosts are hay grasses, not *Poa* or the lab hosts (*Papilio* [N.S.] #6 has these and tons more hosts). P. 110 I think virtually all those *clodius* ssp. are useless synonyms; BC book got them wrong I feel. Pelham suggests that most lepidopterists have difficulty with these *clodius* ssp. names, and thinks the Snake River pops. are derived from uphill pops. in Blue Mts. Pelham suggests that the cryptic larval morph may better define some ssp. P.111, lower left photo is male, upper right female. P. 112 all are synonyms of *smintheus*, except *olympianna* (smaller, a little darker esp. females, due to altitude evidently), Mont. *maximus* (much darker females), and Calif. *sternitzkyi* (misspelled). P. 113, I think *smintheus* is still a ssp. of *phoebus*, look at Shepard & Manley's 1998 fig. 11 micropyle with magnifier and notice it looks quite like *phoebus* (at least intermediate), and their *phoebus* characters of *Sedum rosea* and moist habitat occur in SW Colo. ssp. *pseudorotgeri* as well as *phoebus* (and Evgeni Zakharov finds that a Siberian taxon *bremeri* that Shepard ignored actually belongs to *phoebus* also and intergrades with it). P. 114 C. Hauser (Nota lepid. 16:34-43) wrote that Parnassiinae/Troidini are the same subfamily, because Troidini branched off its base first, then *Cressida-Euryades*, then *Hypermnestra*, then Parnassiini; the other subfamily Papilioninae includes Papilionini and Graphiini. P. 116 there is no evidence suggesting *oregonia* is a distinct species; even the text says everyone agrees it is the same species as *bairdii*; the only experts who wrote about this assigned it to *P. machaon*, and Keith Brown's recent Papilionid book placed it into *machaon*. Yes it is large and beautiful, but that isn't sufficient grounds for distinct species. P. 117 *dodi* TL is Dorothy Alta., allotype Red Deer R. near Nevis, 100 km NW Drumheller (Kondla 1981 Blue Jay 39:144, & Kondla Alta Natur. 26:39). *Pikei* is intermediate to *machaon hudsonianus*, wing pattern more similar to *oregonia*, and host *Artemisia dracunculus*; it differs by having eyespot tending to *machaon*, fw stubbier than *oregonia*, tegulae less yellow, not by the worthless traits listed. P. 120 *shastensis* is a syn. of *indra*; the types of several new supposed ssp. of *indra* (including this and *pygmaeus*) do not match specimens later collected at the type locality, according to Steve Spomer. P. 121 *canadensis* is a ssp. of *P. glaucus* (and *rutulus*), the page struggles with this. Layberry's map isn't as good as Guppy & Shepard's; the book didn't mention Scott & Shepard's paper (Pan-Pac. Ent. 52:23-28) demonstrating intergradation in wing pattern & male & female genitalia in SE BC. By the way *alexiares* wings and valva look like *rutulus*, yet its hybrids with *glaucus* had high fertility and 1:1 sex ratios (J. Scriber & R. Lederhouse J. Res. Lepid. 27:222-232), females are often black like *glaucus*, and its mitochondria are very similar to *glaucus*, and it is accepted as a ssp. of *P. glaucus*, so *canadensis* & *rutulus* should be given the same chance to be included. Perhaps mating is sometimes less than random?, but they sure do produce lots of intermediates in nature. (By the way, Donovan & Scriber J Lep. Soc. 57:25-35 just published data on *glaucus* & *canadensis* in Mich. that show that they lack prezygotic reproductive isolation in the field and mate freely [Deering & Scriber 2002, J. Ethology 20:25-33] and the genes from mixing have spread 200 miles from the mixture zone, there are big step-clines in the genetic traits that distinguish them such as ability to eat *Liriodendron*, they proved they hybridize in nature using numerous allozyme/rearing tests, hybrid larvae survive just as well as parents in the lab [Table 2], hybrid morphotypes are frequent in Wis. [Luebke et al. 1988]; thus they have clearly shown that *glaucus* & *canadensis* are one species, unless their species concept is one of those 23 different species concepts nowadays that has nothing to do with reproductive isolation.) P. 124 *pusillus* is a synonym or very weak ssp., as book photos from BC and Wash. books and Butt. Pacific Northwest resemble Colo.-Mex. adults more than the *pusillus* holotype, and the holotype looks like an unusually pale odd spring form (allotype looks more normal), and in Colo. May adults are small and paler, late summer adults are large (sometimes huge) and darker and females sometimes have orangish inside the black borders. So the only valid ssp. seems to be the oranger *grandiosus* from Guatemala-Chiapas (not SW Mex.) (incidentally the *multicaudata* TL is only "Mexique" the loc. on syntype, the TL should have been restricted). Pp. 130-131 *N. menapia*, top male photo is ssp. *menapia*, and middle and lower photos are of ssp. *tau*; middle photo is typical of *tau* (Pelham told me it is typical of predominant form in Puget Trough) which is characterized by wider upf black (not by unh red of females which is similar to ssp. *menapia*) and is limited to Puget Trough, not quite as melanic as *melanica* from Outer Coast Range of N Calif. (an odd bipartite range of melanism); Scott has normal ssp. *menapia* from Lane Co. Ore.- Lewis Co. Wash.-Mt. Rainier-Mont. P. 133 brain not retina, the brain interprets what we see, often wrongly in the case of those magazine brain-fooler drawings; 30 years ago it was found that the butterfly retina takes info from many ommatidia and combines it into an image before it even gets to the brain, so insects don't see those ludicrous 100-image pictures that Hollywood erroneously projects. Science Jan. 24, 2003 has an article about Sphingid night vision concluding they combine color info from numerous ommatidia to get better color night vision than humans have. P. 134 *flavitincta* is distinguished by dusky ups bases according to Pelham; most females of other ssp. are yellow or yellowish also; *elivata* is a syn. of *sisymbrii* with no

distinguishing features. P. 135 should mention size of posterior black postmedian spot on protodice male upf, etc. P. 136 Shapiro's little cross was done with very few individuals. (The study of *Phyciodes* mtDNA by Wahlberg Oliveira & Scott, Syst. Ent. 28:257-273, showed that of 140 adults studied, there were 99 different sequences of DNA, and *tharos cocyta batesii pulchella* were highly polymorphic and overlapping. It seems that studies that just use one or a few individuals to analyze electrophoretic alleles or DNA, get results that erratically depend on which individuals are randomly tested, thus the results are quite likely to be randomly misleading.) Tuzov's book reports *nelsoni* from E Siberia, but Gorbunov's book suggests *ssp. callidice* ranges from Europe to E Siberia and maps a continuous range from the Urals to E Siberia. P. 138 Kondla notes that *marginalis* is a forest species, it is people who are edge species. There are many unfinished problems in *napi* group in BC book, such as *guppyi/tremblayi* surely belongs to *angelika*, and the hole in map of *marginalis* is nicely filled by *oleracea* implying it's a *ssp.* in an unfinished classification, etc. And Pelham suggests *angelika* might be a syn. of *pseudobryoniae*. P. 139, upper left is male, lower left female. P. 142 eggs are not blue; *insulanus* not *insulana*; *mayi* and *transmontana* are obvious synonyms of *coloradensis*, which isn't very distinct from *ausonides*. N.A. *ausonides* has been treated as a *ssp.* of European *E. ausonia*, but *Euchloe* taxonomy is a mess (see Gorbunov's book etc.). Europeans have switched the usage of the names *simplonia* & *ausonia* recently (*ausonia* Hübner was named for Mt. Ausoni in central Italy, and *simplonia* Freyer was a Boisduval nomen nudum that Freyer properly named but incorrectly listed from Croatia but Bernardi corrected its locale to Simplon Pass, and now it's applied to pops. in the Alps & Pyrenees), so N.A. *ausonides* looks more like what Europeans now call *simplonia* (but the upf cell bar touches costa in *simplonia*, not in *ausonides*). Complicating this mess, our *Yukon naina* is said by Gorbunov's book to be a *ssp.* of *E. ausonia* (and rare high-altitude individuals from Colorado resemble *naina*), and *andrewsi* from San Bern. Mts. looks like *ssp. E. ausonides*, not *hyantis/lotta* which ranges very close to *andrewsi*, leading one to think that someone mixed up their *andrewsi* larvae with *ausonides* to claim larvae are like *hyantis*. So a lot more study will have to be done before we can assign Palearctic *Euchloe* names to American taxa with confidence. P. 144 bottom states that *hyantis* and *lotta* are allopatric and allochronic, but p. 146 left bottom suggests they are separate species because of lack of interbreeding. Actually, *hyantis* has not been proven to be a separate species from *lotta*, and the book's maps clearly show no range overlap between *hyantis* and *lotta* in S Ore., so "lack of interbreeding" has never been tested, and I was informed that they intergrade on Kern Plateau Calif. P. 148 the book is correct about flora, and correct that *stella* is limited to Sierras and has slightly yellowish males. P. 151 Pelham says male *C. occidentalis* photo is correct but photo is too pale. The male & female *pseudocolumbiensis* and the female *occidentalis* are evidently *C. edwardsii*, which ranges into BC according to N. Kondla (*ssp. alexandra* does not occur in Wash. as p. 160 claims). P. 152 keys should not be used, because they involve a sequence of decisions, any one of which if made wrong will result in a totally wrong identification, so the total likelihood of success is small; tables should be used instead, because one can compare every character for the specimen in question, and narrow it down quickly and get a great idea of what it is and how it varies from normal, one can quickly find the most unusual trait in the table and see if your specimen has that feature, and the likelihood of failure is small because a missing character or odd character state will not totally derail the process, plus it's a lot easier to expand a table than a key, variation can be included in a table but not in a key, and a table simultaneously acts as a description. Nobody should ever use or make keys; use TABLES of taxa versus characters. P. 154 I have not noticed any range expansion of *philodice* in Colorado, but Pelham told me that in Wash. it prevails in June from 3000-5000' then evidently expands its range in E Wash. and Puget trough through the summer. P. 156, likewise I am puzzled at how much *eurytheme* migrates. Pelham has seen *eurytheme* migrating through Seattle in Aug., and thinks it expands higher in altitude during the season, and Crispin Guppy saw a migration of thousands in July in Flathead River drainage 6000', and Orley Taylor wrote that it migrates northward during the season. So why have I never seen *C. eurytheme* migrate in Colorado?; in Colo. it doesn't act like a migratory species, and I often find it after strong fall freezes as adults survive freezes just fine unlike the regular migrant butterflies. The theory that adults and larvae die overwinter and it migrates north each spring to repopulate is wrong according to Jack Harry, who caged N Utah alfalfa & found that they survive fine overwinter at 0°F mostly as 2nd-3rd-stage larvae. P. 155 top right is male *C. occidentalis* according to Kondla and myself; Orley Taylor & Robert Silberglied originally studied uv of *philodice/eurytheme*, and its polymorphism is just a single-gene trait that may or may not be associated with reproductive isolation (in *C. occidentalis christina/sacajawea*, some males are orange and some are yellow with the associated uv difference, yet they are one species). P. 160 *columbiensis* females are nearly always white, occasionally yellow; *ssp. alexandra* is not in Wash.-Ore. P. 161 lower photo is *pseudocolumbiensis* based on locale, not *C. o. occidentalis*, though N. Kondla thinks it is *edwardsii* which actually extends into BC. P. 158-61 are confused about *C. occidentalis* and *C. alexandra*. *Papilio* (N. S.) #12 clarified the situation somewhat. The BC book lists a number of *occidentalis* dots from W Wash. and BC. P. 159 cites Scott's book in error, Scott merely lumped all *occidentalis* and *alexandra* into *alexandra*, following Ferris (as for the "ever the iconoclast" bit, "one who attacks established beliefs or persons", Scott often defies convention merely because progress marches on and so much of the old dogma proves to be wrong; people who do the most research disprove the most dogma; after I published my book, I found that a lot of people were annoyed at my name changes, but they rushed out and did research that solved some problems, so stirring the pot does work to speed research. Now we have 23 species concepts and so many name changes made by nearly everyone that there is total chaos in the taxonomy of many butterfly groups, and many people are disconcerted, but progress continues). P. 163 there is actually no SE BC *gigantea* record. P. 167 *iole* has

to migrate across divides to get to the watercourses. P. 169 Gossamer seems too frivolous and misleading to me. P. 170 should mention whether SW Ore. adults are whiter on the uns thus are ssp. *arota* (probable) or not, the unh of the photo female does have dark unh esp. tornus like ssp. *virginiensis*=schellbachi; size is not meaningful as it doesn't vary much that I've seen. P. 171 *alpestris* is not darker-gray beneath. The egg/1st stage overwintering stage is an error according to the info I have; half-grown phlaeas larvae were proved to overwinter by Gordon Pratt, and in general subgenus *Lycaena* species overwinter half-grown, subgenus *Tharsalea* as egg; however alpine Alta. phlaeas would probably be biennial also, so maybe egg and half-grown larva would hibernate, after all phlaeas is the latest-flying arctic butterfly so would not have enough time to grow to half-grown larva. My little note (J. Lep. Soc. 47:253-4) disproved the idea that ssp. *americana* was introduced from Europe, because European phlaeas have browner unh and tails, while the Scandinavian *polaris* has grayer unh and no tails but is said to have brassier ups (see plate in Higgins & Riley) and larger unh spots; I have adults from S Sweden that don't look like *americana* either. P. 172 *henryi* is a syn. of *snowi*; BC book's argument that it is valid because ssp. *artemisia* divides its Alta.-Colo. range is wrong, especially when that book [wrongly] lumps *C. whitneyi* *altalus* with *C. w. damoetas*, which have the same ranges as the *cupreus* ssp. The *Ida*. dot is surely ssp. *artemisia*, which is a valid ssp. that is roughly intermediate between *cupreus* and *snowi* if you add up all characters but is an odd mixture of character traits, the ups color most like *cupreus* in males but females are yellower, the ups spots are larger than either, the unh is grayer more like *snowi*, female uph darker than *cupreus*, habitat is upper sagebrush zone more like *cupreus*. Ssp. *artemisia* supposedly occurs in Lake Co. P. 173 *L. thoe* has two generations. P. 174 *L. dione* is correctly a separate species from *L. xanthoides*, because *dione* is sympatric with *L. xanthoides montana* in LaBonte Can. in Laramie Mts. Wyo., and is much different from the others distinguished by long orange unh band and solid black unh spots. But *L. xanthoides editha* is a ssp. of *xanthoides*, which have short scalloped orange band and tan-centered unh dots and differ basically only in size. *L. x. "editha"* "*pseudonexa*" is actually the intermediate populations between *xanthoides* and *L. x. editha* in N Calif. (note also that Emmel Emmel & Mattoon's *L. xanthoides nigromaculata* photos are rather similar to *pseudonexa* photos), but *pseudonexa* has to be treated as a synonym in my subspecies concept since it has no different characters of its own and is just intermediate *xanthoides X editha* (the factor analysis paper of Pratt et al. [J. Res. Lep. 30:175-195] wrongly lumped these intermediates into *editha*; that paper was a useless black box with no usable extractable data, and proved that they can't even distinguish the larvae of *rubidus* from *xanthoides/editha*; that paper pathetically accused me of mismeasuring the spots). Shapiro's papers on N Calif. butterflies clearly labeled these as intermediate and showed that the intermediates occupy a large range in the Siskiyou Co. area in N Calif., and Warren's Butt. Ore. showed the intermediates in SW Ore. also; they are not just a local thing (obviously the opinions of local lepidopterists should be carefully considered, as Pelham emphasizes). P. 175 middle right photo is *dione* female, lower right is *dione* male. P. 176 *xanthoides* does not remain separate where it comes near *editha*, intermediate pops. also occur at Mather near Yosemite (and also on E slope of Sierra according to that factor analysis paper). *L. x. montana* was declared a homonym of a Eurasian bug by Kocak 1984 (Priamus 3:95), who renamed it *vurali*, TL Broadwater Co. Mont., HT USNM. P. 177 misspelled *jacquelineae* a good ssp. with larger uns spots, but *dorothea* is a syn. of *jacquelineae* (note that Dornfeld's female *dorothea* topotype is not melanic, and his book describes females as having extensive yellow). P. 178 those *duofacies* & *perkinsorum* ssp. are rather worthless, evidently synonyms, and Pelham also questions them; Johnson & Balogh's original paper was not convincing, *sirius* and *duofacies* were described as having both brown & orange females, *rubidus* and *monachensis* were described as having brown females but the illustrated females were orange, and maybe only *perkinsorum* has brown females usually; in Colo. *sirius* is the brown female more common at high altitude while at low altitude the females are mostly orange; better to just call the brown female form *sirius*. P. 180 *klotsi*, *submaculata*, and *rava* are all syns. of *klotsi*, which also can be considered a syn. of *heteronea*, since there are no populations totally lacking unh spots. And *gravenotata* is a good enough name for the Siskiyou thing; the primitive "plesiomorphic" state was surely the spotted unh of all other *Lycaena*, and then the spots decreased in the Great Basin sagebrush where bugs get whiter, so Colo. and Cal. *gravenotata* could share the same name due to presumed shared ancestry. Emmel & Pratt wrote that *gravenotata* is in SW Ore., not *submaculata*, and actually they wrote that *submaculata* is limited to W Sierras N to Shasta Co., and *submaculata* is intermediate between *heteronea* and *gravenotata* thus is a synonym because it lacks any different characters of its own and is just an intergrade; they wrote that ssp. *heteronea* types were darker blue, but that's just because the specimens are old, those 50-year old San Francisco *icarioides pheres* and *lygdamus xerces* in Cal. Acad. Sci. are also darker blue on ups merely because of an unexplained darkening of the blue due to age (blues just seem to get darker blue after 40 years or so in museums). P. 182 the violet tint isn't refraction, it is the violet-wave-length tail of the uv reflection (or diffraction maybe, but not refraction, which is light bending due to its speed slowing when going through a denser substance like water). P. 183, top right & bottom right from Kittitas Co. look like *L. florus*. Pelham informed me that dark forms (like those found in *florus*) occur all over the Wash. steppe to montane habitats and Puget Trough, without any geographic consistency, although dark forms increase in montane univoltine pops. but not to the exclusion of normal females. So, because *florus* is characterized by a gamut of forms from quite melanic to oranger *helooides*-like and a single gen., and ssp. *helooides* is characterized by fewer dark forms (there are dark forms even in Calif.) and several gen., Wash. surely has *L. florus* in montane univoltine pops., and has *L. helooides* (maybe darker N ssp.) elsewhere that vary somewhat (perhaps some lowland pops. along Columbia R. may be *L. h. near helooides*?). (True *L. dorcas* ranges south to Quesnel BC. and C Alta.) P. 184 *bichroma* is a syn. of *nivalis*; actually

browni=praetexta differs by having lilac occupying only the outer 1/3 of unh, vs. outer 1/2 in nivalis=bichroma, the average color of yellow is actually the same, and I have ssp. nivalis females from Loon Lake in higher Sierra for instance that are deep yellow on unh also; and Pelham notes Blue Mts. pops. with significant numbers of individuals with the mostly unicolorous creamy unh, in among the yellower ones; evidently there is widespread polymorphism for yellow/cream unh color; praetexta is a syn. of browni; the photo of "browni" has unh half lilac so would seem to be ssp. nivalis, so Wash. may have ssp. nivalis, not browni; eggs are not blue, they are maybe sl. bluish-green when laid but rapidly turn white like other *Lycaena*. P. 186 penroseae (note spelling) is a syn. of mariposa, and charlottensis differs by having a dark median unh band; Pelham notes a distinctive segregate in the N tier of counties, E of the Okanogan R. in Wash.; uliginosum misspelled. P. 189 estesi is a syn. of haesus, and dolichos=juanita occurs in Ga.-Fla. (see *Papilio* [N. S.] #18). Perpetuating an error such as Great "Purple" Hairstreak is worse than creating it in the first place. P. 190 lorquini is syn. of grunus (and a Baja ssp. "poodiae" I will recognize as a valid species when poodles grow wings & fly). P. 192 occidentalis is a syn. of watsoni; Rosa woodsii is an amazing new hostplant, which Pelham told me is based on ovipositions and larvae he found in Wash. steppe habitats, where Rosa is often the only woody plant present with titus other than *Artemisia*. P. 193 columbia is at best a weak ssp. resembling behrii, as my 8 columbia from N of Chelan Wash. resemble behrii closely although they maybe average a bit larger unh spots; crossii is the only distinctive ssp. (much larger, darker uns, different host). P. 194 tildeni is a syn. of fuliginosa, maculadistinctum a syn. of semiluna; as the photos from Kittitas Co. show, these things are quite variable; ssp. albolineatum from Lake Co. Ore. was missed, it looks valid to me with a row of black unh dots. P. 195 P. sabuleti sabuleti and P. s. ministigma are the same species and the latter has a smaller stigma (Mex. norae with no stigma is a ssp. of sabuleti too). (By the way, I often wonder whether *Oeneis macounii* is a separate species from nevadensis, they are amazingly identical except for the male stigma, if macouni females don't require the stigma, maybe nevadensis females don't either.) The low-altitude fuliginosa may need its stigma to distinguish itself from *S. californica* etc., whereas the high-altitude one without stigma presumably lives with no other *Satyrium*, as other species are absent at high altitude. P. 196 obscurafacies and cygnus are synonyms; "red-capped" should be in bold, not the other stuff. P. 197 note this whole page refers to *S. sylvinus*, not californica as I first thought! P. 198 the *Salix exigua* bug has been spreading in Colo. (from W Colo. to Arkansas River N to Castle Rock), maybe its overwintering eggs on small *Salix exigua* shoots are being mowed and spread in hay bales, much like *Thymelicus lineola*, but the future of an egg in a hay bale would appear to be grim. I have a few adults from N Ida. that seem to be nootka but some have more orange unh lunules, so I thought that coolinensis is really nootka so the latter is a synonym of coolinensis, but Mike Fisher & N. Kondla have looked at the coolinensis types and they were acadica, which must have been mislabeled "Coolin Ida." The pale-uns things with tail can be called ssp. sylvinus everywhere, as typical lowland Cal. sylvinus has a whitish uns (more whitish than grayish); in NW Colo. I found a slightly-paler uns critter in a sage area (the rest of Colo. has darker putnami). P. 200 okanagana probably a synonym; picture in BC book doesn't look much different. It's not "rough-housing", it's just investigative mate-locating behavior. P. 201 lower right is *S. tetra* female, note kinky unh line, and gray scales like Glassberg's tetra photo beyond postmedian line. P. 204, the photo is the real *C. [sheridanii] viridis* "dumetorum" from San Mateo Calif., but the taxon as described is surely a variety of *C. sheridanii* [viridis] lemberti or newcomeri X lemberti perhaps intergrading toward newcomeri, and Pelham says probably so. P. 205, oregonensis is an intergrade between perplexa and affinis so is technically not a valid ssp., and Pelham states that oregonensis is a Lotus-feeding syn. of perplexa and is similar to the Mason Co. Wash. perplexa; these have the unf less orange than Calif. perplexa so are distinctive, though they are intermediate perplexa X affinis. Scott treats perplexa as a ssp. of affinis, because affinis intergrades with homoperplexa (which is very similar to perplexa) in S Wyo.-Neb. ("intergrades [of affinis] with *C. apama* common" *C. Ferris*, 3 mi. E Laramie, Wyo., 1997 *Lepid. News* #2 p.34; & G. Gorelick *J. Lepid. Soc.* 59:181-199)(yet interdigitates altitudinally in S Utah as washingtonia and oregonensis do in S Wash.), and there isn't real evidence (just nearby distributions) that apama/perplexa and affinis are distinct species in Utah or Wash.-Ore. Scott and Justice (*J. Res. Lep.* 20:81-85) tabulated seven characters of colors and spots of these northwestern critters and showed that the Blue Mts. and Okanogan Co. washingtonia actually are somewhat intermediate between affinis and oregonensis, some characters being closer to one and other characters closer to the other; Pelham directed John Justice to most of the localities; Pelham states that Ft. Simcoe has only sheridanii and affinis but not oregonensis, so Scott & Justice's one female "oregonensis" from Ft. Simcoe must have been misidentified, but that doesn't change the results of the paper since 40 other oregonensis were analyzed. P. 207 newcomeri isn't much different from neoperplexa, and the "neoperplexa" photo fits description of newcomeri; neoperplexa has always been described as lacking [actually reduced] a black edging of unh white stripe, and my 6 newcomeri from Okanogan Co. and 2 from Yakima Co. actually have some black edging esp. posteriorly, so these two names do not seem to differ much. *C. sheridanii* males perch on the ground or on low plants because they perch in teeny little swales (most often on hillsides, even near a ridgetop); in contrast male affinis perch mostly on hilltops, thus mostly on sage bushes since it tends to be everywhere in their habitat; Pelham notes the same perching behavior of these two in Wash., where affinis will choose a prominent bush or even fencepost on a ridge; homoperplexa perches in gulch bottoms and on hillside trail bottoms, generally on low plants. P. 210 one could say that johnsoni is rare so has been missed from earlier-succession forests, but Pelham states that all johnsoni colonies were in old growth/virgin forest, and has seen no evidence that it colonizes adjacent earlier-succession forests (but of course all old trees eventually die, so johnsoni must colonize new trees at least locally). P. 211 ninus is a

syn., and pupae probably overwinter under loose tree bark as *Atlides* do in Calif. (perhaps old trees are better for *johnsoni* in providing more loose tree bark for pupae?, and older trees may have more mistletoe?, but the same factors should apply for *C. spinetorum*). P. 212 *barryi* is evidently a diverse wastebasket name for the juniper-feeders, if the uns varies from yellowish-brown to violet and purple, but the latter two colors no doubt refer to the overtone or sheen; as an exercise, the wing pattern variation should be mapped separately from the hostplant variation, and then we could see if there is concordance. (Warren's Butt Ore. has corrected the *barryi* mess; *barryi* is a syn. of *rosneri*.) P. 213 Kondla says the ICZN code states regarding *rosnerae/i* that inadvertent errors must be corrected but incorrect latinization is not an inadvertent error, so *rosneri* must be preserved; but elsewhere the code states that endings must be properly latinized, and patronyms must be given a latinized suffix according to its gender; how do we reconcile these contradictory and rotten articles in the new code? (maybe we change the endings or not, however we wish?!). P. 214 *concava* is a syn. of *augustinus* as it has dark unh base, and the only difference between *iroides* and *augustinus* in my series is the unh base is pale in *iroides* & dark in *augustinus*. P. 215 that chocolate-brown thing in BC book is not *augustinus*, it's a ssp., because eastern N.A. *augustinus* are reddish-brown like *iroides* (*C. Guppy* Jan. 2003 told me it is a new unnamed ssp. of *iroides*). P. 216 my series of Vancouver I. *mossii* are distinguished by having unh red wider and postmedian area less whitish; *windi* has unh white line weak. Males *rait* in gulch bottoms. P. 217 *obscura* and *maritima* are both synonyms, my Colo. females have just as much slight rufous on uph as the E N.A. ones. Evidently Bob did not inadvertently name "teninoensis" in this book, since there is no description/diagnosis (in general, manuscript names should not be mentioned in print because of the possibility of inadvertent valid publication under ICZN rules); *kinnikinnick* misspelled often. P. 218 *sheltonensis* is a good ssp. but *purpurascens* is a syn. P. 221 *atrofasciata* photo is paler than *setonia* photo (the April *atro.* photo in BC book was darker than July photos of both ssp. because of spring date); *setonia* is a syn. of *atrofasciata* and both are evidently characterized by less black on unh line; Pelham agrees that *atrofasciata* and *setonia* refer to spring & summer gen. and only one name is needed; abdomen is orange only on summer males, spring males have no abd. orange. P. 225 *isola* has dozens of legume hosts and does not favor *Melilotis* (*Papilio* [N.S.] #6). P. 226 hw rubbing evolved once I believe, as *Eumaeini* and *Polyommata* are sister taxa (I haven't read *Eliot's* latest classification). P. 226 the *amyntula* ssp. are a mess. *Albrighti* may be a valid local form or ssp. in Mont. and a form in BC, but *maritima* probably occurs over most of Canada. P. 228 top photo could be *E. comyntas* perhaps, but lack of orange spot fits *E. amyntula* better; at that site near Douglas Ariz., *comyntas* occurs in lowlands and *amyntula* in mts. P. 230 Wash. evidently has several species of *Celastrina* including *C. (neglecta?) echo*, and *C. lucia* with all the unh patch forms, and maybe *nigrescens* too (which may be a ssp. of *C. lucia*). Pelham says *Celastrina lucia* occurs in Wash. The BC book didn't make a good case for two species because the sky-blue *lucia* and the darker-blue *echo* are essentially allopatric there, but Kondla claims that *echo* and *nigrescens* are different species sympatric in SW Alta. and NW of Vernon BC. Kondla says *bakeri* & *nigrescens* are not synonyms of *echo*, and states that *bakeri* has a violacea or *marginata* uns and flies earlier, while *nigrescens* flies a little later (overlapping a bit) and has a more maculated *lucimargina* uns (I think these two are probably uns forms of one sp.). A. Warren's Butt Ore. claims to find *echo* & *nigrescens* sympatric in Ore. Pelham states that *echo* always has at least a partial 2nd gen. in Wash., but *echo* has just one gen. in BC as Kondla says the "summer form *echo*" female photo in BC book is the only known summer female of small 2nd brood of *echo* in southern interior BC., and the BC phenogram shows mostly one gen. Chris Guppy now suspects there are 3 species in BC, *lucia*, *echo*, and *nigrescens*, which all fly in the N Okanogan Valley, *nigrescens* occurs in the E half of S BC, and he has found a new ssp. of *nigrescens* in the C interior of BC. So, I conclude that there are surely two species in BC and evidently southward, and there are lots of opinions, but someone must start observing ovipositions and rearing larvae to determine how many real species there are. P. 231 Warren's Butt Ore. says the photo is *C. nigrescens* ovip. on *Holodiscus*. Surely the ability to express the unh patch of form *lucia* is genetic, even if the patch is induced by photoperiod/temp., so there's not much difference in the theories. In Colo. Front Range the *lucia*-patch form of *C. lucia sidara* is rare but occurs and the patch is small, while in W Colo. in seemingly identical temperature regimes and not much more moisture, the *lucia* patch form of *C. lucia lumarco* is common and the patch is twice the size, so surely this difference is genetic as photoperiod is the same and temperature similar. David Wright informed me that *Cherry Gall Azure* and *Hops Azure* and *argentata* may be same species, eating mostly *Deervilla* in the north (Maine to Minn. etc.); if that species got to Colo. it perhaps ranges to BC and even Ore. as *nigrescens*?? (what does *nigrescens* eat?). P. 231 Paul Ehrlich told me years ago that "Sooty Azure" was a bad name because *Cel. nigra* has no blue; he was right so I named it *Spring Sooty* in my book. P. 235 top right photo maybe is *E. enoptes*? but is from N Wash., one would have to get series and find hosts to determine this. Concerning the "probable and presumptuous" comment, too bad this characterizes most of the *Eriogonum* feeders, including *Euphilotes*, *Plebejus acmon/alupini*, and *Apodemia mormo*. P. 236 *leona* is obviously a ssp. of *P. speciosa*, and possibly a syn. of *bohartorum*. P. 238 Alaska-Yukon-Alta. mts. has ssp. *kurnakovi*, not *couperi*; *xerces* blues are obviously a ssp. of *G. lygdamus* (half of them [form "antiacis" etc.] had unh black dots like *pardalis* rather than white blobs, the ups is darker blue on dead museum specimens due to darkening of the blue, hosts were identical, etc. etc.). P. 239 *toxeuma* a syn. of *daunia*; isn't a "lunker" a fish?. Plate 240 *icarioides* (lowland) painting is ssp. *fulla*, not ssp. *icarioides* which has distinct black unh postmedian dots, and lower right *saepiolus* is female not male. P. 241 *spangelatus* misspelled. P. 242-3 I think *ricei* is a syn. of *anna*, as the amount of orange varies from little to much even in Calif., and Pelham finds the same great variation in Wash. Scott (*Papilio* [N.S.] #12) revised this group and documented sympatries

between species, and for the bugs in Cascadia used the names *Pleb. anna* (ssp. *anna* & *vancouverensis*), *Pleb. atrapraetextus* (ssp. *atra.*, near-*longinus*, & *benwarner*) & *P. melissa melissa*. Pelham states that *melissa* and *idas atrapraetextus* are sympatric in Wash. & Ore. at least later in the season, and “ricei” has near sympatry with *atrapraetextus* in Ochoco Mts. *Glycyrrhiza lepidota* is shunned by *melissa* in Colo. *Fridayi* is a ssp. of *Pleb. atrapraetextus* that occurs only in higher Sierra Nevada of Calif. P. 246 *P. saepiolus* is a complicated mess that Scott corrected in *Papilio* [N.S.] #12 & #18. *Ssp. aehaja* does not occur in “Cascadia”. *Ssp. rufescens* evidently isn’t in it either. *Ssp. saepiolus* probably occurs over most of it, as I doubt that Marin Co. females are much bluer as noted above (in Syst. WNA Butt. review, which shows that *maculosus* from E Nevada has ordinary uns spots and resembles *saepiolus*). *Littoralis* is evidently a weak ssp. or form of *saepiolus*, distinguished by white rings around unh spots, but I have some like that from Siskiyou Co. (Scott Camp Crk. and S of Copper Ore.) where that phenotype is a form. *Ssp. kodiak* & *insulanus* (a syn. of *amica*) aren’t in Cascadia (*amica* may be in NE Wash.). P. 247 lower right photo is male not female. I fixed *P. icarioides* in #12 also. P. 248 I thought *montis* & *pembina* were synonymous, but *montis* is higher-altitude montane and *pembina* lower steppe-semi-steppe according to Pelham; but Pyle’s book states upf border of *montis* is narrower, but it is wider in BC book photos; is it valid? *Ssp. icarioides* has whitish uns so rings disappear and uns looks like *saepiolus*. *Helios* is a syn. of *icarioides* (see Dornfeld’s Klamath Co. photo with unh rings not visible). *Ssp. fenderi* is identical to ssp. *icarioides* but a bit darker below. P. 250 *icarioides* is the Lupine Blue obviously, and preoccupation refers to homonymy in ICZN code, NOT to common names, please. To repeat an error is twice as bad as making it in the first place; after all the original creator of the error thought it was correct, whereas the repeater knows it is wrong yet repeats the error, making two wrongs (printing an error, and failing to correct it). P. 250 lower right photo is male *P. melissa*, note narrow ups border and pointy fw. P. 251 *pallidissima* is a syn. of *minnehaha*. P. 252 lower photo is *P. alupini* (“acmon” in this book) *lutzi*, not ssp. *alupini*. P. 253 Scott’s “true acmon” from Chelan Wash were collected July-Aug., and Pyle reversed Pelham’s findings; Pelham finds that all along E slope of Cascades exists a zone where some colonies are univoltine and others bivoltine at least partially (in Columbia Basin there is only one gen.), and the second gen. is smaller and more “acmon”-like than the spring brood, so these populations look like “*alupini*” in the spring and *acmon* in summer and fall. In Calif. pure *acmon*, early spring adults also look more *alupini*-like (darker-inner edging to uph orange band), so all of these E slope Cascades multivoltines are most likely true *P. acmon* and not *P. alupini lutzi*. And Warren’s Butt. Ore. finds true *acmon* over much of Ore. Scott’s published note actually contained lots of data, considering that it used all of Goodpasture’s results plus more. Scott’s 1986 book published no map of anything in Willapa Hills, his dot was in Olympic Mts. P. 253 the *E. ovalifolium* host of Olympic Mts. *spangelatus* was discovered by Paul Opler and printed in Butt. Cascadia without mentioning him. P. 262 mating pair from Lincoln, Mont. (same pair uns on p. 275) is obviously not *brico*, and seems to be *S. zerene picta*, note the reddish uns & pale unh submarginal band & lenticular silver submarginal spots & wide dorsal black bars etc. that suggest *zerene*. P. 262, Steve Spomer and Sterling Mattoon and I are just as qualified as national *Speyeria* authorities as Paul Hammond. (And *S. edwardsii* is definitely not directly evolved from *S. callippe semivirida*, contrary to Hammond J. Res. Lep. 29:63). P. 263 pl. 9, *coronis* photo is definitely *coronis* according to Pelham & Scott & Spomer (though N. Kondla thinks it is *zerene*; the unh submarginal silver spots have pointed caps like *zerene*, and I have specimens of *zerene* like this, though my *zerene* have narrower median black upf spots and oranger ups than this photo). Pl. 9 middle row right *zerene* has uns photo that is identical to 4th row left uns photo, and is *picta*=*garretti*, not *bremnerii* which has more solid russet all over unh. The right photo in 4th row is most like *picta*, and Pelham agrees it is *picta*, so unfortunately there is no uns photo of *bremnerii*. 4th row middle “*picta*” specimen is a *picta* in Burke Museum according to Pelham, but is a pale variant that resembles ssp. *gunderi*. 5th row left is *S. hesperis beani*. 5th row middle photo from Tiger Mdws. Pend Oreille Co. looks darker somewhat like *S. atlantis hollandi* to me and Kondla (and Kondla told me he caught both *hollandi* and *S. hesperis* near-*beani* in forest beside Tiger Mdws., then found only *hollandi* at sedge fen/grassy marsh W of Ione). Also, Steve Kohler has found both in NW Montana. However, this specimen is probably *S. hesperis* near *brico*, because S. Spomer told me he reared specimens from Tiger Meadows in 2002 (livestock sent by Hammond) and although some were very dark like *hollandi*, they all had the wing shape and size of *beani*, and larvae were typical dull-black like ssp. *hesperis*, not even “intermediate” like the *beani* from BC Spomer has reared before; Spomer’s reared adults from there have the disc reddish-brown but some are dark-brown like *hollandi*, the ups border are oranger like *brico*, the unh submarginal band mostly rather narrow like *brico* (not quite as narrow as *hollandi*), and the unh disc is usually connected to silver spot at tornus like *brico*, although the unf tornus spots are strong and black like *hollandi*. Thus this photo is evidently *S. hesperis* near *brico*. Spomer thinks the one “*atlantis*” phenotype Hammond reared from a *hesperis* female from N of Missoula Mont. that produced a gamut of phenotypes, was just a dark *beani* (or maybe a galloping larva got returned to the wrong box), as these bugs vary. More rearings should be done here, as the larva color evidently is an important key to identity in this region as only *S. hesperis* has the blacker larva (*S. atlantis hollandi* has a paler larva than the black *hesperis*-type larva of near-*brico* here). Thus this confusion of *S. atlantis* & *S. hesperis* in BC and N Wash.-NW Mont. is an identification problem, not a species problem, as there are clearly two species *S. atlantis* and *S. hesperis* (many individuals in *Speyeria*, *Phyciodes*, *Colias*, *Euphilotes*, *Pieris*, etc. cannot be identified, even when the species are valid). P. 263 5th row right is *S. hesperis dodgei*. P. 264 *eileenae* is a syn. of *pugetensis*, which is a valid ssp. my book should have included. *Ssp. leto* is linked with *cybele* through the sequence *cybele*>*carpenteri*>*charlotti*>*leto*. Rod Davis once reared *cybele* from Bear Paw Mts. Mont.

which apparently showed a nice cline between cybele and leto, and Moeck mentions intergrades from there and in Alta. P. 265 lower right photo is male. P. 266 bottom photo is *S. hesperis hesperis* from Jefferson Co. Colo., one of the occasional variants that is 2/3 silvered. Ethne is a syn. of *whitehousei*, while Spomer & I now think that *columbia* is a distinct ssp. that is small with dark disc and darker ups bases (Scott only has 5 *columbia*, which have ordinary disc, and a bit darker ups bases, and have paler ups; it isn't strikingly distinct). P. 267 upper left callippe is male not female, bottom photo is female. P. 268 ssp. *hippolyta* evidently occurs all the way from Olympic Mts. to Wash. coast and Ore. coast and Point Reyes Cal. (as *myrtleae*). P. 269 top right *picta* "male" is female, middle right *picta* "female" is male (and S. Spomer says this could be *zerene* or *hydaspe*). P. 270 Pelham says photo is *S. coronis simaetha*, so that must be what it is, it's greener than the small numbers I've seen. P. 271 top left Pelham says is *S. zerene*, I think that's correct; bottom left I and Pelham and Kondla agree is *zerene* also; right is male not female. P. 273 top left is female not male, top right maybe is male; bottom right Pelham and Spomer say is *S. hydaspe* and I think that's correct, the uph submarginal crescents are M-shaped which is frequent in *hydaspe*. P. 274 *viola* and *dodgei* are vars. of *irene*. Hammond's theory that the *irene* type is *S. egleis* is interesting, so I looked at Emmel Mattoon's photos of type figs. 252-3, and find that they are actually *S. hesperis*, not *S. egleis* (note especially the clear pale median spots in unh disc of their female fig. 253, the clear pale spot at base of unh cell CuA_1 is characteristic of all *irene* [and is clear on male *dodgei* on Pyle's p. 275], but is absent on all *egleis* in which that area is the color of the disc or the whole area is paler; and the unh submarginal pale band is not suffused with a little brownish as is *egleis*, although the posterior part of this pale band has an infusion of disc color in my *irene* more than the photo; the upf black bars are a little darker like *irene* rather than the weaker bars of *egleis*; the unf of females of both *irene* and *egleis* is darker redder on basal area than is this photo, but the photo is horribly bad too pale and too yellow; also L. P. Grey surely examined this specimen and called it *atlantis*, and was surely aware of the *egleis* possibility because of the words *egleis* and *irene* on the label, so obviously Grey identified it as *atlantis* also. So I reject this argument, *cottlei* is an *irene* bug (those unh median spots are very clear and obvious identifiers). The book is rather confused about *brico* because the *brico* photos on p. 275 & 271 are misidentified: p. 275 top right is female *S. hesperis beani*, not *brico*; bottom right same mating pair as p. 262 is *S. zerene picta*. P. 275 bottom, as noted above Hammond surely had females of *S. hesperis* and his reared "chocolate...atlantis" proved to be dark "hutchinsi" by larval color; and from the book statement, the two females could have been separate species; anyway his statement that they "intergrade" is just difficulty of identification. Scott, Kondla, & Spomer wrote the 1998 paper, and Ron Hooper invented the name Northwestern Fritillary. (Note: in the *Speyeria* paper by Scott Kondla & Spomer, *Papilio* [N.S.] #8, *dennisi* is older than *helena* so is the correct name for that ssp.) P. 276 *S. hydaspe*, Kondla has since written his paper showing *minor* is the proper name for coastal small often-silvered ssp., and *rhodope* is the correct name for what was called *sakuntala*. The unh in N Cal. still looks darker to me making *purpurascens* valid, but I haven't examined hundreds. P. 279 *washingtonia* generally has a darker-green disc than those shown. Some *S. mormonia* adults esp. females descend to low alt. in late summer. P. 278 top is *S. coronis* male, not *mormonia*, note fw shape and ups color and black spot size and little marginal black. P. 283, diapause is 4th stage; *sierra* is a syn. of *chermocki*. P. 285 *astarte* has fw more pointed (not squared) than *distincta* and unh median band is narrowed posteriorly. P. 286 *ranieri* is a syn. as it is similar to *grandis* (looks nothing like arctic *chariclea* to me), while Mont. "ingens" is a syn. similar to *helena* (and an intergrade anyway thus not a valid ssp.). And *grandis* is very similar in wing pattern to European *titania* in many characters, which overpower the one character of slightly-wider little valva rod, making N.A. bugs *B. titania* contrary to Shepard. *Vaccinium* is the most common of many *titania* hosts in Colo. (*Papilio* [N.S.] #6 again). P. 289 *oregonensis* is a syn. of *leanira*; both photos are *C. l. daviesi* from Cal. Sierras; *basinensis* is a syn. of *alma*. P. 290 *carlota* a syn. of *gorgone*. P. 291 *manchada* a syn. of *segregata*, note the male ups photo on p. 296 is identical to Dornfeld's Butt. Ore. male ups of *segregata*. P. 292 photos are Colo. ssp. *damoetas* (note convex fw etc.), whereas ssp. *altalus* with concave fw etc. occurs in B.C., contrary to Shepard; habitat is alpine AND SUBALPINE talus in Colo. and Alta. and Wyo. and Calif. P. 293 female *dorothyi* are dark, males often dark on ups too, and *dorothyi* is very variable though no females are quite as light as *acastus*; Kondla reads ICZN details to suggest *dorothyi* is correct even though ending was wrong sex; I personally refuse to change sex of endings as that rule is bad (for example the original author may have not intended the names to be latin, and non-latin names should not be latinized—all my names were not latinized because I do not know latin and I spelled the names to make them sound pleasing, and thus any changes of my names are contrary to the code because the names were originally not latin), and the rule causes endless instability and will be changed sooner or later. P. 293-4, evidently *sterope* actually is a ssp. of *acastus*, not *palla*, and *dorothyi* links the two, according to info from Pelham, as explained in A. Warren's Butt. Ore. *C. acastus dorothyi* is extremely variable, and is closer to *acastus* though no females are as light as *acastus*; *dorothyi* TL is from an area where paler *acastus*-like and creamy *sterope*-like females are mixed together, and not far southward the predominant female form is *acastus*. *Sterope* has 100% creamy "eremita" females, and males can be much more heavily marked as well. The ventral bands are slightly yellowish but some can be whiter or rarely even pearly. *Sterope* occurs in steppe desert of Columbia Basin, barely penetrating the forest-steppe ecotype, habitats like *acastus*. The hostplants of *dorothyi* and *sterope* are the same (*Chrys. viscidiflorus*), and behavior [mate-locating] in steppe-desert draws is the same. The creamier females and creamier uns spots may have entered *sterope/dorothyi* from *C. palla*, just as introgression evidently introduced the yellower unh spots into *C. acastus vallismortis*. *Sterope* is marginally sympatric but largely allochronic with *palla* in the Simcoe Mts. of S-C Wash. and Blue

Mts. of Wash.-Ore. [the sterope records in 7 cos. in N-C Ore. in John Hinchliff's checklist belong to *palla*?, and the Columbia Co. Ore. record is an error evidently]. *C. palla* does not extend up the E slope of the Cascades, where it is replaced by *sterope*. Creamy females are absent in *C. palla* in NE Wash., and some are creamy in S Wash. but nowhere do they predominate. *Sterope* females are never reddish. Thus from Pelham's info and the situation in Calif., it seems that *C. palla* and *C. acastus* are just barely distinct species and introgression has been frequent, overlapping evidently only very narrowly in the S Sierra (where *C. palla australomontana*—which is very similar to *C. acastus vallismortis*—and *C. acastus neumoegei* barely overlap), (*acastus* and *palla flavula* overlap in W Colorado, at Basalt Eagle Co., S of Glenwood Springs in Garfield Co., and Minturn in Eagle Co.). The larval scoli on *sterope* should be examined and compared to *palla* and *acastus*, as they may differ in length in Calif. Norbert Kondla has found specimens in BC that may be *C. acastus sterope*. P. 294 in Calif. the Coast Range *C. palla eremita* has females mostly cream, and in lower Sierras (TL of *palla*) females are supposed to be most often fulvous but are often intermediate or cream, then in high Sierra ssp. *altasierra* has females orange (rarely cream/orange), thus the proportion of cream females in *palla* is intermediate between *eremita* and *altasierra* and there are no other usable characters (except the male ups which may be less dark in foothill and high Sierra), so one of the three names must be synonymized. Nitpickingly it would be *altasierra* since it is younger than *palla* and both have most females orange and male ups may be slightly paler; however cream females are frequent in the Sierra foothills so perhaps we should call ssp. *palla* the whole mess from lowland Sierras and Coast Ranges to S-C and SE Wash. as Pyle's book lists, realizing that the percentage of orange females varies geographically somewhat; then *altasierra* is the paler orange high-altitude bug from Sierras & Crater Lake to Mt. Jefferson, and *calydon* in the Rockies has male ups margins darker. The hosts are doubtfully sufficient to put *altasierra* into *acastus*, because *acastus* eats *Chrysothamnus* often, yet I found it in Colo. and Wyo. on *Aster glaucodes*, and *palla* eats *Erigeron* in Colo. and *Aster*, *Chrysothamnus*, *Solidago*, *Senecio*, elsewhere, so neither species is very specific. P. 297 left middle is *P. mylitta* from "Madeira" (Madera Co. evidently) Calif. Jack A. Levy, not *oriseis herlani*, and note in particular it has only a couple dark postmedian dots (*herlani* has a whole row of those dots and a whole different unh); middle right is *P. pulchella* female also from "Madeira", evidently ssp. *pulchella* from the bit of dark upf showing but Madera Co. goes from San Joaquin Valley to alpine so this could be *deltarufa* or *montana* maybe. P. 298 *pascoensis* is probably not a valid ssp., and the brown patch is large on p. 299 photo. P. 298, *Erigeron philadelphicus* should not be considered to be a host, as the larva probably wandered onto it, especially considering it pupated the next day and all larvae wander just before pupation. P. 299 I examined all the specimens that Porter & Mueller (J. Lepid. Soc. 52:182-205) used for electrophoresis in their paper (from Mich. and N Ohio), and found them all to be *P. cocyta*-group (see *Papilio* [N.S.] #13, and this was discussed briefly in a paper analyzing mtDNA of *Phyciodes* [Wahlberg, Oliviera, & Scott, Syst. Ent. 28:257-273], thus they proved only that *P. cocyta selenis* has mtDNA similar to *P. diminutor*). P. 300 all *Phyciodes* diapause in 4th stage. P. 301 both photos, again from "Madeira" (high altitude Madera Co. Cal. evidently, not Glenbrook Nev. as text implies with "TL Glenbrook...shown in the photographs") Jack N. Levy, are female (not male) *P. pulchella montana*; note that *montana* females usually have uph median band orange but yellower at rear like these two photos; real *P. oriseis herlani* females have this band all tawny (seldom all yellower like a few *montana*), & ups & uph is more uniformly orange with black lines (the "herlani" in Opler's field guide is evidently *montana* also). *P. mylitta* is commoner because it is a weedy multivoltine species. *P. oriseis*'s preference for riparian banks is only the raiting (perching) site for mate-locating males; the hosts are way up on hillsides etc. The uns of ssp. *herlani* is tawny with a network of fine lines; *mylitta* unh is mottled with brown areas more like ssp. *oriseis*. P. 302 lower photo evidently male. Variation last line should read SE individuals. *Melitiine* should be *Melitaeine*. Adult *barnesi* don't differ from *pallida*; the larvae are darker; larvae wander away before pupation so pupa would almost never be on host. P. 303 top photo is female not male; is there any data for *Centaurea diffusa* & *solstitialis*?, Shapiro's *solstitialis* record was merely "suspected"; Pelham told me that he gathered larvae on two occasions on *Centaurea* but believes they wandered from *Cirsium*, and he has never seep ovip. nor found eggs on *Centaurea*. (By the way, I caught the beautiful *Melitaea phoebe* in Spain, and read that its hostplants are 7 sp. of *Centaurea*, so we certainly must introduce it into Colo.-Mont.-Wash. etc. to help control *Centaurea diffusa* a bad Eurasian weed; Mongolian *phoebe* might have the right climatic adaptation for our area.) P. 304 pupa is evidently *E. chalcedona chalcedona* as it is from Josephine Co. Ore. P. 306 *ancia* TL not Banff; *howlandi* is a syn. of *ancia*; *veazieae*=*macyi* is a ssp. of *E. bernadetta*. Pelham notes that *bakeri* is a member of a guild with *hopfingeri* to N and *effi* to E (*hopfingeri* & *bakeri* are riparian and use more herbaceous *Penstemon*), and is assoc. with Snake R. & Columbia R. riparian habitats, as opposed to sandy habitats of *macyi* and lithosol habitats of *veazieae*. (But the adults and larvae and host of *effi* seem to place it as a synonym of *E. anicia maria*.) (Many ssp. in *Euphydryas* have geographic differences in host usage, and if the ssp. of *E. editha* were assigned by host use as found by the Ehrlich group, the ssp. would be totally different.) N. Kondla & S. Kohler in *Papilio* [N.S.] #12 showed that *E. bernadetta* is a distinct species (which includes *veazieae* and *macyi*), while Pelham has found that in hostplant use etc. *bakeri* seems to belong to *E. anicia*. P. 307 top right *veazieae* specimen looks aberrant with enlarged white spots, because none of my series of *veazieae* nor any book photo (or photos Robert E. Hardwick sent from Kittitas Co. and Klickitat Co. Wash.) looks like that photo; p. 401 states the photo is Harney Co. Ore., where such a specimen would be an aberration. Something interesting to consider: Dennis Murphy hybridized *chalcedona* and *ancia* for several generations and the results were similar to *veazieae*; maybe that says something as to its origin? The bottom sentences on p. 307 are false, Austin & Murphy 1998 actually described

and illustrated clear and obvious intergradation between *chalcedona* & *ancia* in Nev. and fully stated that it involved intergradation, and they treated *chalcedona* and *ancia* as one species (in NE Nevada, they are barely sympatric, but are mostly allochronic and occupy mostly different altitudes, yet in the few places and times where and when they do meet Murphy found that they interbreed more than randomly, thus there is no reproductive isolation at all)(later Austin changed his mind based on NW Nev.). Scott rehashed the *chalcedona*-*colon*-*ancia* intergradation vs. sympatry mess in *Papilio*(N.S.) #12. Note that Brussard et al. found *colon* more similar to *ancia* in its isozymes. P. 308 *chalcedona* wings are not more rounded. Pelham says Shepard was premature and *perdiccas* TL may be Vancouver I. Ssp. *wallacensis* from Missoula Mont. is same as *colon* except valva prong is intermediate to *ancia*. P. 309 *rassenkreis* & *wallacensis* misspelled. P. 310 *edithana* should be boldface. P. 311 the “baroni” pictures in Dornfeld’s book do look like *rubicunda* as Pyle says they must be (true *baroni* is a very distinctive local Mendocino Co. critter with much more red uns and the distal blue “editha line” often absent and ups often a bit darker)(by the way the blue “editha line” is not one line it is the “editha area”). Ssp. *mattooni* (not *mattoonorum* which is a *Hesperia* comma) and *owyheensis* are syns. of *edithana*. My series from Deschutes Co. Ore. appears identical to Calif. *nubigena*, thus *remingtoni* seems to be a syn. of *nubigena*. P. 312 Scot’s Broom is really Scotch Broom? Pl. 11 *paradoxa* female is male. P. 316 *neomarsyas* is syn. of *satyrus*. P. 317 ssp. *fulvescens* is a syn. of *rusticus*, as the Sierra uns is just as fulvous as Coast Range uns. True ssp. *rusticus* from Calif. has a lighter browner uns than *hylas* and the light-brown color is especially obvious in unf cell (Comstock’s uns from Plumas Co. is grayer than any other Cal. specimen I’ve seen), as brown as eastern ssp. *faunus* though the uns green spots are paler; the northern extent of *rusticus* is questionable, as Dornfeld wrote that Ore. adults have gray uns (maybe his desc. wasn’t accurate enough for this purpose?), though the p. 315 Clallam Co. male has brownish uns and Pelham states that NW males are fairly brownish (but has he seen Calif. *rusticus*?) and he is satisfied that NW pops. are *rusticus*, so maybe near-*rusticus* does extend north through Wash., but BC and Ida. have the gray-uns bug which I named *cenveray* in *Papilio* (N.S.) #12; someone should compare these with Calif. ones, and determine the limits of the browner *rusticus* (my 3 BC adults and the BC book photo are gray on uns like *cenveray*). Female *faunus* uns varies from unmarked (form *silvius*=*orpheus*) all the way to mottled almost as strongly as males, and the mottled ones are often mistaken for males. P. 318, I have never seen intermediates between *gracilis* & *progne*. P. g. *gracilis* is in E.N.A., not in Wash., though some specimens may look like it. We judge species by actual characters as our only estimate of “true genetic compatibility”; in this case there is no difference between *gracilis* and *zephyrus* except the contrast between uns base and distal area, and this contrast does intergrade on specimens we look at. True splitters take a nearly perfect cline like *leanira*-*cerrita*-*alma*-*pariaensis*-*fulvia*, and divide it by cutting the cline right down the middle in an orgy of pigeonholing that violates all fairness. P. 319 *P. oreas* and *P. progne* are separate species because no intermediates have been found (see #12 again). I haven’t seen Okanogan/Pend Oreille adults, but NW Mont. adults are the *threatfuli* type; *threatfuli* seems good in BC-NW Mont. (not quite a uniform entity), but when you add the SC Mont. bugs that are a little paler on uns (and this book says Blue/Wallowa/Ochoco adults are darker on uns), I think these are just degrees of intermediacy between *silenus* and *oreas*, so *threatfuli* is an intermediate therefore a synonym. I wonder how fast *silenus* really is, as *nigrozephyrus* isn’t fast. P. 320 *Aglais* seems a valid genus for *milberti* & *urticae* based on various characters and cladistic reasoning. *Roddia* seems to be the proper genus for *l-album*=*vaualbum*: in egg cluster size, *Roddia* has 18+ eggs, *Polygonia* 1-several, *Nymphalis* very large clusters; in larval horns *Roddia*’s are small (large *Polygonia*, absent *Nymphalis*); in pupal shape, *Roddia* is like *Nymphalis*; pupa of *Roddia* has silver spots in saddle like *Polygonia*, not *Nymphalis*; in genitalia, *Roddia* resembles *Nymphalis polychloros* but valva prong is a bit more pointed; in mtDNA *Roddia* is closer to *Polygonia* (work of Soren Nylin’s group in Sweden, Zool. J. Linn. Soc. 132:441-468); *Roddia unh* has a silver comma like *Polygonia*, not *Nymphalis*; *Roddia* are sexually dimorphic like *Polygonia*, not *Nymphalis*; wing shape of *Roddia* is like *Nymphalis*, not *Polygonia*; so the total score is 5 traits closer to *Polygonia*, 3 to *Nymphalis*, if you ignore the intermediate egg count and put the horn size into the *Polygonia* score, so I think we can conclude that *vau-album* is not a *Nymphalis*, so I call it *Roddia* (the alternative would be to lump *Polygonia* and *Roddia* into *Nymphalis*, the sister genus to *Aglais*). But now Wahlberg et al. have lumped it back into *Nymphalis* for unknown reasons; I call it *Roddia*. P. 321 *watsoni* is a syn. of *j-album*; *j-album* has larger uph submarginal yellowish spots than Eurasian *l-album*. P. 323 Red Admiral is used, but *Admirable* is used elsewhere in book. P. 324 *hyperborea* is a syn., as hinted. Eggs may be in double layer too. P. 326 *furcillata* & *subpallida* are syns. P. 327 the book missed “the Nut”, while quoting my original note in *Lepid. News*; the proper quote was: “And who was Milbert of *Nymphalis* fame, was he the brother of Filbert the Nut?”; we can’t leave out the Nut! P. 328 range is wrong, they must occupy the intervening white areas if they shift over the landscape as the book says. P. 329 is an unusual dark *virginiensis* larva, back east they are paler with rings around body. P. 330 *annabella* differs from *carye* by only a teeny tit on the valva, like *titania*/*grandis*, I caught them in Colombia and can’t tell the adults apart, so I lump them. P. 332 Australia has ssp. *kershawi*. Absent in most of S Amer. R. R. Baker found evidence of a little overwintering in Britain, and surely a little overwintering occurs in Colo. as T. Cockerell found one in Jan. P. 334 I don’t like the name *Admirable*; it’s a couple pages from *Limenitis* Admirals which have wide stripes. (*Atalanta* has *raiting*=perching mate-locating behavior which the ignorami call aggressive, so Redneck would be just as good.) P. 336 *grisea* is a syn., and its *unh* is tan to russet. P. 338 if Weidemeyer was German, it is pronounced Vye-day-my-air. P. 338 *fridayi* are real intergrades due to introgression: N Nev. has *weidemeyerii* and *fridayi* and no *lorquini*, Mono Lake & W Nev. has *fridayi* as *weid.* & *lorquini* almost interbreed randomly, & Dornfeld and this

book note *fridayi* in SE Ore. in the *weidemeyerii* pop.; only in Mont. does S. Kohler find reproductive isolation. P. 339 *arthemis* may have red spots on unh submargin, or a wider red band—this is a weak character. The bit about *arthemis* and *astyanax* being separate species was thoroughly disproved by Austin Platt, who in numerous quality papers amply demonstrated *astyanax* and *arthemis* are one species that intergrade fully in three genes that even change in different latitudes. Hybridization with *lorquini* in BC would logically favor more lumping, not more splitting. *L. arthemis* (with *astyanax*)-*weidemeyeri*-*lorquini* is a *stencospecies*. Suture zones of Remington were discredited in bird papers etc. 30 years ago. Pelham notes that there are a lot of hybrids of *archippus* X *lorquini* along the Snake River, and suggests that if one used hybridization to define species, there would be only one N.A. *Limenitis*. Looking at the big picture, *archippus* is quite distinct and I have never found a hybrid, whereas *arthemis/astyanax* broadly intergrade and have been thoroughly proven to be one species, and they are mostly allopatric to *weidemeyerii* and *lorquini*, except *weidemeyerii* overlaps *arthemis arizonensis* in Ariz.-N.M. (*weidemeyerii* X *arthemis* hybridize at Police Coulee in Alta. and in Sweetgrass Hills of N Mont., but not in 4 N.D. cos. where they are sympatric) so we can evidently treat *weidemeyerii* as a bookkeeping species. BC book says there are many hybrid *arthemis rubrofasciata* X *lorquini*, but are there enough to make them one species?, not according to the map dots, so we can treat *lorquini* as a mostly separate bookkeeping species from *arthemis*. P. 341 top two photos look like *ilgae* on unh, but locale says Pierce Co. Wash. well within range of *itelkae*, so are these ssp. weak? The the *burrisonii*-like hybrid in BC book looks like female not male. P. 342 intermediates without different characters of their own should not be named, so we should just call *idaho lahontani* X *archippus* intergrades if they tend toward *lahontani* a little, & *idaho* is a syn. of *archippus*. P. 343 how often does *archippus* eat apples?, not often, they haven't colonized Denver with its common apple trees and superabundant crabapple trees. Pelham states he has never observed *archippus* eating apple either, and *archippus* are not in apple orchards in S Okanogan Valley now. P. 346 *eryngii* is not really distinct from *california*, *columbiana* looks like a synonym of *ampelos* to me, the coastal *insulanus* is only a bit darker than *ampelos* on ups but unh is much darker than *ampelos* (Pelham notes *ampelos* is somewhat variable and occupies the steppe & semi-steppe), *swadneri* from NW Mont. is ochre-cream; *yontocket* is usually ochre above though it varies (and is mostly intermediate between *insulanus* and *california*, suggesting it cannot be a valid ssp.?, but series of *yontocket* seem to show an influx from *eunomia* also, plus some odd redder-unf adults, so *yontocket* can be kept for now since it is not quite intermediate). P. 347 Scot's Broom misspelled Scotch broom here. Most are not ochre so Ochre Ringlet is misleading; what's wrong with just simple Ringlet? It was Demorest Davenport—not Thomas Emmel—who showed that they all belong to *C. tullia*. P. 348 *boopis* TL Point Richmond Cal. is a good ssp. limited to Calif. as book says, it has only 0-1 unh ocelli, whereas Ottawa-Colo. *nephele* has the full set of unh ocelli; a bunch from Ochoco Mts. have lots of ocelli so look like *nephele*, but many others in my coll. from Ida.-Wash.-Mont. have varying numbers of ocelli from few to many and perhaps could be called near *nephele* maybe; *ariane* and *incana* are near *nephele*; *paucilineatus* is basically a syn. of *gabbii*. A huge statistical study of geographic variation would be useful. Lots of hosts in *Papilio* [N.S.] #6 never made it into this book. P. 350 Great Basin Wood Nymph isn't a good name because it occurs in Calif. P. 351, *phocus* (TL Lac la Hache BC) is a syn. of *oetus*; in BC book it is said to have the pattern of *oetus* but the unh smoothly dark-brown, but what are the odds of their ranges changing exactly all along the BC-Wash. border as the BC book map shows?, zero!, so it would seem that they cannot be distinguished by actual characters and are synonyms. But Pelham suggests that some boreal pops. in N Cascades have suppressed unh markings and smooth ground color, if *phocus* applies to those; however he notes *oetus* is extremely variable. P. 352 lower photo of "incana" is same photo of "ariane" on p. 249. P. 353 photo looks like *C. palla* more than *hoffmanni*, and maps of *Chlosyne* and *episodea* show it more likely to be *palla* than *sterope* or *hoffmanni*. P. 354 TL Rock Lake, Alta., N. Kondla 1996 Alta. Natur. 26:40. Bean did not send the types to Edwards because the bug was named 20 years before Bean collected; Kondla is looking into this. *Hopfingeri* is syn. of *episodea*. P. 356 larva is 3rd or 4th-stage *O. calais valerata*. P. 356 *O. uhleri uhleri* probably is annual in Colo. foothills, as I haven't noticed differences between years (it's possible that two cohorts would produce a population every year, but *uhleri* larvae grow on grasses on open S-facing slopes that are warm, versus sedges in the shade of trees for Front Range Colo. *O. chryxus*). P. 357 the bit about the king's new clothes is amusing. Actually *wyomingo* in E edge Rocky Mtn. Nat. Park occurs there alone without any *ridingsii* flying earlier in the summer, so can't be just the second generation. Scott & Fisher show (*Papilio* [N.S.] #18) that *wyomingo* is a late-flying ssp. that has color of *ridingsii* & pattern lines of *stretchi* & *raits* in swales. (In Spring Mts. Nev., there are two cohorts of *Euphilotes ancilla purpura* that are sometimes sympatric but always allochronic by a month, that do not differ by mating site or diapause stage, and feed on different varieties of *E. umbellatum* [Austin et al. 2008 (J. Lepid. Soc. 62:148-60) named it a new ssp. *cryptica* but it's a syn. of *purpura* to me], and a certain lepidopterist told me that they must be two separate species because he insists sympatric allochronic entities have to be separate species; in that case Pyle might say that there are not only no clothes, but no emperor)(actually I feel naked without my net, and the butterfly watchers trade their net and any possibility of certain identification for binoculars, so I think they are the ones running around naked). *Wyomingo* hibernates as young larva. P. 357 only 2 sites in text, 3 counties on map, so one site is on county line? P. 358 Vancouver I. missed on map. P. 359 top is confusing as it is *O. nevadensis*, and isn't labeled Great Arctic like bottom photo is; they do not overlap at all & may be ssp. P. 360 *beani* is blacker than other ssp. P. 361 *valerata* is very variable and a ssp. of *O. calais*, and the Mt. Hull stuff might be *O. calais altacordillera* (see *Papilio* [N.S.] #12). P. 365 Bt corn has proven to be not an important threat. Since the monarchs mate randomly by male capturing female in S

Mexico in Feb.-March, the SW-fall-flying ones out of E U.S. would interbreed with the SE-fall-flying ones out of Wash., canceling out their adaptation, wouldn't they? P. 396, JAL is JNL=Jack N. Levy. P. 397, 72:bottom no data.

THE BUTTERFLIES OF THE PACIFIC NORTHWEST. James R. Christensen.

1981. University Press of Idaho. 116 p.

This book is old and insignificant, but many misidentified photos should be noted: Pl. 2, 13, 2nd not 1st brood, 19 E. h. lotta. Pl. 9, 65 & 64 are *P. gracilis zephyrus*. Pl. 11, 70 *S. coronis simaetha*, 72 *S. hydaspe purpurascens*, 70 elaine. Pl. 13, 90 *P. cocyta*. Pl. 14 & 15, 97 *C. acastus* I think, 96 E. editha both plates. Pl. 16, 109 L. c. snowi, 110 L. p. arethusa, 112 could be xanthoides X editha. Pl. 17, 112 could be xanthoides X editha, 119 cupreus. Pl. 18, 117 male-female switched. Pl. 19, 117 idas anna, 130 E. amyntula. Pl. 21, columns reversed. Pl. 27, #28 & #30 switched. Pl. 30, 16 & 17 switched. Pl. 43, 90 *P. cocyta*. Pl. 44, 109 cupreus. Pl. 45, 112, xanthoides X editha. Pl. 47, 129 *S. sylvinus*, 147 mossii. Pl. 48, 164 centaureae, 181 H. comma idaho.

WATCHING WASHINGTON BUTTERFLIES. R. M. Pyle. 1974.

Seattle Audubon Society, Seattle, Washington. 109 p.

Misidentified Photos: Fig. 7, *Euchloe hyantis lotta*, 12 *Colias philodice*, 15 *Satyrium sylvinus nootka*, 30-31 *Cupido amyntula*, 40-41 *Polygonia faunus*, 64 *Erynnis persius* (more likely than *pacuvius* or *proportius*).

BUTTERFLIES OF OREGON. THEIR TAXONOMY, DISTRIBUTION, AND BIOLOGY.

Andrew D. Warren. 2005. Contributions Gillette Museum of Arthropod Diversity, Colorado State University, Fort Collins, Colorado. 408 p.

This is a good contribution to the taxonomy of Oregon butterflies, aimed at scientist-lepidopterists. It is mostly concerned with studying the geographic variation and distribution of the various species and subspecies. Warren notes that there have been numerous misidentifications of Ore. butterflies, so he mostly rejects other people's identifications and uses mostly his own identifications. There are no illustrations of the butterflies, so it is not possible to check his identification ability here, but it is quite good (though I disagree with the sex or species of eight of his corrections of identifications of photos in Bob Pyle's Butt. Cascadia book, as noted above). The book also has a useful massive compilation of citations of the biological studies conducted on Oregon butterflies anywhere in or near Oregon, no doubt assisted by the Pelham Catalogue bibliography. In contrast to the rejection of others' identifications, Warren accepts a lot of hostplant data & studies from Jonathan Pelham, even though none of that data has been published thus cannot be verified. The altitudes and flight times are combined & summarized for the whole state, which isn't too useful when there is such great variation between the climate of rainy coast, desert, and alpine. The use of the terms "guarding perches", "defending perches", "dueling", etc. for mate-locating behavior are rather annoying anthropomorphisms to me. P. 106 claims that *Lycaena heteronea* "frantically patrol between flower heads of their *Eriogonum* foodplants, and defend perches on *Eriogonum* flowers.", which is self-contradictory in mate-locating terminology (the bug flecks over the hosts in Colo.). (Raiting & flecking are now used for perching & patrolling mate-locating behavior, see *Papilio* [N. S.] #14.) Scott (1992, *Papilio* [N. S.] #6) reported life histories of many species that were not cited in the book (*Pyrgus communis*, *Pholisora catullus*, *Papilio zelicaon*, *Pontia beckerii*, *protodice*, *Pieris "napi"*, *Lycaena helloides*, *Plebejus melissa*, etc.), maybe because they were far from Ore. Warren does not like subspecies, yet still uses them throughout in discussing geographic variation. Many of the ssp. discussed I think are synonyms, as discussed in the review of Syst. W N.A. Butt. above, so I have not repeated all of those here.

An Ore. curiosity is Elmer Griepentrog, who evidently singlehandedly mislabeled a thousand specimens, resulting in dozens of erroneous records noted in the book. One would think that someone with that name was invented by mischievous Entomology I students at Oregon State, but Michael Fisher told me that Elmer was a real collector who lived in Seaside, Ore., and exchanged butterflies with him in the late 1960s, and that material was identified correctly to species. Evidently something later happened with Elmer.

Details: Male *E. proportius* rait on gulch banks and just off the top of hilltops. *Ricara* isn't a very good ssp. of *Pyrgus ruralis*. Scott's paper on *Hesperia* (*uncas*, *comma colorado*, *comma ochracea*) immatures was missed (*J. Lepid. Soc.* 29:163-7, 1975). Warren documents migration of *Hesperia juba* and refutes adult overwintering, which Scott also refuted in *Papilio* (N.S.) #6 & in *Lepid. News* 1982#1 p. 5. *H. comma oregonia* is a variable worthless ssp. name. Pyle's Butt. Cascadia p. 81 lower left *peckius* actually is a male. *Alkaliensis* is a syn. of *Polites s. sabuleti*, my series from Lake Co. Ore. is clearly ssp. *sabuleti*, and Austin (1987 also mapped ssp. *sabuleti* from NW & N-C Nevada (near Ore.)). The *Polites "mardon"* in Butt. Cascadia p. 89 lower right, Warren says is actually a female, while I think it is *O. sylvanoides orecoasta* female. *Lutea* may be a good ssp. of *O. yuma* with slightly yellower unh. Male *yuma* in N.M. that have strayed from the *Phragmites* patch also rait near the ground on river banks. *Osceola* & *kiowa* are syns. of *Euphyes vestris*. *Cynodon* & *Poa pratensis* are errors for *Amblyscirtes vialis*.

The ssp. in *Parnassius clodius* are mostly useless. *Magnus* & *xanthus* are syns. of *Parnassius phoebus smintheus*. May to Aug. is several generations of *Papilio multicaudatus*. *Neominois menapia tau* from Wash. actually is a good ssp. with darker apical black upf marks, similar but not as black as *N. m. melanica* from coastal Calif.; red unh markings are variable and haven't been shown to have much geographic variation thus far. *Elivata* is a syn. of *sisymbrii*. *Transmontana* & *mayi* are syns. of *coloradensis*, which isn't much different from *Euchloe ausonides*. Ken Davenport has evidently found intermediates between *Euchloe hyantis* & *E. h. lotta* on the Kern Plateau. The black upf bar and uph black dots should have been mentioned in the treatment of *Anthocharis "sara"*. Ssp. *stella* has yellowish males, so the Ore. bugs aren't typical *stella*. *Colias philodice eriphyle* is limited to BC pops. with some orangish-laved adults. If *Colias eurytheme* is migratory, why have I never seen migrants in Colo.? The treatment of *Colias occidentalis* & *christina* is good, except that they are obviously conspecific. The *P. sennae* female in Butt. Cascadia looks too yellow to be *philea* to me.

The ssp. of *Lycaena arota* are defined wrongly here and will have to be restudied: ssp. *arota* actually has whiter unh & the male unf is just ochre-tan with no orangish; *virginiensis*=*schellbachi* has brown unh & postmedian-submarginal orange on male unf; the male upf margins don't differ. The *Lycaena cupreus* ssp. are messed up also, as Scott (*Papilio* [N. S.] #12 wrote: SE Ore. evidently has ssp. *artemisia*, while SW Ore. evidently has near-*cupreus*. Warren notes intermediates between *L. xanthoides xanthoides* and *L. x. editha* in Ore.; Pratt et al. 1993 actually could not even distinguish *L. rubidus* larvae from *xanthoides-editha*; Scott in *Papilio* [N.S.] #14 p. 35-36 discusses *xanthoides-editha* further. The heavily-spotted unh *Lycaena heteronea* from N Cal. & S Ore. are *gravenotata*, as they look the same and there are no other differences; excuses such as "most likely does not indicate close genetic similarity" are just hot air in the absence of data, as we have to go on known facts to decide, not speculations, and the facts say they are *gravenotata*; and the primitive condition throughout *Lycaena* is a fully-dotted unh, so it's the spotless Great Basin types that are genetically different; above I dealt with lousy names *submaculata* & *rava* etc. *Penrosae* is a syn. of *L. mariposa*. *Habrodais grunus* males don't "guard perches", they fleek (patrol to seek females) late in afternoon as the previous sentence on p. 113 says.

P. 115, a mile away is NOT sympatry (close but no cigar, folks), and *perplexa* is a ssp. of *C. affinis* (and the *perplexa*-clone *homoperplexa* from Colo. intergrades with *affinis* in S Wyo. & Neb. says G. Gorelick). P. 116, Scott's 1986 book reported that *viridis* is a syn. of "dumetorum" but the latter will no doubt be banished by ICZN under its plenary powers. P. 117, only one of Scott's specimens from near Satus Pass was misidentified for certain; and Warren discusses the problems of distinguishing *sheridanii* from *perplexa* & *affinis* in Ore., so one supposes that many specimens were misidentified in Warren's study samples also. Glenn Gorelick wrote that *pseudodumetorum* (from Stanislaus Co.; my series from there fits the O.D. description of specimens from Trinity Co. TL) is related to *viridis*, and Warren seems to show that it belongs to *C. viridis* (as *sheridanii*) (near *lemberti*) (Ken Hansen found that it is sympatric with *perplexa* in Trinity Co., and Univ. Colo. museum specimens show *perplexa* sympatric with *pseudodumetorum* in Del Puerto Can. in Stanislaus Co., so can't belong to that), and so it seems that *viridis* is conspecific with *C. sheridanii* (*dumetorum* is older than *sheridanii*, but I have petitioned ICZN to suppress it and protect *sheridanii* from *viridis*). My reared toptype *newcomeri* and nearby specimens are small with a thin white unh line, so I have difficulty believing that populations in Polk & Josephine Cos. in W Ore. that are nearly immaculate can be intermediate *newcomeri*X*viridis* as middle of p. 121 & top of p. 122 says (Warren notes that true *newcomeri* are often large, females are often tawny, and uns is somewhat bluish-green, but do they have few white spots?), maybe the W Ore. bugs are better called *newcomeri*X*lemberti*, as *lemberti* averages fewer spots (the next sentence on p. 121 calls them near-*viridis*, which is even more dubious). P. 123 claims that *interrupta* is similar to *neoperplexa*, which is also dubious, as the unh line is variable and sparse in the former, complete in the latter. P. 124 then claims that *pseudodumetorum* could be treated as a syn. of *newcomeri*, which is quite wrong, and the same paragraph claims that *lemberti* is broadly similar to *neoperplexa*, which is again wrong. We need more study of larvae of these bugs, which show good differences (Ballmer & Pratt in *J. Res. Lepid.* 27:40 says that *comstocki*, *viridis* "dumetorum", & *lemberti* have saw-tooth-profile big ridges on most older larvae, whereas *perplexa* is more rounded), and less study of adults which in Ore. can often not even be identified. Anyway, *viridis* "dumetorum" and *pseudodumetorum* are not *newcomeri*, and this book's *Callophrys* section suffers from sloppy writing. But it does seem that Warren has shown that *viridis* "dumetorum" and *sheridanii* (and *lemberti* & *interrupta*) are likely all one species, so we do need the ICZN to suppress the name *dumetorum*, as *dumetorum* has the awful history of stomping on *viridis* and causing vast confusion of *viridis* vs. *dumetorum* and *dumetorum* vs. *perplexa*, and now causing even more confusion of *dumetorum* vs. *sheridanii*; the disgusting name *dumetorum* must be banished forever! And *sheridanii* must be given priority over *viridis*.

The treatment of *Callophrys gryneus* in this book is excellent, as Warren corrects a lot of errors, and finds that the type of *barryi* was mislabeled and was from a *Thuja* pop. near Portland, so he synonymizes it to *plicataria*, & finds that *plicataria* & *barryi* & *byrnei* are all syns. of *rosneri*, and synonymizes *acuminata* to *nelsoni*, & redemonstrates my 1986 book finding that the *gryneus* ssp. (incl. *siva*, *nelsoni*, *loki*, etc.) can switch hosts in nature and they all eat each other's hosts in lab, and Warren shows that my book's lumping of all these into one sp. *gryneus* was correct. *Ninus* is a syn. of *C. spinetorum*. The book missed my note that *Callophrys johnsoni* flies down-gulch to find moisture (Scott 1973b, *J. Lepid. Soc.* 27:283-7). I have 3 *Callophrys mossii* from Morrow Co. that are *mossii*X*schryveri*. *Obscurus* is a syn. of *polios*, and *maritima* doesn't look different either. My 1986 book described *C. polios* larva & pupa. *Purpurascens* is a syn. of *sheltonensis*, a good ssp. with darker purple uns. The book separates *Satyrium fuliginosa* & *S. semiluna* as distinct species, but does not give

supporting information, as they are not sympatric, supposed differences in structure and wing shape are not presented, and high-Cascades pops. are not on ridges & have vestigial stigma intermediate between the species (& the book claims that Dornfeld's 1980 p. 196 #4c-d female is *S. semiluna* but it looks exactly like *S. fulig. tildeni* and nothing like *semiluna* to me). *Maculadistinctum* is a syn. of *semiluna*. *Columbia* is a syn. of *Satyrium behrii*. *Occidentalis* doesn't seem much different from *Satyrium titus watsoni=winteri*. *Obscurafacies* is basically a syn. of *Sat. californica*, as is *cygnus*. Scott 1992 showed that *S. californica* is polyphagous on many shrubs, so maybe Dyar's 1894 Yosemite rearing on *Salix* was *californica*. Pyle's Butt. Cascadia p.197 upper right photo actually is a female. *Okanogana* is a syn. of *Satyrium saepium*. *Setonia* is a syn. of *atrofasciata*, *pudica* a syn. of *Strymon melinus*, whose males have orange abdomens after spring, and which doubtfully immigrates.

Cupido comyntas in Ore. are not *sissona*, which was explained in *Papilio* (New Series) #12 with the description of *C. c. pacnowe*. Wright (1885) reared *C. amyntula* from inside *Astragalus* pods in Calif. The treatment of *Celastrina echo* also suggests there is a 2nd species *Cel. nigrescens* over much of Ore. (*nigrescens* with small, shorter wings [vs. larger, longer wings in *echo*], hazy dusky blue ups. [vs. darker clear purplish-blue in *echo*], unh variable but most have larger darker spots [vs. poorly-maculated clear-white, often bluish unh], fringes strongly checkered [vs. white]). But I do not understand the difference between them, as *nigrescens* has violet ups in BC also. Rearing from natural hosts is needed (but beware, my adults of *Cel. neglecta* reared from *Prunus virginiana* leaf galls in Minn. were much more silvery than wild adults). *Echo* is supposedly assoc. with *Ceanothus velutinus*, *nigrescens* with *Holodiscus*, but I wonder if those are real main hosts, or just possible occasional hosts? (*Celastrina* taxa such as *sidara* & *ladon* & *neglecta* may have main hosts and lots of secondary hosts). Much more work needed here; based on maculation & fringe is *nigrescens* the S ssp. of *Cel. lucia*? *Toxeuma* is a syn. of *Glaucop. piasus daunia*. *Leona* is obviously a ssp. of *Philotiella speciosa*, and maybe a syn. of *bohartorum*; *septentrionalis* is a syn. of *speciosa*. The *Euphilotes* on *Eriog. marifolium* (fig. on front book cover) is *E. battoides* ssp. near *battoides* (which also eats *E. marifolium*). Warren raises *E. glaucon* on *Eriog. umbellatum* (with *oregonensis* & *intermedia*) to bookkeeping species status, within stenochospecies *E. battoides*. He treats the similar bug on *E. heracleoides* as a separate species having paler uns & later flight, but it seems to be a new ssp. of *E. glaucon*, which has a couple apparent relatives that are "sympatric or narrowly parapatric [apparently the latter, as proof is not given of the former]". *Euphilotes "baueri"* seems to be an unnatural catchall mess. Evidently *E. columbiae* should be called *E. ancilla columbiae*? *Plebejus "idas"* in Ore. I treated as *Pleb. atrapraetextus* in *Papilio* (N.S.) #12; the Ochocos to Wallowas pops. are *P. atrap. atrap. Xlonginus*, and I named the Warner Mts. bug *P. atrap. benwarner*. In Colo., male *P. atrap. sublivens* and *P. melissa* often rait (perch to await females) on little trail-like depressions in valley bottoms. The BC book statement that *idas* feed only on *Vaccinium* was an error. *Ricei* is a syn. of *P. anna anna*; the amount of uns and female ups orange varies greatly in these. Both of Shapiro's records of host *Astragalus whitneyi* var. *siskiyouensis* on p. 189 top (for *P. anna*) & also p. 190 top (for *P. melissa*) were for the same Ball Mtn. bug that later proved to be *P. atrap. benwarner*, not *P. anna* or *P. melissa* (see *Papilio* [N.S.] #12). Real *P. melissa* in Colo. was NOT the high-altitude bug with reduced orange (an error by F. M. Brown, which I fixed in #12). *Glycyrrhiza lepidota* is a dubious host for *P. melissa*; it is ignored in Colo. Austin's (1998i) conclusions on *Plebejus saepiolus aehaja* & *rufescens* were wrong, and *insulanus* is a syn. of *amica*, as I explained in *Papilio* (New Series) #12; evidently Oregon has ssp. *saepiolus* over most of the state, and weak ssp. *littoralis* (which actually has smaller uns spots with white rings) on the SW coast. Colo. Front Range *P. s. near-gertschi* eat *Astragalus* (Scott 1992). *P. 193, fenderi* does not resemble *pardalis*. *Montis* is a syn. of *Pleb. icarioides pembina*. Pyle's Butt. Cascadia p. 250 photo actually looks most like a male *Pleb. melissa*. *Calchas* is a syn. of *Plebejus shasta*, & *pallidissima* is a syn. of *minnehaha*. *P. 200 top, Pelham's bivoltine P. alupini* in Wash. may refer to what he now considers to be *P. acmon*, which is bivoltine there (at the Chelan site Scott 1998b referred to, etc.). *P. 201 middle, real lutzi* doesn't have large uns spots. *P. alupini spengelatus* also occurs in Alta. (*Papilio* [N.S.] #12), so may occur in high Wallowa mts. too. The *Plebejus* on *Eriog. pyrolifolium* in the high Cascades from Mt. Shasta to Mt. Adams is evidently a nice distinctive unnamed ssp. of *P. alupini*, which is sympatric with *P. acmon* but not with other *P. alupini* varieties.

Scott (1986) book had photos of *Euptoietia* immatures. *P. 207, Pyle's pugetensis* photo was on p. 265, not 256. *Speyeria zerene hippolyta* differ from *bremneri* on ups median band & unh disc & submargin. Scott's 1986 book lumped *S. z. garretti* into *picta*, following Grey & Moeck 1962. NE Ore. pops. are evidently *S. egleis linda*. Pyle's p. 273 lower right photo is a male *S. hydaspe*, not *zerene*. *Dodgei* is a syn. of *S. hesperis cottlei* (in turn a weak variety of *irene*). *Speyeria atlantis* was a mess until Scott et al. (1998) fixed it & divided it into two sp. & named 3 ssp. Study of the types clearly indicates that *irene* is NOT a syn. of *egleis*, as noted in Butt. Cascadia review. Pyle's p. 278 upper left fig. is *S. coronis* I think. All those *Boloria selene* ssp. except *atrocostalis* are very weak. The *B. bellona* ssp. are very dubious; Scott 1992 showed a partial 2nd gen. in Colo. *Uslui* & *borealis* are syns. of *B. epithore chermocki*. *Oregonensis* is evidently a syn. of *Chlosyne leanira*, & *basinensis* a syn. of *alma*. Scott 1992 found "basinensis" on *Castilleja chromosa*. *Chlosyne palla* ssp. *eremita-palla-sierra* is a cline, and only 2 names can be used for a cline, so *eremita* has to be considered a syn. of *palla*. The treatment of *Chlosyne palla* & *acastus* is good, with J. Pelham's finding that *sterope* belongs to *C. acastus* (which becomes *C. sterope acastus*). *P. 231, Ferris 1981b dorsal female* at far right is on p. 324. *Manchada* is a syn. of *C. hoffmanni segregata*. *Hoffmanni's "patrol territories"* evidently means flaiting behavior? I add two more misidentifications of *Phyciodes orseis* in Pyle's Butt. Cascadia: the middle left male on p. 297 is *P. mylitta*, and BOTH females on Pyle's p. 301

are *P. pulchella montana*. P. 238, Aster “*Eurybia*” *glaucodes* is the host of *Phyc. batesii anasazi*, which is not conspecific with *P. cocyta* as this page claims (see *Papilio* [N.S.] #13). P. 241, *owimba* & *inornatus* are not synonymous, the antenna is orange in former & black in latter, & latter is an invalid intergrade anyway (see *Papilio* [N.S.] #13). Jack Harry gave lots of hosts of *Euphydryas gillettii* in Scott’s 1986 book. P. 244 notes apparent intermediates between *E. anicia bakeri* and what is now considered a separate sp. *E. bernadetta veazieae* along Snake River in Baker Co. Scott 1981a found no difference between *colon* & *chalcedona* valvae. SE Jackson, S Klamath & S Lake Cos. seem to have *E. chalcedona colon* with mixture of *macglashanii*; Austin & Murphy showed *macglashanii* is extra-creamy on ups, with little extra red. Pyle’s Butt. Cascadia p. 312 upper right photo looks like a male to me. *E. editha baroni* is a local Mendocino Co. Calif. bug with extreme red uns. *Polygonia faunus rusticus* has a brownish uns, and evidently does not extend northward into Ore., while *fulvescens* is doubtfully much different from *rusticus* in nature (most known specimens are reared). The *zephyrus* fig. by Guppy & Shepard p.252 looks like a female in wing colors & shape to me. *P. oreas* and *P. progne* are closest relatives in uns color, hosts, larvae, rarity, weak flight, & gnathos (see *Papilio* [N.S.] #12), despite mtDNA (*progne* evidently stole its mtDNA from *P. interrogationis/P. comma*) which has been proven recently to be rather useless for the study of phylogeny. SW Ore. pops. are *P. oreas near-silenus*, based on my Siskiyou Co. series. *Threatfuli* is not intermediate *silenus*X*nigrozephyrus*, they are intermediate *silenus*X*oreas*, nearer *silenus*. *Nymphalis milberti* migrates altitudinally. Pre-hibernation Sept. *Nymphalis antiopa* frequently feed on *Chrysothamnus* flowers in Colo. *Vanessa cardui kershawi* is geographic variation. Scott (J. Res. Lepid.) showed that spring *cardui* migrants into Colo. fly to NNE all day long, so how does *cardui* get to Ore.? Scott’s 1986 book figured the *Vanessa virginiensis* larva. Idaho is basically a syn. of *Limenitis archippus*. Warren indicates that *Adelpha eulalia* is a distinct species, but provides no evidence that *californica* is a separate sp. from *A. bredowii*, so it’s *A. bredowii californica* perhaps.

Only half of my *Coen. tullia ampelos* have one ocelli (on rear of unh). *Eryngii* is a syn. of *C. t. californica* (a slight intergrade thus invalid). *Yontocket* isn’t similar to *insulanus* (which is like *ampelos*), it looks like *california*X*ampelos* then adding strong uns markings. *Ariane* & *incana* are synonyms of *nephele*, *paucilineatus* is a syn. of *gabbii*. Scott 1992 reported that *Cerc. pegala* do NOT drop their eggs in flight, they always land and extrude an egg, and half the time the egg falls parabolically into the litter and sticks to the first thing it hits (the other half the egg sticks to the plant the abdomen touches). *Erebia epipsodea* has no valid ssp. *Neominois* mating period is early-to-mid morning, so the afternoon patrollers must have been flying for another reason such as predator avoidance.

The references are 99 pages!, and with their info cited in text make the book useful. (There is a space between entries and the type is large, so with hanging paragraphs and smaller type they could have been put into 50 p. to save \$) P. 292, it’s J. V. Z. Brower. P. 296, move *Chermock* & *Frechin* to p. 297. P. 299 spelled *Cockerell*. P. 358, add Scott, J. A. 1975. Early stages of seven Colorado *Hesperia*. J. Lepid. Soc. 29:163-7.

THE BUTTERFLIES OF OREGON. Ernst J. Dornfeld. 1980.

Timber Press, Forest Grove, Oregon. 276 p.

Warren (above) corrected this book, so I won’t repeat most of that, and won’t repeat the ssp. corrections already noted. It was a good book for its time, but is somewhat out of date as Warren notes.

P. 40 *oregonia* is obviously *P. machaon oregonia*, & has 2-3 gen. *Maple* is dubious for *Papilio glaucus rutulus*. *Parnassius hibernate* as eggs. *Neophasia men. menapia* occurs in Ore. *Limenitis lorquini* doesn’t occur in Wyo. or Alta. *Polygonia oreas* don’t eat *azalea* & *rhododendrons*. *Castilleja* is an error for *Chlosyne palla*. *C. palla* & *C. sterope* & *Callophrys* etc. were updated by Warren. *Euph. editha baroni* is limited to Mendocino Co. Calif. Adult *Call. eryphon*--not larvae--were on willow catkins. *Polygonum* is an error for *Lycaena mariposa*. *Lycaena helloides* hibernates as eggs. The gnathos illustrated for *Plebejus melissa* looks like “*idas*”, and the treatment includes *Pleb. atrapraetextus* (maybe some photos too). *Avena* is a lab food only of *Amblyscirtes vialis*. *Symphoricarpus* & *Ambosia* are errors for *Pholisora catullus*, *Amaranthus* an error for *Heliopetes ericetorum*, *Sidalcea* an error for *Pyrgus ruralis*, *Sitanion* an error for *Thorybes mexicana*. *Th. diversus* does not occur in Colo. or Wyo.

I list some of the misidentified photos here (the major corrections--very many ssp. names have to be updated due to progress, & the sex is wrong on a dozen as Warren notes). Pl. 4#1a-c are *N. m. menapia*. Pl. 6#4b-c-5a-c are *Col. occidentalis occidentalis*X*wasatchia* females. Pl. 10#4a-c are *Cerc. pegala nephele*. Pl. 17#4c is probably *Polyg. faunus*. Pl. 18#1a-d *Phyc. cocyta selenis*. Pl. 19#5a-c is *Chl. sterope sterope*. Pl. 22#2a-c are evidently *E. editha rubicunda*. Pl. 34#4a-b is *Sat. (fuliginosa?) semiluna*. Pl. 36#8a-d *Call. [viridis] sheridanii newcomeri*. Pl. 39#2a-d *Lyc. nivalis browni*. Pl. 40#2a-d *Pleb. anna anna*. Pl. 41#5a-d *Cupido comyntas pacnowe*. Pl. 43#4a-d *Och. sylvanoides orecoasta*. Pl. 46#3a-d are *Hesperopsis libya near-libya*. Pl. 48#2a-c are *Th. mexicana aemilia*.

The maps need correcting in recently-revised groups such as *Colias occidentalis/edwardsii*, *Chlosyne palla/sterope/acastus*, *Plebejus acmon/alupini*, *Euphilotes*, *Callophrys [viridis] sheridanii/perplexa*, *C. gryneus*, etc.

ALBERTA BUTTERFLIES. C. Bird, G. Hilchie, N. Kondla, E. Pike, & F. Sperling. 1995. Provincial Museum of Alberta. Edmonton, Alberta. 349 p.

This book has a nice appearance, with glossy pages and pretty maps and photos (the little North American maps were taken from my book). Unfortunately, the book has many errors. The text appears to have been written a decade prior and poorly updated since. Evidently the book was written mostly by Charles Bird, a long-retired botanist rather than a lepidopterist; p. 2 suggests that Albert Finnamore helped mix up some of the photographs. Gross misidentifications abound in the photos. Dozens of photos are misidentified as to sex, including many not corrected in the errata. Surprisingly little new biological information such as hostplants is presented, and numerous hostplant errors from the literature are repeated. Species are arranged alphabetically rather than by taxonomic similarity. Many good common names are ignored and bad ones used. The little maps showing the North American range are lifted directly from Scott's (1986) *Butterflies of North America*, frequently without adjusting for changed species concepts. The dot maps seem fine, but the purple mountainous area is too small on all dot maps so a quarter of the dots for alpine species are mistakenly outside the purple area.

Detailed comments on each species: *Bouteloua curtipendula* occurs N to Mont. at least, and is probably the host of *Amblyscirtes oslari* in Alta. *Amblyscirtes vialis* doubtfully eats *Poa* since its hosts are hay-type-grasses, my larvae died eating it, and *Avena* is a lab food only. *Ancyloxypha numitor* doesn't eat *Poa* for the same reason; Minn. females fly right over *Poa* lawns to oviposit on hay grasses beyond. Scott (1992, *Papilio* New Series #6) paper clearly showed that *logan* belongs in *Anatrytone*, whereas *Atrytone arogos* is vastly different. *Schizachyrium* is an error for *A. logan*, as the original host record merely stated "*Andropogon*" and only *A. scoparius* is *Schizachyrium*. The photo of male *Hesperia comma manitoba* is rather pale on uns & looks like *assiniboia*. The TL of *Hesperia comma manitoba* is Lac la Hache B.C., not Man. as the inappropriate name implies. There is no proof that *H. comma* contains more than one species in Alta.; in Colo. we have proof that the *manitoba*-like *colorado* is the same ssp. as the lowland *ochracea* that looks like *assiniboia*. The photo of *Hesperia nevada* female is *H. comma* female. *Oarisma garita* males fleek, they do not perch or defend territories. *Poa* is dubious for *Poanes hobomok*; photo of female is *Polites peckius*. The *Polites rhesus* male photo is female. The *Carterocephalus palaemon* photo is ssp. *magnus*, not *mandan*, according to N. Kondla; *Poa* is a lab host only, *palaemon* would probably eat *Bromus inermis* but doubtfully *Bromus tectorum* which is not a suitable hay grass. Presumably *Erynnis afranius* occurs in Alta. *Thermopsis rhombifolia* was just a "likely" host of *afranius* in N.D., not a confirmed host. *Erynnis persius fredericki* was named after R. C. Frederick not R. C. Fredericki, and this ssp. is doubtfully distinguishable from *borealis* or *persius*. *Malva* is the same old error for *Pholisora catullus*. The text says *Pyrgus centaureae freija* is darker than *loki*, but the photos look the same. Scott never observed oviposition on *Potentilla diversifolia* in Colo., only on *Vaccinium* near it, and Scott's recent research shows that lab larvae eat *Vaccinium*, as well as *Potentilla* and *Fragaria* etc.--another example of Scott's law that all bog butterflies are polyphagous. *Pyrgus communis* has only 2 gen. on the Colo. plains, so has two in Alta. The *Pyrgus scriptura* photos have black streaks extending to edge of the hindwing fringe, and the unh is darker (both traits like *Pyrgus xanthus*), so are oddly atypical—why? *Thorybes pylades* pupae do not overwinter. Several populations of *Parnassius phoebus* occur in Colo. on hills at the plains edge; and males fly only 1/4 m above ground. Shepard's claim that *Parnassius phoebus* and *smintheus* are separate species lacks proof, as the primitive egg of *smintheus* matches the European egg, and several of Shepard's "Alaskan" traits also occur in SW Colo. *pseudorotgeri* (occurrence in moist swales on *Sedum rosea*, dark submarginal band on fw), and the micropyle photos are rather similar with a magnifying glass. *Papilio machaon dodi* is a syn. of *brucei*. Two of my maps for *machaon* are mislabeled. Oakley Shields (J. Res. Lepid. 6:69) reported lots of mating behavior of *Papilio zelicaon*. The book separates *Papilio glaucus* and *canadensis* and *rutilus* wrongly, as Hagen et al.'s reasons for separating *glaucus* from *canadensis* are very weak, they intergrade over a broad zone of a hundred miles or more, and *canadensis* and *rutilus* intergrade broadly in SE B.C. and Mont. (J. Scott and J. Shepard, *Pan-Pacific Ent.* 52:23-28, and the Black Hills (L. Brower, *Evolution* 13:40-63); even in Colo., *glaucus/rutilus* intergrades are found along with a few stray *glaucus*; there is little or no reproductive isolation within the *glaucus*-group, whereas *P. eurymedon* and *P. multicaudata* show total reproductive isolation). The domestic crabapple record for *P. eurymedon* is an error as previous records mentioned only apple. Species *alnifolia* is an error for *P. multicaudata*, as previous records mentioned only *Amelanchier*. The *Pieris "marginalis male"* photos on p. 117 are obviously female & same photo wrongly put on p. 119. Text states that *marginalis* may intergrade with *Pieris oleracea* along Bow River, so why are they split? (They state that Geiger & Shapiro also claim that *Anthocharis sara* and *stella* are differentiated at the species level, yet inconsistently do not split them). *Pieris rapae* occurs in some wild areas in Colo., up to 11,800'. *Pontia callidice occidentalis* does have a spring form, but the book fails to mention that *Pontia protodice* is just a temporary immigrant in Alta. late in the season, thus is summer form. The upf markings of *Pontia sisymbrii* photos are wide. Alta. bugs evidently are *Anthocharis julia columbia*. The second *Colias canadensis* photo is male ventral. Migration of *Colias eurymedon* is rare. *Colias gigantea* is surely a ssp. of *scudderii*; biological differences between them are dubious, *scudderii* is semipolyphagous on *Salix* & *Vaccinium* & *Polygonum*, so *gigantea* will probably prove to be also. *Colias nastes* larvae lack pink stripes in Alta. according to Clyde Gillette; the pink description comes from European larvae, which recent research suggests is a different species *Colias tyche* that includes *thula* & *boothii*. Spring *Colias philodice* also generally have submarginal unh dots; pupae do not overwinter. The *Colias pelidne* male photos are *Colias philodice* male spring form; the female *pelidne* photo is a female *C. occidentalis christina*. The "niphon" photo on the key on p. 152 is *eryphon*. The photos of *Lycaena "dorcias" florus* male look like *L. helloides* male. Photos of female *L. "dorcias" florus* are *L. d. dorcias* (the authors missed the opportunity to contribute anything regarding the relationship between *L. helloides*, *L. florus*,

and *L. dorcas*). *Papilio* (N.S.) #12 shows only *Potentilla palustris* is used by *L. d. dorcas*, and *L. florus* is a separate sp. Two hostplant errors published by Lembert (*Gayophytum* and *Oxytheca*) are repeated; *helooides* overwinters as egg not pupa. The N.A. maps for both *dorcas* and *helooides* were taken from Scott (1986) without adjusting for the book's species concepts. *Lycaena mariposa* eats *Vaccinium* not *Polygonum douglasii* (an error corrected long ago). The *Lycaena nivalis* female is male. *Ivesia* & *Horkelia* are errors for *Lycaena xanthoides editha* hosts. P. 163 misquotes Wright & Ballmer who actually found gene flow between *xanthoides* and *editha* even on the E side of the Sierras; and A. Warren's Butt. Ore. now adds *xanthoidesXeditha* intergrades in Ore. I still use the name *Lycaena thoe* instead of *L. hyllus*, as the *hyllus* figure was evidently *L. thersamon*. The book fails to list any distinguishing features of *Lycaena cupreus henryae*. *L. phlaeas* doesn't occur in Colo., and overwinters as larvae not pupae. Both photos of *Callophrys [viridis] sheridanii* uns seem to be ssp. *newcomeri* not *sheridanii*. The photos of *Satyrrium*=*Harkenclenus titus* are evidently ssp. *watsoni*, not *immaculosus*, though Alta. evidently has an intergrade between them. *Callophrys polios* doesn't always oviposit on dorsal leaf surfaces. The upper right "*Callophrys niphon*" photo is *C. eryphon*; upper left *clarki* is female. Do they intergrade in Alta?, the text mentions intergradation, 2 photos are misidentified as noted above, a key couplet is wrong, and Rob Robbins told me that *eryphon* and *niphon* intergrade. *C. niphon* doubtfully patrol to find females. *Callophrys gryneus* and *siva* intergrade S of Gothenburg Neb., and in the Sacramento Mts. N.M. and W Tex. *Satyrrium acadica* has no valid ssp., and N. Kondla found that the 3 dots in SW Alta. are actually Ted Pike specimens of *S. sylvinus nootka*, which belong on p. 184, where the photo looks like *nootka*. S Alta. evidently does have *Satyrrium liparops aliparops* if only some specimens have an orange upf patch (some have orange in Colo.). *Cercocarpus* is an error for *Satyrrium saepium*. *Strymon melinus* eats fruits, not leaves; photo is male (abd. is orangish on summer males). Poison ivy is an error for *Celastrina*, as the published record was *Rhus typhina*. *Trifolium* and *Thermopsis* are errors for *Everes amyntula*, which oviposits mostly on pedicels; *albrighti* is maybe a valid form in Mont. & BC with *grayer* uns, and maybe ssp. *maritima* occurs in Alta. (ssp. are weak and difficult in this sp.). The top photos of *Glaucoopsyche lygdamus couperi* are of an aberration, middle photos are ssp. *oro*, bottom female looks like ssp. *afra*. *Plebejus "idas"* is evidently two sp. in Alta. (see *Papilio* [N.S.] #12): *P. scudderi* which eats *Ericaceae* (*Ledum groenlandicum* is not a confirmed host as it was reported as only "probable" in Man.), & *P. atrapraetextus* which eats legumes. *Plebejus melissa* eggs--not larvae--overwinter; the male may be *P. atrapraetextus*. *Lotus corniculatus* is an error for *Plebejus acmon*, a lab food only. *P. alupini lutzii* occurs in Alta., not *P. acmon*, and *P. alupini spangelatus* occurs in the alpine zone at two known Alta. sites. *P. icarioides pembina* & *P. shasta minnehaha* occur in Alta. P. 212 3a is *Chlosyne palla* not *damoetas*. *Helianthus* and *Salix* and *Ulmus* are old bad errors for *Nymphalis milberti*. The book fails to mention that *Onobrychis viciaefolia* (not *viciifolia*) was published as a "fortuitous" host of *N. antiopa*. *Nymphalis californica* and *N. vau-album* have no ssp. The keys to *Polygonia zephyrus/oreas/faunus* are misleading. *Polygonia faunus rusticus* only occurs in Calif.; Alta. specimens should evidently be called ssp. *cenveray*, & *arcticus* is evidently similar to *cenveray* as the Yukon photo of "arcticus" female (not male) looks like that. *Polygonia gracilis* and *P. zephyrus* are both treated even though they say they cannot be distinguished in Alta. (actually they can, *zephyrus* has less-two-toned uns), and they repeat the same dot map and the same Scott N. Amer. blotch map of *gracilis* for both species! The Alta. *Polygonia interrogationis* was obviously a vagrant. *Polygonia progne progne* does not occur in Alaska & is rare stray in Colo. Alta. has *P. oreas near-silenus*. *Ulmus* is an error for *P. gracilis zephyrus*. *Lupinus* is dubious for *Vanessa carye annabella*. *Vanessa atalanta* overwinter as adults not pupae. Most of Scott's (*Papilio* [N.S.] #6) hostplant data for butterflies such as *Boloria* never made it into this book, even though that paper was cited in Bibliography. The *Boloria alberta "male"* is a female; male *alberta* do not hilltop, they patrol the hillside *Dryas* mats, and *B. alberta* does not occur on "barren" ridges except as singletons rapidly trying to find their proper *Dryas* mat habitat. *Boloria bellona jenistae* is a syn. of *bellona*. Alta. has *B. titania grandis* (not *chariclea*); *Salix herbacea* may be an error for *B. titania chariclea*. Alta. has *B. epithore chermocki=uslui*. *Thalictrum* is an oviposition error for *Boloria eunomia*; larvae lack blue dots. The photo is too pale to be *B. e. dawsoni*. *Boloria improba aberration youngi* is a syn. of *improba*; the Alta. ssp. is ssp. *nunatak*, which flies on lush tundra, not barren land. The Wyo. range of *Boloria alaskensis "napaea"* is missed on map; *Polygonum viviparum=Bistorta vivipara*. *B. selene* must have 2 gen. occasionally. In *Euptoieta claudia*, *Portulaca oleracea* is questionable (listed in a Brazil paper). Alta. has *S. atlantis hollandi* (upper photo)(N. Kondla notes it also from SW Alta. corner & Cypress Hills), *S. hesperis beani* (middle photo), & *S. hesperis dennisi=lais* (bottom photo). P. 252 photo looks like *S. coronis snyderi*. The *Speyeria egleis* photos look like ssp. *mcDunnoughi*, not ssp. *albrighti* the ssp. that would occur in Alta. P. 257 maybe they missed the small *Viola labradorica* on Horn Ridge. *Speyeria zerene* doesn't hilltop in Scott's experience. Paul Grey's map has so many "*garretti-picta*"s that *garretti* should be considered a syn. of the older *picta*. The genus *Charidryas* was used by Miller/Brown for no good reason. *C. acastus* has two gen. in Alta. according to N. Kondla. The photos of *Chlosyne damoetus* are *C. palla*. Alta. *C. damoetas* are actually *C. whitneyi altalus*, with brighter ups & concave fw margin. *Aster* is an error for *C. gorgone*, which has no ssp. Cypress Hills Alta. has *E. anicia near-anicia* and *E. bernadetta*. *Euphydryas editha* overwinters half-grown, not in final stage. *E. gillettii* 2nd photo is uns. The Alta. ssp. was later named *P. batesii saskatchewan*; young larvae live on, rarely under, a silk web. All *Phyciodes* overwinter as unfed 4th--not 3rd--stage larvae. *Phyciodes cocyta* was never known as *pratensis*. Pupae of *Phyciodes pulchella* are notably less-bumpy. Alta. has *P. pulchella owimba* & *P. tharos orantain*, both with orange antenna clubs. Alta. has *L. lorquini itelkae*. The *Limenitis weidemeyerii* illustrated has wide bands, not the narrow ones supposedly characteristic of *oberfoelli*. *Lethe*

anthoned overwinters as 3rd and 4th stage larvae. *Cercyonis silvestris* is misspelled, and is a ssp. of *C. sthenele* not *oetus*; ssp. *phocus* was characterized by grayer unh, *charon* by darker unh. *Ino* is a syn. of *C. pegala nephele*. *Coenonympha tullia* likes flowers; it's silly to claim that *C. ochracea* jumps from the S Rockies to N.W.T. with *C. inornata* in between, and there is no evidence that there is more than one species *tullia* in N.A. The N.A. map of *C. inornata* was unaltered from Scott without the necessary change to fit their species concept. W. Krivda wrote that *Erebia discoidalis pupae* overwinter. A large paragraph is devoted to apologizing why they can't decide their ssp. of *Erebia epipsodea*; the answer is simple, there aren't any valid ssp., but there is much individual variation, including a Colo. form *brucei* that lacks ocelli. *E. magdalena saxicola* is a dubious ssp. based only on a few paler body hairs; *E. magdalena* lays eggs on boulders, not under rocks, courtship occurs all day, and females occur near the males (males flait [patrol and sometimes perch] in rockslides nooks, while females occur on more vegetated areas nearby); the book fails to note that Yukon pops. are intermediate in red flush between *magdalena* and *mckinleyensis*. If the *Erebia stubbendorffii theano* record from Alta. is an error, then why is it on the N.A. map? (because those maps were taken from Scott's book unchanged); *Salix* was only an oviposition site, not a host. *Oeneis alberta* females are active in morning too; I haven't noticed any hilltopping. Later research in *Papilio* (N.S.) #12 shows that Alta. has *O. chryxus chryxus* (the photos) in mts. (*chryxus* oviposits on fallen or still-attached twigs above *Carex* mats under pine [usually] trees in Colo.), *O. calais caryi* in N Alta., & *O. calais altacordillera* in mts. It is doubtful if any butterfly eats *Juncaceae* in nature, even *Oeneis jutta*. *Oeneis melissa* overwinter as larvae, not pupae. *Oeneis polixenes* larvae cannot possibly mature the first year; the mts. have males near-*brucei* but females nearer *subhyalina*; but what ssp. occurs in extreme N Alta? The life history of *Oeneis taygete* is known at Churchill; I still lump it into *O. bore*. Female *Oeneis uhleri* are active in warm mornings also. *Neominois* actively mate-locate in early morning. Why so few dots for *Danaus plexippus*?, it must be a rare migrant to Alta. *Acerates* and *Apocynum* are dubious *plexippus* hosts. P. 313, *Viceroy*s are also semi-unpalatable.

BUTTERFLIES OF THE ROCKY MOUNTAIN STATES. C. Ferris & F. Brown, eds.

1981. 8 authors. University of Oklahoma Press, Norman, Oklahoma. 442 p.

I already published full corrections/review of this book in *J. Res. Lepid.* 20:58-64, though that review is somewhat dated. So this is brief, and corrects some errors in that review, and includes all the photo corrections. P. xiii, Cockerell lived in Las Vegas New Mex. P. 16, *Oeneis melissa* & *Papilio zelicaon* are rare in alpine meadows. *Neophasia menapia tau* is a local NW Wash. ssp. with more upf apical black. The first *Pontia sisymbrii* photo is female. P. 161 "astraea" f is christina f. P. 176 *pseudorotgeri* photo is *smintheus m* (*pseudorotgeri* is only in San Juan Mts.). *Apodemia mormo pueblo* occurs in SE Colo. *Celastrina lucia sidara* occurs on E slope of the Colo. mts., *C. l. lumarco* on W slope, *C. (argentata?) humulus* in Front Range foothills, *Cel. neglecta* on E plains, & *Cel. neglecta cinerea* in southern Sangre de Cristo Mts. R. Bailowitz later published in *J. Lepid. Soc.* that *herri* is a S Ariz. ssp. of *Cupido*=*Everes amyntula*. The area has multivoltine *Lycaena helloides*, & single-gen. *L. florus (castro)* in mts. incl. *L. florus megaloceras* in Bighorn Mts. & *L. f. near-megaloceras* in C Mont. (see *Papilio* [N.S.] #18). The area has *Satyrium titus watsoni* E of continental divide, and *S. t. immaculosus* W of it & in the NW Great Plains. The book should have figured *Callophrys [viridis] sheridani* *neoperplexa*, as nobody seems to know what it is or how it is distinguished from *newcomeri*. The first *Cyllopsis pyracmon henshawi* photo is female. P. 277 *damei* are *Cercy. pegala nephele*. P. 285 upper left male is *Oeneis uhleri*. P. 310 upper left male & female are *S. atlantis sorocko*. P. 319 photos are *P. batesii* near *batesii* from Syracuse, Onondaga Co. N.Y. P. 324 right female is *C. palla calydon* female. P. 331 *alena* is the later-named *E. anicia wecoeut*. P. 332 the *maria* & *windi* paintings in Howe (1975) are correct, as Todd Stout finds that *maria* is mostly black, and the orange *windi* would seem to be a syn. of *E. anicia eurytion* (or *eurytionXancia* thus invalid) which this book claims ranges N at moderate elevations into Wyo. & Montana. P. 340 brown female is the later-named *P. oreas nigrozephyrus* female. Some migrants of both *Libythea carinenta larvata* & *L. carinenta bachmanii* are known in Colo.

FLORISSANT BUTTERFLIES. T. C. Emmel, M. C. Minno, B. Drummond.

1992. Stanford University Press, Stanford, California. 290 p.

Florissant is a relatively flat wooded area of the Colo. mts. that has a rather dull fauna for Colorado, except it has a lake-bed shale deposit with lots of fossil tree stumps & fossil plants & bugs. The butterfly fossils are illustrated (some are spectacular, some are vague wrecks). The best feature of the book is the many b/w photos of larvae & a few pupae, most by M. Minno. The book has 8 color plates of adults. The text is rather skimpy for each species, and there are a lot of errors.

Color plates: Pl.I#4a-c are *E. persius*, probably coll. by me. Pl.I#7b is *Pyrgus ruralis*. Pl.II#23a-b are *P. zelicaon*; 23c-d are *P. zel. form nitra*. Pl.IV#42a&c are *Lycaena florus near-sangreamar*. Pl.V *Oeneis chryxus* live adult is *O. calais altacordillera*. Pl.VI#82a-c are *Chlosyne palla flavula*. Pl.VI#83b-c are *Chlosyne leanira alma*. Pl.VI#77a-c are *Phyciodes cocyta selenis*. Pl.VII#75c is *Spey. hesperis hesperis*. Pl.VII#75d is *Spey. coronis halcyone*. Pl.IX#80a[same specimen as fig. 47]-b are *Phyc. cocyta selenis* from Big Spring Ranch.

Detailed comments: P. 2, the first Lepidoptera probably ate detritus, as Micropterygidae do today. P. 13 Colo. has more butterfly species than Calif. P. 32 photo is *Rumex crispus*. P. 33 *Megathyms yuccae* occurs at a similar elevation at Tarryall River in Park Co. *Erynnis afranius* surely does not occur in the area, and the plate figs. are *E. persius*. *Pyrgus xanthus* is a mtn. species, but not alpine. Mature larvae of *Pyrgus scriptura* probably hibernate. *Stinga* eggs hatch quickly, & mature larvae hibernate. *Hesperia comma ochracea* most often ovip. on turflike sedges. *Polites sonora* eggs are laid without glue and fall into the litter. *Paratrytone snowi* host is only *Muhlenbergia montana* (*Blepharoneuron* was Scott's error). *Parnassius phoebus* adults often feed; eggs hibernate. *Nitra* is just the black form of *P. zelicaon*. *Papilio glaucus rutulus* ovip. on ups of leaves. *Neophasia* lay a row of eggs on the pine leaf. *Eriphyle* is limited to S BC. P. 56, maybe the record of *Rumex triangulivalvis* is an error, as fig. 23 is not that but is *R. crispus*. The *Lycaena* "helooides" in the area have just one gen. in July so are surely *Lycaena florus*, the mtn. sp. with one gen. (*L. helooides* has 2-3 gen. and lives at lower alt. on the plains & ovip. on detritus below the hosts). *Satyrium californica* ovip. on stems. *Callophrys affinis homoperplexa* surely eats *Ceanothus fendleri* often in the area. *Strymon melinus* abdomen is orange only on summer males. *Trifolium* is an error for *Cupido* "Everes" *amyntula*. *Celastrina lucia sidara* (not *cinerea*) occurs in the area (main host *Jamesia*). *Plebejus melissa* ovip. on lower stems of host, & eggs hibernate. *Plebejus saepiolus* has just one gen. L June-M Aug., & in the Front Range of Colo. *Astragalus agrestis* & *A. alpinus* are preferred over *Trifolium* (*Papilio* [N.S.] #6). *Plebejus* (*Agriades*) *glandon* eats *Androsace* in Europe. *Apodemia mormo pueblo* was later named from the area. *Polygonia faunus rusticus* is a Calif. bug. *Polygonia oreas nigrozephyrus* surely occurs in the area as it occurs all around it. *Vanessa virginiensis* adults hibernate. The return migration of *Vanessa cardui* is usually weak. *Speyeria* "atlantis" has been split into two sp. (*Papilio* [New Series] #8), and text doesn't say whether the local bugs are unsilvered; but I have seen specimens of *Spey. hesperis hesperis* & *Spey. atlantis sorocko* (not "electa") in Teller Co.; *hesperis* occurs in aspeny wooded areas, *sorocko* in wet meadow/heavily forested areas averaging higher in altitude (75d on pl. VII is *Spey. coronis*). *P. cocyta selenis* is the proper name for "tharos pascoensis", and *P. pulchella camillus* for *P. campestris*. *Phyc. pallida* does not occur in the area, though it evidently occurs northward along the South Platte River into Park Co.; the fig. 47 and Pl. IX#80a-b photos are all *P. cocyta selenis*. The *Euphydryas* treatment is confused; they describe the early bug as like *E. anicia* "capella" (probably just *eurytion*), and the later bug as yellower (evidently *eurytion*)(doubtfully *Euph. bernadetta* which has recently been separated as a distinct sp., *Papilio* [N.S.] #12). They say the hostplant isn't known, but say that larva & pupa were found on Big Spring Ranch, and below the Pl. I legend are 2 photos of a near-"capella" female from near Florissant and a white "capella" larva on Castilleja, which is probably the larva from Big Spring Ranch which is 4 mi. SW Florissant, as Emmel's (1963) description is like this photo; based on the habitat & altitude & known range of *eurytion*, *eurytion* occurs here, not *capella*; *E. anicia eurytion*=*carolae* occurs in NW South Park, see *Papilio* [N.S.] #12). *Cyllopsis pertepida* larva has two long head horns. *Coen. tullia ochracea* often feeds on flowers. *Erebia epipsodea* males surely can mate within a day or so after mating, and their flight is rather even compared to other *Satyrinae*. *Neominois ridingsii* sometimes visit flowers, and males are active in early morning ~7:50-11:00. *Oeneis* "chryxus" in Colo. is really two species, *O. chryxus* which flies in even-numbered years and oviposits on tree twigs above turflike sedges under conifer trees, and *O. calais altacordillera* which occurs in even- and sometimes in odd-numbered years and oviposits on grasses (see *Papilio* [N.S.] #12). *Florissant* has true *O. chryxus* (which does not ovip. on grasses as book claims), while *O. calais* occurs at the S end of Rampart Range NE of Woodland Park, and surely occurs on Pikes Peak (Pl. V live adult is *altacordillera* from Mosquito Range). The larva photo p. 87 doesn't show the heart-band enough (the heart-band is generally solid brown in *O. chryxus*, solid to just a row of dashes in *O. calais*).

THE BUTTERFLIES OF MANITOBA. P. Klassen, A. Westwood, W. Preston, W. McKillop. 1989. Manitoba Museum of Man & Nature, Winnipeg, Manitoba. 290 p.

This book is surprisingly good and professional, considering that only Klassen was a well-known lepidopterist. There are almost no misidentifications! The treatment of difficult bugs such as *Polygonia* & *Phyciodes* & *Argynnis* (*Speyeria*) "atlantis" is very good. The color plates are nice, the text has nice dot maps, detailed records are listed, many references are given, and a conversion table of common to scientific hostplant names is given.

On the plates I note mostly just taxonomic progress: Pl. 3 fig. 32, Chaffee; Pl. 5f4 is *P. polyxenes kahli* (connected eyespot); 8f1-3 are *P. glaucus canadensis*; 10f11-15 Col. *occidentalis christina*; 12f16 is odd for *dorcas* as it looks like *florus*; 14f11-12&17 *Cel. lucia*; 14f16&18 *Cel. arg. argentata*; 14f19-22 *Pleb. scud. scudderi*; 14f35-37 *Plebejus glandon suttoni*; 17f1-2&4 *Spey. atlantis* near *hollandi*; 17f3&5-6 *S. hesperis dennisi*; 19f7 ssp. *harrisi*; 19f8-10 ssp. *hanhami*; 19f15-18 *P. cocyta selenis*; 19f19-22 *P. batesii saskatchewan*; 19f25-26 *E. anicia helvia*; 23f11-14 *C. tullia benjamini*; 23f15-17 *C. t. inornata*; 25f4-6 *O. calais calais*; 25f11-12&14 *O. alberta ojbwe*; 26f10-12 *O. polixenes luteus*.

Epargyreus clarus pupae hibernate. *Hesperia comma assiniboia* does not occur in N.M. (ssp. *ochracea* in Colo. is similar). Klassen wrote to me that *H. comma* from Thompson & Gillam are somewhat intermediate *manitoba*/*laurentina*. I doubt that *Polites peckius*=*coras* hibernates as pupa. Klassen told me that *Anatrytone l. logan* occurs in Man., but *Culross* has intermediates to ssp. *lagus*. *Poa* is a lab host for *Amblyscirtes vialis*. *Kahli* is a ssp. of *Pap. polyxenes* mostly limited to Riding & Duck Mts. *Euchloe ausonides* eats flower buds. Man. has *Colias occidentalis christina*, not *C. alex. chr.* *Eurema lisa* record at Sandilands, p. 255. *Lycaena helooides* hibernates as egg. Man. evidently has *Satyrium titus watsoni*=*winteri*

in SE (but pl. fig. 27 has tiny dots), intergrades *watsoniXimmaculosus* in SW. P. 94 photo is *Satyrium calanus*. *Callophrys polios ovips.* at bases of leaf buds. Man. evidently has *Celastrina lucia* all over Man. (pl. 14 figs. 11,12,17) and *Cel. argentata* in S Man. (pl. 14 figs. 16, 18, evidently same sp. as *serotina* & *humulus*)(maybe the two gen. stated in S Man. is a combination of the two sp., or ?maybe *Cel. neglecta* creeps into S Man.). *G. lygdamus afra* is in Riding Mts. & has smaller unh spots. *Plebejus melissa* eggs hibernate. Churchill evidently has *P. glandon suttoni*. Man. has *Speyeria hesperis dennisi*, not *electa*. *Boloria eunomia* from Churchill is a new ssp. with unh silver spots small & submarginal pale rectangles redder. *Boloria frigga* hibernates as 4th-stage larva. P. 139 *boisduvali*. *Lysimachia* is an error for *Chlosyne gorgone*. *Phyciodes* hibernate as unfed 4th-stage larvae. Man. has *P. cocyta selenis=morpheus*; the two-gen. bugs in S Man. may be *P. diminutor*. *P. batesii saskatchewan* occurs in SW Man. at least, maybe *P. b. lakota* in SE Man. (I have not seen those). *E. anicia* is an error for Man. (J. May also mislabeled *Mellicta athalia* from BC). *Polygonia faunus* lay many eggs/day. P. 218 *Vernonia*. *V. cardui kershawi* occurs in Australia. *Coen. tullia* from The Pas show more unh postbasal bars, maybe an influence from another ssp. *Erebia rossii* dot from The Pas is dubious according to P. Klassen letter. *Erebia mancinus* is in Man. *Erebia epipsodea* has no ssp. Neominois males actively mate-locate in early morning. Man. has *Oeneis calais calais*; larvae hibernate twice, as young and old larvae in two winters. *O. uhleri* hibernate as 2, 3, 4, 5th stage (young & old) larvae. The *O. alberta* E of Red River were later named *O. alberta ojibwe*. P. 249 *misandra*. P. 253 *Lintner*. P. 261 *Leslie* 1980 missing (cited p. 72).

BUTTERFLIES OF SASKATCHEWAN. Ronald R. Hooper. 1973.

Museum Natural History, Regina, Saskatchewan. 216 p.

A good little book that is quite likeable, despite being out of date. Nice b/w photos, good mostly-accurate though skimpy text (not much was known about Sask. butterflies), not too many errors. *Hesperia dacotae* was later found in SE Sask. *Longleyi* is a syn. of *Ancyloxypha numitor*. P. 61 *kahli* is a ssp. of *P. polyxenes*. P. 63 *dodi* is a syn. of *P. machaon "bairdii" brucei*; the photo is maybe an odd *P. zelicaon*. The Badlands Old World Swt. photos are *P. m. brucei=dodi*. P. 67 the *bairdii* from N.D. must be *P. polyxenes*. P. 68 bottom looks like *Colias philodice* male. *Pontia protodice* & *P. callidice occidentalis* are wrongly lumped; the photos are *occidentalis*; the text gives the range of both. The book wrongly lumps *Colias eurytheme* & *C. philodice*, and the text does not distinguish them. *Christina* is a ssp. of *C. occidentalis*. The book lumps *Euchloe ausonides* & *creusa* (Butt. Canada lists 5 *creusa* dots across C Sask.). *Apodemia mormo* has been found SE of Killdeer. *Callophrys nippon* is not in Cypress Hills. *Sheltonensis* is a *C. eryphon* ssp. in NW Wash., not Sask. Sask. *Satyrium titus* uns dots are rather tiny in N.D.-Sask.-Alta. so evidently they are *watsoniXimmaculosus*. The stray Sask. *Lycaena phlaeas* was surely *hypophlaeas* from E U.S. P. 111 *scudderi*. Sask. has *Plebejus alupini lutzi*. Albright has gray uns, so Sask. evidently has *Cupido amyntula maritima*. Sask. has *Celastrina lucia* all over, *Cel. neglecta* in the S half (see Butt. Canada), and probably also *C. argentata* in S Sask.? *Zephyrus* is a ssp. of *Polygonia gracilis*, and the *gracilis* figures look like *gracilisXzephyrus*. The Woodland Pearl Crescent is *Phyciodes cocyta selenis*, & the Prairie Pearl Crescent is *P. tharos orantain*; Hooper was one of the first people to publish this distinction. *P. "campestris camillus"* does not occur in Sask., the photos are Alta. *P. pulchella owimba*, and the Sask. records are *P. batesii saskatchewan* which is the only *batesii* ssp. in Sask. Photos & text are *Chlosyne sterope acastus* (which does not eat paintbrush), not *C. palla*. *Euphydryas bernadetta* & *E. anicia* near-*anicia* fly together in Cypress Hills, *bernadetta* averaging a little earlier. Sask. has *Boloria freija freija*, not *natazhati*. Sask. has *Speyeria zerene picta*. *S. mormonia* has one gen. *S. atlantis hollandii* and *S. hesperis dennisi=helena* do not interbreed (the cross-bred specimens on p. 171 were misidentified according to N. Kondla); *dennisi* also is at Duck Mtn.; the Cypress Hills evidently has *S. hesperis* near-*beani*. *Mayae* is a syn. of *S. aphrodite manitoba*. *Ino* is a syn. of *Cercyonis pegala nephele*. Sask. has *Oeneis calais caryi*. *O. uhleri* has one gen.

BUTTERFLIES OF NORTH DAKOTA. AN ATLAS AND GUIDE. Ronald Alan Royer.

2003. Minot State University Science Monograph #2. Minot, North Dakota. 192 p.

This is a nice little book. The color photos of each butterfly are at the top of each text page, and the map is at the bottom, so everything can be seen at once, rather than flipping through a book for photos and maps as in most books. And only 3 photos are misidentified. It is not out of print, despite internet \$ gouging caused by its pleasing appearance (send \$40+\$5-in-U.S. shipping to Minot State Univ. Bookstore, 500 Univ. Ave. W, Minot, ND 58707). Notes: *Epargyreus clarus* males rait (perch to await females) only in morning, and rest in the shade in the afternoon. *Erynnis icelus* in SD is only in the Black Hills, so *icelus* is no doubt absent in SC & SE N.D. P. 19, the hw fringe is not white-tipped on the *afranius* photos. For *Erynnis persius*, "aspens and willows" are errors based on *icelus*. Scott knows of no evidence for *Pyrgus communis* immigration. *Pholisora catullus* usually eats *Amaranthus*. *Ancyloxypha numitor* eats various hay grasses. *Hesperia comma* hibernates only as eggs. *Hesperia ottoe* "flourishes only in ungrazed native prairies" and *Polites origenes* occurs in "undisturbed native prairies" and *Atrytone arogos* is in "undisturbed native prairie" (Butt. S.D. also notes this for *arogos*) because larvae of all three are aerial on the plants—unlike all other *Hesperia* and *Polites* which make nests in plant base/soil litter—so cows can eat them (see J. Scott, *Papilio* [New Series] #6, who found *Andropogon gerardii* the main and only host for the three). *Poa* is dubious for *Hesperia leonardus pawnee*, and larvae hibernate in 1st stage. *Polites*

peckius in ND is ssp. peckius, as photos and text indicate a larger unh patch than farther south. For *Polites mystic*, *Agropyron* & *Phleum* are H. Tietz errors, based on *Polites themistocles*. The female *Wallengrenia egeremet* photos are really *Polites origenes* (egeremet females seem to lack any upf orange, whereas the photo has orange on the costa; *themistocles* females always have orange, and *origenes* females rarely have a little; egeremet females have the unh spots more elongate basodistally, and the spot in cell M_2 is positioned more basally; *themistocles* usually have a more ochrey unh with no spots though there are often a few anteriorly or occasionally small traces; *origenes origenes* females do often have the tiny row of pale spots on darker unh as on the photo). *Poa* is rather dubious for *Poanes hobomok* as the bug is expected to be a hay-grass feeder like *Poanes taxiles*. *Cyperus esculentus* is a lab host only for *Euphyes vestris*, the native hosts are turflike dryland sedges in Colo. *Bouteloua curtispindula* is the main host for *Amblyscirtes oslari* in Colo. *Poa* is an error for *Amb. vialis* as *P.* larvae died eating it. *Leussleri* is a syn. of *Megathymus streckeri texana* or a weak ssp. *Kahli* is evidently a ssp. of *P. polyxenes* as it has a definite geographic range, though many individuals resemble *asterias* so maybe the ssp. is weak. ND has *Papilio machaon brucei*=*dodi*, as the predominantly-black ssp. *bairdii* stops in C Colo. *P. zelicaon f. nitra* is produced by a dominant gene; its frequency should have been stated (it is ~5% in Front Range Colo., but up to 30-50% in W Neb.). Obviously *glaucus* and *rutulus* are ssp. of *Papilio glaucus*, and ssp. *glaucus* occupies SE N.D., and the “*canadensis*” in W N.D. are *glaucus*X*rutulus* intergrades, just as they are in the Black Hills of S.D., while ssp. *rutulus* does not occur in N.D. The male *Pontia protodice* photo is actually *P. occidentalis*. *P. protodice* also hilltops, and the spring form also has darker tan-green unh. Early spring *Colias eurytheme* have much less ups orange (sometimes none) like the hybrids with *C. philodice*. *C. alexandra altamont* occurs in N.D. *Longi* is a syn. of *Lycaena rubidus rubidus* in Scott’s opinion. *Satyrium acadica* males rait only in late afternoon-sunset. *S. titus* have tiny uns dots in N.D. so are near ssp. *immaculosus*; oaks is an error. For *Satyrium edwardsii*, *Q. macrocarpa* is Bur Oak. *Satyrium liparops aliparops* has weak uns white lines (a photo from Minneapolis also has weak lines) and little to considerable ups orange flush, while ssp. *fletcheri* seems to have weak uns lines and much orange flush, while ssp. *strigosa* has stronger unh lines and no flush, so Royer seems to be correct in the ssp. The photo of *Callophrys gryneus siva* female from Slope Co. has an offset unh line, so has intergraded with ssp. *gryneus* some. *Obscurus* is a syn. of *Callophrys polios* in Scott’s opinion, and *franki*=*humuli* are syns. of *Strymon melinus*. N.D. *Cupido amyntula* are evidently a ssp. (different from ssp. *amyntula*) with whiter uns ranging to Man. & even Mich., which evidently should be called ssp. *maritima*, as the N. Kondla’s photo of lectotype of *maritima* is whitish on uns and spotted like the photos. Ssp. *albrighti* is evidently a ssp. in Little Belt Mts. of C Mont. (and a ssp. or form in BC, where some have whiter rings around the black spots); Steve Kohler notes that uns spot size is quite variable in both *albrighti* & *amyntula*, but *albrighti* has uns more gray (whitish-gray in ssp. *amyntula*), is smaller, with male ups more purplish-blue (vs. lighter blue in *amyntula*), and female ups darker gray-black with little blue. There is lots of variation and current confusion among *C. amyntula* ssp., but N.D. does not have *albrighti*, and *maritima* seems the best name now. The photos of *Glaucopsyche lygdamus* are definitely not oro, they fit *afra* (TL Brandon Man. due S of Riding Mts., the neotype female designated by F.M. Brown has tiny unh dots) which seems to be a valid ssp. in Scott’s series from Riding Mts. S Man. & Nipigon Ont., which has tiny unh dots & larger unf dots. *Glycyrrhiza* is a recorded host of *Plebejus melissa* but is not a host in Colo. N.D. evidently has *Plebejus saepiolus amica*? N.D. has *Plebejus icarioides* near *lycea*, which Steve Kohler finds in SE Mont. *Plebejus alupini lutzii* is evidently the correct name for N.D. “*acmon*”, as the upf border is narrow on the photos here and in Sask. (though there are two gen. as in ssp. *texanus* farther S); *Astragalus* & *Lupinus* are errors, & apply to true Calif. *P. acmon*. Chris Schmidt finds that Canada *Plebejus* “*glandon*” “*rusticus*” is a new ssp. related to *megalos*. N.D. evidently has *Apodemus mormo mormo*. *Desmodium* is a dubious C. Abbot host for *Euptoieta claudia*. N. D. evidently has *Argynnis* (*Speyeria*) *aphrodite manitoba* based on p. 121 photos (not ssp. *aphrodite*), but the p. 125 female is like *whitehousei*=*ethne*, so if the darker ones have been misid. as *S. atlantis*, then N.D. has ssp. *manitoba*X*whitehousei*. *Speyeria aphrodite* & *edwardsii* females also aestivate. The female *Speyeria atlantis* photo is really *S. aphrodite whitehousei*. N N.D. evidently has *S. atlantis hollandii*, while the two SW N.D. counties evidently have ?*S. hesperis lurana*X*dennisi*? *Speyeria mormonia eurynome* evidently is the N.D. ssp. The ssp. of *Boloria selene* are very weak (only *atrocostalis* is distinctive), so ssp. *sabulocollis* would supposedly occur in SW N.D., *myrina* in the remainder. *Aster* is not known to be a *Chlosyne gorgone* host to Scott, while *Rudbeckia hirta* has one Colo. host record. Evidently *Chlosyne nycteis reversa* occurs in Bottineau Co., ssp. *nycteis* elsewhere. N.D. has *Phyciodes tharos orantain*, though SE N.D. might have ssp. *tharos* with black & white antenna club. *P. cocyta selenis* occurs in N.D., though as noted SW records may be *tharos*. *P. batesii lakota* occurs in N.D., and the unh brown marginal patch is still present and conspicuous. The first *Polygonia comma* and *P. progne* photo is of course spring form, the others summer form. The band of *Nymphalis californica* records across E N. Amer. does not fit migration, it is more like the *Polygonia satyrus* distribution in the east. *Nymphalis antiopa* is bivoltine in Colo. *Vanessa hibernation* isn’t well known, but *V. atalanta* hibernates as adults regularly, and *V. virginensis* and *V. carye annabella* hibernate much farther north than *V. cardui*. *Limenitis archippus* is somewhat unpalatable too. *Oberfoelli* is a syn. of *Limenitis weid. weidemeyerii*. N.D. seems to have *Asterocampa celtis celtis*, and *Asterocampa* are natives, not strong migrants. *Lethe anthedon* hibernates as 3rd-4th stage larva. N.D. has *Cercyonis pegala nephele*. *Neominois ridingsii* rait early in the morning, when they are very active. *Varuna* isn’t a good ssp. of *Oeneis uhleri*, being merely a bit smaller.

FIELD GUIDE TO BUTTERFLIES OF SOUTH DAKOTA. Gary M. Marrone. 2002.

South Dakota Department of Game, Fish, & Parks. Pierre, South Dakota. 478 p.

This is a useful book. It has nice color photos of each species, though many require flipping a page to be viewed. Few photos are misidentified. The map is with the text, which is nice. Sometimes the text seems skimpy, but it is much larger than the Michigan & Missouri books. My only major complaint is that many hostplants are given obscure common names that cannot be found in any major flora, as noted below; there should have been an appendix to convert these names to scientific names, but G. Marrone sent me a list deciphering the questionable hosts, as noted below with the asterisk*. Nearly all of these hostplants are from the literature, and few are from S.D. Detailed comments: For *Parnassius phoebus* *smintheus* (*sayi* is a syn. of *smintheus*) p. 26 spearleaf stonecrop is *Sedum lanceolatum**. S. D. has *Papilio machaon* *brucei*=*dodi*, not ssp. *bairdii* which is predominantly black and stops in C Colo.; males flit and rait on hilltops to seek females. The male photo of *brucei* may be *P. zelicaon*, as the abdomen uns is black and the tegulae aren't very yellow. The female f. "hollandii" is really *Papilio polyxenes* f. *pseudoamericanus*, as the unh base is yellow making it a yellow form (*hollandii* is a black form with yellower abdomen), and the many orange spots on unh postmedian band are more than most *machaon* and are typical of *pseudoamericanus*, and tegulae are blacker evidently (broken off?), and Marshall Co. is out of range for *P. machaon* which only occurs in W S.D. *Nitra* is the black form of *P. zelicaon*, not a ssp., and its frequency should have been more precisely stated; *gothica* is a syn. of *zelicaon*. For *P. zelicaon* p. 42, wild parsley is *Lomatium foeniculaceum**, and heartleaf alexanders is *Zizia aptera**. Obviously *P. glaucus* *glaucus* occurs in most of S.D., and SW S.D. has *glaucus* *glaucus*X*glaucus* *rutulus* *intergrades* (not *canadensis*), and *rutulus* is just a rare form in SW S.D. The male *glaucus* photo has a solid unf submarginal band, so is the spring form evidently. The male on p. 60 has little unh orange, as it is an intergrade. Note that L. Brower proved intergradation from *glaucus* to *rutulus* in Black Hills due to input of genes from the latter (Evolution 13:40-63). For *Papilio eurymedon* p. 66, snowbrush Ceanothus is *Ceanothus velutinus** (*C. herbaceus*, *C. fendleri*, & *C. velutinus* all occur in Black Hills). *Magnolia* is an error in past literature for *Papilio troilus*. *Nordini* is a syn. of *Pontia* s. *sisymbrii*. The female photo of *Pontia* *protodice* is actually *P. callidice* *occidentalis*. For *P. protodice* p. 78, common pepperweed is *Lepidium densiflorum**. *Pieris* "marginalis" and *Anthocharis* "stella" are known only from single records coll. by Edwin M. Perkins in Allyn Mus. Entomology (G. Marrone e-mail Oct. 10, 2007); according to S. Spomer, Kilian Roever coll. an *Anthocharis* in the Black Hills in nearby Wyo., so the S.D. Black Hills sara is probably valid, but could the S.D. "marginalis" be mislabeled? The photo of Manitoba *marginalis* is *Pieris oleracea* according to dogma and the maps in Guppy/Shepard BC book, and anyway Scott is not certain that *pseudonapi*=*mcdunnoughi* belongs to *P. marginalis*, though S Bowden's lab crosses determined that it could be. The male female "oleracea" photos from Colo. are *P. marginalis* *pseudonapi*=*mcdunnoughi* according to the usual dogma. *Palaeoreios* is a syn. of *Euchloe* *aus. coloradensis*. For *Euchloe* *olympia* p. 93, blue mustard is *Chorispora tenella*, a new host (*Marrone observed ovip. in Black Hills). *Anthocharis julia* *julia* is the bug in S.D. The male photo is *A. j. julia*; the female photo looks like the new ssp. *Anth. sara* *colorado* from SW Colo. *Colias philodice* & *C. eurytheme* *hibernate* as 3rd (sometimes 2nd-4th)-stage larva. *Colias occidentalis* does not occur in Baja Cal., that is *C. harfordii*. For *Col. occidentalis* *krauthii* p. 103, alpine sweetvetch is *Hedysarum alpinum**. For *krauthi* and "alexandra", northern sweetvetch is *Hedysarum boreale**. The *Colias* "alexandra" is now known as *C. edwardsi* *altiplano*. For *Zerene cesonia* p. 107, false indigo means *Amorpha fruticosa**, but that common name shouldn't be used because in floras it also means *Baptisia*. *Americana* is now considered a syn. of *Lycaena phlaeas* *hypophlaeas*, which did not immigrate from Eurasia (Scott J. Lepid. Soc. 47:253-4, & S. Kohler Taxon. Rept. 7[1]) because Eurasian bugs differ and have more uniform unh, more diffuse unh red line, and European *phlaeas* have longer tails and browner unh & little notches on basal edge of uph orange band; *hypophlaeas* evidently arrived from Siberia or evolved from *L. p. feildeni* long ago. For *Lycaena* *thoe*=*hyllus* p. 131 & *helloides* p. 135, marsh smartweed is *Polygonum amphibium*=*coccineum**. For *helloides* p. 135, prostrate knotweed is *Polygonum aviculare*. *Longi* is a syn. of *Lycaena* *r. rubidus*. *Polygonum douglasii* is an occasional host of *Lycaena* *florus*, which is now considered a separate sp. from *L. helloides*. *Satyrium titus* ssp. are mostly weak (Gatrelle's sample sizes in Taxonomic Report 4[6] were deplorably minuscule), but evidently the ssp. in SD is *watsoni* (TL Kerrville Tex., one paratype has small red spots but O.D. only claimed uns was paler)=(winteri Gatrelle TL Sherborn Mass.)(=campus Gatrelle TL Shelby Co. Iowa) which occupies most of E U.S. and has larger black dots and redder spots (=occidentalis Austin & Emmel TL Pershing Co. Nev.), while ssp. *titus*=*mopsus* (both TL Screven Co. Ga.) from the coastal plain of Fla.-Ga. has whiter edging to unh marks, *immaculosus* (TL Provo Utah) has tiny or absent uns dots and occurs W of the continental divide and in N.D., while the E-slope Colo. bug has small black dots and oranger spots evidently *watsoni*X*immaculosus*. *Satyrium liparops* *aliparops* in S.D. surely has some adults with orangish ups flush? *C. gryneus* *gryneus* has the postmedian unh line kinkier than *C. g. siva*. *Obscura* is a syn. of *Callophrys* *p. polios*. *Franki* is a syn. of *Strymon* *m. melinus*. *Everes amyntula* *valeriae* also occurs in E slope Colo. For *Cupido* (*Everes*) *amyntula* p. 166, *Glaucopsyche lygdamus* p. 177, & *Thorybes pylades* p. 350, cream-colored vetchling is *Astragalus ochroleucus** (also in Britton/Brown flora). *Celastrina lucia* *sidara* also has f. *lucimargina* with dark margin & patch; how frequent is the patch? H. Pavulaan states that Black Hills have intergrades *sidara*X*nigrescens* etc. For *Celastrina neglecta* p. 170 & *Erynnis martialis* p. 362, inland ceanothus is *Ceanothus herbaceus** (p. 362 says it's the same as New Jersey Tea, but both *Ceanothus americanus* & *C. herbaceus* are N.J. Tea in Great Plains Flora)(Marrone states that *C. herbaceus* [ssp. *pubescens*] is common in Black Hills & *C. fendleri* is

uncommon in W Black Hills). *Glaucopsyche lygdamus afra* evidently occurs in NE S.D. as the bugs there have smaller unh dots like *afra*. For *Plebejus melissa* p. 179 white crazyweed is *Oxytropis sericea**. S.D. evidently has *Plebejus saepiolus saepiolus*, not the small-dotted *amica*. S.D. has *Plebejus icarioides lycea* (not *pembina*), which also occurs in Bighorn Mts. & SE Mont. For *Plebejus shasta* P. 185 tufted milkvetch is *Astragalus spatulatus**. S.D. evidently has *Plebejus alupini near-texanus*, as there are two gen. and the upf border is wide on the photo. *Plebejus glandon rustica* hibernates as 2nd-stage larva. *Apodemia mormo* female photo from Colo. has some uph orange, so could be a very dark ssp. *pueblo* from SE Colo, but is probably a variant ssp. *mormo* from W Colo. Most of S.D. evidently has *Argynnis (Speyeria) aphrodite near-whitehousei=ethne* instead of *manitoba*, though the male looks a little paler (& a female from Spearfish Can. Lawrence Co. I examined was *whitehousei*). For *Speyeria edwardsii* p. 214, *S. zerene* p. 221, *S. hesperis* p. 229, *S. mormonia* p. 231, longspur violet is *Viola adunca** (*adunca* is called “hook-spurred violet” in Great Plains Flora). The *Speyeria “edwardsii”* nectaring on p. 215 is really *S. aphrodite*. The female *coronis* photo is *Speyeria coronis halcyone*, but some *snyderi*-like adults occur in Wyo. and the 5m4f I have examined from Black Hills had red-brown discs except 1m had green disc & 1f had green-over-brown disc like *snyderi*, so the S.D. ssp. is near-*halcyone*. S.D. evidently has the pale-brown-disc *Speyeria zerene platina* or near-*platina*, not ssp. *sinope*; *platina* also occurs in Wind River and Bighorn Mts. Wyo. (though some dark-brown disc variants like ssp. *picta* occur in Bighorns), while the greenish-brown-disc *sinope* occurs only in S Wyo.-Colo. Maybe S.D. has ssp. *harmonia*, but it is basically a syn. of *Speyeria callippe calgariana* anyway. E Alaska, C Calif, & Ariz. have only *Speyeria hesperis*, not *atlantis*. For *S. atlantis* p. 227 meadow violet is *Viola papilionacea** in Britton/Brown. Bighorn Mts. *S. hesperis* may perhaps best be called *luranaXdennisi*, as the unh submarginal band is wider than *lurana* and unh disc has more pale tan areas and the black bars are narrower. The Black Hills have a new ssp. *S. mormonia kimimela* described by Marrone, Spomer, & Scott. The ssp. of *Boloria selene* are very weak, although elsewhere *nebraskensis* is larger, & *atrocotalis* has darker ups borders. *Carlota* is a syn. of *Chlosyne gorgone*. For *C.nycteis* p. 240 cutleaf coneflower is *Rudbeckia laciniata**. The female *Chlosyne nycteis* photo is a male. For *Chlosyne acastus* p. 242 showy aster is *Aster conspicuus**, but is an error for *acastus* according to Marrone (it is a known host of *Chl. hoffmanni*), and rubber rabbitbrush on p. 215 & 242 is *Chrysothamnus nauseosus** (p. 215 shows the white tomentose carpet on the twigs)(Marrone wrote me that this is merely a likely host due to association); readers should note the odd two generations in S.D.-N.D.-Alta. For *Phyciodes tharos* p. 246 & *P. pulchella camillus* p. 252 heath aster is *Rose Heath Aster=Aster ericoides**. W S.D. surely has *Phyciodes tharos orantain* with orange *nudum*, and ssp. *tharos* evidently occurs in the far E with black *nudum*. *Phyciodes cocyta selenis* larvae don't have pinkish spines as Charles Oliver wrote. *Phyciodes batesii lakota* has a definite brown unh marginal patch (the unmarked unh describes ssp. *batesii*); *P. batesii* also occurs W to NWT and Ariz.; the female photo is a male. All *Phyciodes* hibernate as unfed 4th-stage larvae. For *Phyciodes pulchella* (not *pratensis*) *camillus* p. 252 marsh aster is *Aster hesperius**. *Phyciodes pallida pallida* (not *barnesi*) evidently occurs in S.D., as the photos resemble that (the larva description fits *pallida*, as *barnesi* larva is blacker, though the desc. is no doubt from Colo.). For *Euphydryas bernadetta bernadetta* p. 256 clustered broomrape is *Orobancha fasciculata**, and beeplant is an error. *Polygonia interrogationis* and *P. comma* have winter and summer forms, which surely indicates that they must overwinter northward. Ssp. *cenveray* may occur in SD, as the book describes 2 rows of green unh spots. The *Polygonia “faunus”* photos are *Polygonia progne* winter form. For *Polygonia gracilis zephyrus* p. 268 swamp currant is *Ribes lacustre**. *Vanessa atalanta* is claimed in both Dakotas books to be just an immigrant, but I wonder if it (as well as *Vanessa virginiana* and *V. carye annabella*) often overwinters? *Oberfoelli* is a syn. of *Limenitis weid. weidemeyerii*. The Mountain Emperor is the new ssp. *Asterocampa celtis jeffermont* from Front Range of Colo. named by Scott & M. Fisher. *Lethe eurydice fumosus* surely occurs in most of the state (it occurs in S-C Minn.) except maybe the NE, and the photos are *fumosus*. In *Cercyonis pegala*, Porcupine grass *Stipa spartea* is a dubious H. Tietz host. In Colo., *Cerc. meadii* eats turflike sedges, while *C. oetus* uses *Poa pratensis/agassizensis* etc. *Neominois ridingsii* males are very active in early morning when they mate-locate by raiting on ridgetops. Ssp. *uhleri* (not *varuna*) occurs in S.D. evidently (Neb. has large ssp. *uhleri*). For *Erynnis afranius* p. 364, American deervetch is *Astragalus americanus** & should be called American vetch. For *Erynnis afranius* p. 364 & *E. persius* p. 366, goldenpea is *Thermopsis rhombifolia**. *Fredericki* is a syn. of *Erynnis persius*. For *Pyrgus scriptura* p. 368 & *P. communis* p. 370, scarlet globemallow is *Sphaeralcea coccinea**. *Amaranthus* is preferred by *Pholisora catullus*, and *Chenopodium berlandieri* is greatly preferred over *C. album*, if the latter is eaten at all. *Ancyloxypha numitor* eats hay grasses, so *Poa* is an error; *Spartina* is a known host, but *pectinata* was not known to me, and prairie cordgrass is *S. pectinata*, and S.D. has two sp. of *Spartina*, so is this a valid record or just data creep? *Stipa* is a known host of *Oarisma garita*, but *Stipa columbiana=Columbia needlegrass* is a literature error. Ssp. *phyleus* and ssp. *uncas* occur in S.D. The photos of *Hesperia comma assiniboia* look like *H. comma idaho* instead, leading one to believe that *assiniboia* doesn't occur in S.D., but I have a long series of typical *H. comma assiniboia* (unh tawnier & spots yellow) from Slim Buttes in Harding Co. Aug. 23 1970 (and 4 from Day Co. ½ mi. W Bitter Lake Aug. 25, 1971), so ssp. *idaho* doesn't occur in county unless it sneaks into the S part (maybe global warming is shifting *idaho* north and replacing *assiniboia* in Harding Co.??). The sheep fescue=*Festuca ovina* host for *Hesperia nevada* is really *Festuca idahoensis*. *Polites rhesus* hibernates as older larva. S.D. has *Polites peckius surllano*, with a small unh patch as shown on the photo. For *P. mystic*, Quackgrass=*Agropyron repens*, timothy=*Phleum pratense*, & *Echinochloa crusgalli* are all errors by H. Tietz based on *Polites themistocles*. Ssp. *logan* evidently occurs in E S.D., as Scott's specimens from Logan Co. in

NE Colo. are lagus, from Lincoln & Dawson Cos. in SW Neb. & Davis Mts. Tex. are lagusXlogan, and from Lincoln Co. on the W edge of Minn. are typical logan, while the male & female photos in this book look like lagus and near-lagus, so W S.D. evidently has lagus or near-lagus, while E S.D. surely has ssp. logan, just like N.D. Napa is a syn. of *Ochlodes s. sylvanoides*, and it eats hay grasses, so the literature host *Cynodon dactylon* is dubious; 1st-stage larvae hibernate. *Poanes hobomok* probably is a hay-grass feeder like *P. taxiles*, thus *Poa* is a dubious host, and is an error for *Poanes taxiles*. *P. taxiles* males rait in gulches. *Ssp. viator* occurs in S.D. The ssp. of *Euphyes vestris* are pathetically weak and worthless; *metacomet* may be a weak ssp. by having a slightly stronger unh lighter band, and would be the only ssp. in S.D. For *Euphyes vestris* p. 437, sun sedge is *Carex heliophila**. *Notamblyscirtes simius* immatures were desc. by Scott in *Papilio* (New Series) #6, unfed 1st-stage larvae hibernate, and males rait on hilltops in early morning. *Bouteloua curtipendula* is the main host of *Amblyscirtes oslari*. All the *Amblyscirtes* are basically hay-grass feeders. *Poa pratensis* is an error for *Amb. vialis* as my larvae died eating it, and *Cynodon dactylon* is an old lab-record error. *Leussleri* is a syn. of *Meg. streckeri texana* (though S. Spomer notes that Nick Grishin's Kerrville *texana* topotypes have sl. smaller spots and sl. thinner uph marginal band).

BUTTERFLIES EAST OF THE GREAT PLAINS. An Illustrated Natural History.

Paul A. Opler, George Krizek. 1984. Johns Hopkins Press, Baltimore, Maryland. 294 p.

This book is welcome, because it emphasizes natural history. It has more info on natural history than the outdated Klots Field Guide. It is a nice book, though there are some errors. It has 54 p. of 324 nice large color photos of living butterflies, but most of the less-common species have no photo, so identification will be difficult just using this book. The text has shaded distribution maps for each species, which are mostly accurate. Contrary to the title, it covers only E U.S., and not E Canada.

There is at least one misidentified photo. Photo 156 is *P. comma* not *interrogationis*; 166 is odd-looking and looks like *P. faunus* on ups but the margins are straighter than *faunus* so it evidently is *satyrus*; 174 is *Junonia genoveva* (the name was *genoveva* in my book, later switched to *evarete*, now switched back again); the postmedian unh lines of 204 are atypical of *Lethe appalachia* though the photo is correctly identified; 209 is evidently *N. helicta septentrionalis*; 272 looks like *P. themistocles*; 273 is probably *Polites themistocles* as *P. origenes* females usually have little upf costal orange.

Some published info on hibernation stage (*J. Res. Lepid.* 18:171-200) was not included. Based on Paul Opler's experience with *Eurema daira* etc. in Costa Rica, it is hypothesized that many southern U.S. species overwinter as diapausing adults, which deserves further research. Such species as *Zerene cesonia*, *Eurema nicippe*, *E. lisa*, *Phoebis sennae*, *Libythea*, & *Vanessa cardui* are often reported to overwinter as adults, yet they are also reported to overwinter as pupae (see *J. Res. Lepid.* 18:171-200); maybe their development is just slowed in winter in warm southern regions when they can be found in any stage. The exact northern extent of adult winter diapause is still not well studied in many *Vanessa* & *Nymphalis* etc.

Detailed comments: There are lots of distribution improvements and recent taxonomic changes that I won't mention. Times are given for mating and oviposition, as if these are genetic preferences, when mostly it just represents small sampling size of only part of the daily activity period. p. 12 female butterflies are larger partly because they have to emerge a few days later than males in order to synchronize virgin females with the maximum population of males (*J. Scott* 1977, *J. Animal Ecol.* 46:909-924). P. 14, S Fla. is subtropical, not tropical. P. 25 perching species visit mud too. P. 31 adults need more nectar in summer because it is hotter. *Eurytides marcellus* is a stray in Wis., where *Asimina* is absent. P. 50 "balanced polymorphism" of *Papilio glaucus* black females was disproved (*Science* 156:3774; *Ann. Ent. Soc. Amer.* 65:1267; *J. Lepid. Soc.* 26:109; *Evolution* 27:257), and the best theory is that black females benefit from mimicry, while frequency-dependent selection (birds learn to eat the black form if it becomes common), immigration of yellow females, and perhaps survival differences, cause the polymorphism. *Magnolia* is not eaten by *Papilio palamedes* larvae (*J. Brooks* 1962, *J. Lepid. Soc.* 16:198). F. Chew found that *Pieris oleracea* & *P. rapae* don't really compete much, so *rapae* did not displace *oleracea*. *Pieris virginiensis* doesn't occur in Ala. *Colias philodice* hibernates as 3rd-stage larvae. *C. interior* does not blend with *C. pelidne* in Man. *Phoebis sennae*: p. 70 *Chamaecrista* is a genus, a syn. of *Cassia*. *Phoebis agarithe* is identified by the straight unf line; *Pithecellobium guadalupense* is a syn. of *P. keyense* in Long & Lakela flora; *P. unguiscati* is an error of *H. Tietz* (based on *H. Dyar's* [*Proc. Ent. Soc. Wash.* 4:448] *P. guadeloupense*, which is a syn. of *P. keyense*). *Cassia bicapsularis* is a lab-only plant eaten by *Eurema nicippe*, based on *R. Kendall* (*J. Lepid. Soc.* 17:22). "Wild currant" is an error published before *Feniseca's* carnivory was known. *Lycaena phlaeas* was not introduced from Europe; half-grown larvae hibernate (*J. Res. Lepid.* 18:171). *Lycaena dione* & *thoe* mate-locate all day. For *L. thoe*, change the error *Rumex orbiculatus* to *R. verticillatus* (*orbiculatus* actually derives from *Shapiro's Butt.* Delaware Valley "Water Dock", in *Gray Flora Water Dock* is *R. verticillatus* & is just the 2nd name for *R. orbiculatus*, & *Shapiro Butt.* N.Y. uses only *R. verticillatus*). E Iowa is an error for *Lycaena epixanthe*. P. 88 summer *Strymon melinus* males have orange abdomens. *Aronia* was only suspected for *Satyrium titus* (by *A. Shapiro, Butt.* N.Y.). I think *Satyrium acadica* mate only late afternoon-dusk, and the few mating pairs I found in early morning I think just mated through the night. *Lysiloma* was just a lab host of *Ministrymon azia* (*R. Boscoe*), and is an assoc. record in Mex. (*S. Roman*). Oaks are dubious for *Calycopis cecrops*, & lack documentation. *Pinus clausa* was an assoc. host only for *Callophrys niphon* (based on *C. Zeiger*,

1980 Lepid. News #2 p.22, Fla.). *Fixsenia favonius ontario* is a ssp., which often lacks the orange up patches esp. northward. Move the W Wis. dot for *Erora* to E Wis. *Suriana maritima* may be an error for *Strymon istapa=columella*, that has been copied often since (Frank Fee [pers. comm.] did validly record *S. martialis* on it from Big Pine Key). *Hemiargus hanno antibubastus*: *Pithecellobium guadalupense* is a syn. of *P. keyense* in Long & Lakela flora; *P. unguiscati* is an error of H. Tietz (based on H. Dyar's [Proc. Ent. Soc. Wash. 4:448] *P. guadeloupense*, which is a syn. of *P. keyense*), and the record is based on H. *thomasi* (reported as *ammon*, from Lake Worth, Palm Beach Co. Fla.) so should be transferred to H. *thomasi*. *Cupido "Everes" comyntas*, change *Medicago sativa* to *M. lupulina* (Shapiro reported *lupulina* in Butt. N.Y., & Butt Rocky Mtn. States wrongly changed it to "alfalfa", which led to *M. sativa*). *Thermopsis* for *Cupido amyntula maritima* (not *valeriae*) was just an assoc. record from same book. *Celastrina* proved later to be 4 sp.: *Cel. lucia*, *C. ladon* (the 1st brood hosts listed in N Va.), *C. serotina*, *C. neglecta* (the 2nd brood listed in N Va.). *Cimicifuga* is the host of *C. neglectamajor* & sometimes *C. neglecta*. *Glaucopsyche*, spelled *Vicia caroliniana* in Gray flora. *Plebejus scudderii nabokovi* is all over N Wis. & N. Mich. I found *Libythea* mating pairs in daytime. Shapiro Butt. N.Y. wrote that *Viola septentrionalis* was only assoc. with *Speyeria atlantis*. *Salix herbacea* was assoc. only (*S. Scudder*, Butt. New England), and *Polygonum viviparum* only speculation, for *Boloria titania*. *Lysimachia* is an error for *Chlosyne gorgone*. *Phyciodes tharos* oviposits all day. *Phyciodes batesii* hibernates as unfed 4th-stage larvae, larvae are not pink, a partial gen. is rare if it exists anywhere, and several S Wis. records are errors (11 more errors in print from Iowa, Ill, Ind., Ohio). *Lonicera japonica* was a lab-only food of *Euph. phaeton* (A. Clark Proc. USNM 71 #2683; W. Saunders' Butterflies of Allegheny Park). The *Polygonia comma* black form occurs only in summer. *Vanessa cardui* usually pupates many meters from its host nest. *V. atalanta* ovip. all day, & hibernate as adults. P. 163 spelled *Ruellia nudiflora*. P. 167 *Quercus velutina* (Shapiro Butt. N.Y.) refers to ssp. *arthemis*, & *Vaccinium stamineum* is a dubious record of Abbot & Smith. Austin Platt (pers. comm.) told me *Lim. archippus* larvae are not colored differently on willow & cottonwood. *Apaturinae* is an older name than *Charaxinae* (the book combines them). P. 173 TL of *alicia* is New Orleans. W. Edwards described *N. areolata* immatures. P. 184 spelled *Carex alopecoidea*. If the two gen. for *Megisto cymela* are allochronic then they cannot interbreed and wouldn't be separate species anyway; *M. "cymela"* is still an uncertain mess, even though R. Gatrell declared that two sp. *M. cymela* & *M. eurytris* Maynard 1891 occur in S.C.; my specimens say that *viola* is a good ssp. with wider brown uns lines (J. Calhoun found that *viola* intergrades to *cymela* in Fla. panhandle); but I can't separate anything else despite my topotypes of *eurytris* & *cymela* caught by Gatrell, thus *eurytris* still looks like it is probably a syn. of *cymela*. Clark & Clark Butt. Va. wrote that male *Cercyonis pegala* fly L June-L Aug. and females fly E July-L Sep., while in Colo. males live to at least Sep. 7 & females to Sep. 13 though sex ratio after Aug. 15 is about 1:4; and Marc Epstein got Colo. females to oviposit in L July. *C. pegala* often visits flowers. *Erebia disa* just likes shade, and truly crepuscular butterflies are extremely rare or absent. *Strigulosa* is a ssp. of *O. calais*; leaning of adults is thermoregulation in *Hipparchia* (R. Findlay etc. Ecol. Ent. 8:145-153). Bigelow's sedge *C. rigida* is a syn. of *C. bigelowii* in Gray & Britton/Brown floras. P. 195 spelled *Asclepias exaltata*, & *Acerates* is a syn. of *Asclepias*. *Epargyreus clarus* usually ovip. on the host; males roost under leaves every afternoon after their mate-locating period; pupae hibernate. *Cogia outis* is native in Ark.-Mo. *Robinia pseudoacacia* is correct for *Erynnis icelus* (J. Burns' rearings reported in Butt. Ga. & Univ. Calif. Pub. Ent. v. 37). *Erynnis zarucco*: *Sesbania longifolia* is error, correct name is *Daubentonia longifolia* (from Mather & Mather J. Lepid. Soc. 13:72). *Salix* & *Populus* are errors for *E. persius*. *Althaea rosea*, *Abutilon*, & *Malva* are errors for *Pyrgus oileus* (they are based on H. Tietz' "syrichthus", strictly a syn. of *oileus*, but the name *syrichthus* was mistakenly used for *P. communis* when these hosts were first published, thus they really belong to *P. communis*). *Origanum* & *Marrubium* are errors for *Pholisora catullus* (& my larvae refused *Marrubium*); mating occurs all day. *Nastra neamathla* occurs from Fla. to C Tex. & *julia* from Tex. to Calif., and are ssp. according to Butt. Houston. P. 223 spelled *chicory*. *Hesperia uncas* is resident in Sherburne Co. Minn. near Minneapolis. *Hesperia comma* larvae are brown; eggs hibernate. *Hesperia leonardus pawnee* occurs in SW Minn. & W Iowa in typical form, and intergrades to *leonardus* are in E Minn. & W Wis. & NW Iowa; Ala. is error. *Hesperia metea*: *glomeratus* is *A. virginicus* var. *abbreviatus*, not a var. of *scoparius*. *H. sassacus*: *Festuca rubra* seems to refer to Shapiro's Butt. N.Y. host *F. nutans*, a syn. of *F. obtusa* in Gray flora. *Polites themistocles* eats many grasses. *Polites mystic* mates all day long. *Polites vibex*: *Stenotaphrum* was eaten in lab only (J. Lepid. Soc. 19:3). *Wallengrenia egeremet* mate-locates all day. *Atrytone arogos* once flew commonly in July in Fla. *Problema bulenta antenna clubs* are like *byssus*. Mich. *Euphyes dukesi* flies in July. *Cyperus* was lab food only for Heitzman's *Euphyes vestris*. *Linda* is a ssp. of *A. aenus*. The Ohio Amb. *aesculapius* dot is error. Amb. *reversa* has the same seasonality as *A. carolina*. Amb. *nysa*: *Echinochloa crusgalli* is error, J. R. Heitzman published *Ech. pungens* (J. Res. Lepid. 3:154), a syn. of *E. muricata* in Mo. flora, however Gray flora uses *E. pungens=muricata*. *Avena* (*S. Scudder*) & *Cynodon* (J. Lepid. Soc. 19:31) were eaten only in lab by Amb. *vialis*. *Calpodus ethlius*: *Thalia geniculata* may be an error, C. Zeiger (1964 Lepid. News #4) reported *T. dealbata* from "Fla."; *T. dealbata* occurs in N. Fla., and *T. geniculata* is a different introduced sp. in S Fla. The *Panoquina panoquinoides* early stages desc. is from Calif. *P. pan. errans*. The immatures desc. for *P. ocola* perhaps refer to *P. lucas=sylvicola*.

BUTTERFLIES AND MOTHS OF MISSOURI. J. Richard Heitzman & Joan Heitzman.

1987. Missouri Department of Conservation. Jefferson City, Missouri. 385 p.

Almost no misidentifications of the butterflies! For what this book aims to do, it is a very good book. It covers all the butterflies, and some of the more beautiful, common, or pestiferous moths. Each has color photos, and comments about distribution status, hostplants, larva description, and misc. comments. Ssp. are given only if they are not the nomenotypical. There are 7 color pages of larvae/pupae photos. My main gripe is that the text is skimpy for most species. Richard Heitzman is a skilled experienced lepidopterist, who has lots of interesting information, which I wish had been included.

Detailed comments: P. 34 the “lucilius” photos are Sask. E. afranius. P. 48, Mo. has the later-named P. peckius surllano. P. 68 linda is ssp. of A. aenus. P. 85 joanae is a ssp. of P. machaon, being a bairdii-bug that has introgressed with P. polyxenes a lot, leading some to consider it a syn. of polyxenes, but it is evidently valid, with eyespot & mtDNA of bairdii. P. 366-7 has photos of joanae larvae, which (S. Spomer & Heitzman pers. comm.) have unicolorous light-green or bluish-green forms (polyx. larvae are paler ventrally) with paler/absent spots (polyx. spots orange) & young larvae with a light-yellow saddle (polyx. whiter). P. 96 menapia must have been an unnatural truck/plane import. P. 122 mopsus is a syn. of titus and Mo. has Satyrium titus watsoni=campus. P. 135 Phaseolus misspelled. P. 138 H. thomasi was surely misidentified. P. 141 Mo. has Cel. ladon (spring), C. neglecta several generations later in year, & C. neglectamajor in Ozarks. P. 164 photos show later-named P. phaon jalapeno. P. 169 progne photo is summer form female. P. 174 Cirsium. P. 175 atalanta adults hibernate. P. 183 A. troglodyta floridalis is very dubious (misidentified?). P. 192 hermes is just a stray if it only occurs July-Sept. SW Mo. has C. pegala texana if you think it differs from C. pegala maritima “alope”; why doesn't the text explain the creamy Salem Uplift form which looks like cream C. p. carolina from the Carolina Mts.?, much too skimpy. P. 198 Cynanchum is error (it is a host of D. gilippus).

THE BUTTERFLIES OF IOWA. Dennis W. Schlicht, John C. Downey, Jeffrey C. Nekola. 2007. University of Iowa Press, Iowa City, Iowa. 233 p.

This book has some interesting features. Schlicht & Nekola are mostly interested in butterfly conservation, and have scoured Iowa to find what is left of the fauna, an admirable and arduous task because most of Iowa has been scraped bare to plant soybeans and corn. They have found 170 fens in NE Iowa for instance (160 remain, p. 100), and state that 90% of Iowa's fens have been drained (p. 98). A good lesson for lepidopterists and land managers is repeated often here, as they write that the new practice of burning small prairie tracts (to prevent shrub and tree succession) has burned up colonies of *Hesperia dacotae*, *Oarisma poweshiek*, *Atrytone arogos*, & *Coenonympha tullia*, & fire is stated to reduce *Speyeria idalia* pops. Also, *Hesperia ottoe* was “exterminated by management practices”, and *Plebejus melissa* declined on managed reserves. Obviously, to remove shrubs/small trees land managers should **MOW** these small prairies in the fall instead of burning them, as I wrote in my 1986 book for *Hesperia dacotae* (& grazing is said to help *S. idalia*). P. 22 notes that 50000 of A. F. Porter's butterflies were eaten evidently by dermestids (the fate of the Stallings & Turner megathymid collection--lepidopterists, give your collections only to large museums with guaranteed endowments!).

The main trouble with the book is that the authors did not consult experts or research the literature enough. In particular, my two hostplant papers (in *Papilio* [New Series] # 6 in 1992 & #13 in 2006) gave many hostplants and life histories that I found in about 20 yearly visits to my sister's farm near Albert Lea, Freeborn Co. Minnesota, only a few mi. N of Iowa. Many of the questions that they ask in this book were already answered in those two papers. (At the end of each species account, they ask Questions, which is rather annoying—in a book I want answers, not questions.) When I first got the book I turned to the *Phyciodes* with grim expectation, and there it was: *Phyciodes diminutor* Scott 1998 was missing, even though its TL (in *Papilio* [N.S.] #10) was 10 mi. N of Iowa, and I caught it at the Minnesota Welcome Center on I-35 just a stone's throw N of Iowa, and then I turned to the *Phyciodes* plate p. 167, where I saw that the upper two photos are actually a male *Phyciodes diminutor*, which p. 213 writes is from Rowley Fen, Buchanan Co. Iowa June 15 1985 D. Schlicht; that male is obviously *diminutor* as the antenna nudum is orange, size is large, there is a large uph space without line, and unh has large russet marginal patch. Obviously *P. diminutor* occurs all over NE Iowa, where it must eat mostly *Aster simplex*; true *P. tharos* occurs all over the state.

The book skimps on taxonomy, as the ssp. names of many sp. are not given. The nomenotypical ssp. occurs in Iowa in *Epargyreus clarus*, *Thorybes pylades*, *Erynnis brizo*, *juvenalis*, *Hylephila phyleus*, *Polites origenes*, *Poanes hobomok*, *massasoit*, *viator*, *Euphyes dion*, *Atrytonopsis hianna*, *Battus philenor*, *Papilio glaucus*, *troilus*, *Pontia protodice*, *Colias philodice*, *Zerene cesonia*, *Callophrys gryneus*, *henrici*, *Strymon melinus*, *Danaus plexippus*, *Speyeria cybele*, *Boloria bellona*, *Chlosyne nycteis*, *Phyciodes tharos*, *Vanessa cardui*, *Limenitis archippus*, *Asterocampa celtis*, *clyton*, *Hesperia metea*, *Eurema mexicana*. Other missing ssp. are added below.

There is a list converting plant common names to scientific names, but wild beans p. 45 isn't on the list. J. Burns' histograms seem to show just two gen. of *Erynnis martialis* (Apr.-M June being the 1st). *E. baptisiae* has three gen. in Neb. according to Steve Spomer, not four, and Burns lists Sept. records for Ill. & Ohio where there must be 3. Mint is an error host of *Pholisora catullus*. *Ancy. numitor* eats 5 sp. of hay grasses in S Minn., and bluegrass is an error. *Polites peckius surllano* is the ssp. in Iowa, as the photos show; I think turfgrasses such as *Poa pratensis* are the main hosts, as the eggs are laid without glue. Scott gave several hosts for *Polites themistocles*, and he showed that *Andropogon gerardii* is the main host for *P. origenes*. The types of *A. arogos iowa*—not the genus--were from Iowa. *Anatrytone logan* larvae overwinter

half-grown. *Problepsis byssa kumskaka* occurs in Iowa. Bluegrass is dubious as a host of *Poanes hobomok*, which probably eats hay grasses as does *P. taxiles*. *P. zabulon* surely has 2 gen. at most. It's very interesting that *Poanes massasoit* has a disjunct NW range. Scott found *Poanes viator* & *Euphyes dion* larvae on the same parts of *Carex lacustris* just N of Iowa. *Buchholzi* is a syn. of *E. conspicua*, based on my Neb. & Minn. series. *E. bimacula illinois* occurs in Iowa. *Euphyes vestris* has two gen., and Scott found it eating dryland *Carex pensylvanica* which occurs in Iowa. Kentucky bluegrass is dubious for *Amb. hegon*, which surely eats hay-grasses like all other *Amblyscirtes*. *Amb. vialis* overwinters as larvae, which eat various hay grasses. *Pontia protodice* is rare in S Minn., and it evidently strays N in spring. *Phoebis sennae marcellina* occurs in Iowa. Based on wing pattern & DNA, *Lycaena phlaeas hypophlaeas* is native in E U.S. and evolved from arctic or Siberian ancestors; it was not introduced. The 3rd L. *helloides* gen. is probably Sept., and it is Sept. in Neb. also according to Spomer. Evidently *Satyrium titus watsoni=winteri=campus* occurs in Iowa. *S. calanus falacer* & *S. liparops strigosa* occur in Iowa. Scott reported other hosts of *Cupido comyntas comyntas* in Iowa. The photo looks like *Glaucopsyche lygdamus afra* rather than ssp. *couperi*. *Libytheana carinenta bachmanii* occurs in Iowa. *Speyeria cybele alcestis* (a ssp.) & *aphrodite* & *idalia* (the weak ssp. *occidentalis*) surely just have one gen., with females aestivating until later in summer. *Boloria selene nebraskensis* occurs in Iowa. Both *Phyciodes* overwinter as unfed 4th-stage larva. *Polygonia progne* 2nd gen. is probably L July overwintering to E May. *Vanessa atalanta rubria* occurs in Iowa. *Iowa arthemis* is just a rare polymorph in the *Lim. arth. astyanax* population. *Lethe eurydice fumosus* is widespread from Iowa to Colo., and also occurs in wet roadside ditches in S Minn., where its host is *Carex aquatilis*. *Coenonympha tullia benjamini* occurs in Iowa. *Cercyonis pegala nephele* occurs in Iowa, and probably has just one gen. Six of the species listed in "Occasional Strays & Old Records" are uncommon strays but validly recorded. *Leptotes marina* surely overwinters much farther north than coastal Texas (probably in Colo.). *Danaus gilippus thersippus* occurs in Iowa. *Dione vanillae incarnata* occurs in Iowa.

The maps are nice, and old and new records are distinguished by symbols to help document the disappearance of some species. A map of county names would have been useful.

The color plates are nice, and only one was misidentified (the *Phyc. diminutor*)!, although *Hesperia leonardus pawnee* & *leonardus* photos are transposed. The 2nd *Erynnis lucilius* & *Oarisma poweshiek* & *Euphydryas ph. phaeton* are also males. The female *Ancy. numitor* is above the male. Both *Call. gryneus* are females. The lower left *Sat. acadica* is male, lower right female. The *Polygonia interrogationis* photos are summer, then winter form; *P. comma* are winter, then summer f; *P. progne* all summer f.

BUTTERFLIES OF WISCONSIN. James A. Ebner. 1970.

Milwaukee Public Museum, Popular Science Handbook #12. Milwaukee, Wisconsin. 205 p.

This book is somewhat out of date, so I won't make many comments. There are not many distribution records, there are no maps and just a few counties are mentioned in text for most species. The book has good b/w photos, and there are almost no misidentifications. P. 58 the *Euchloe olympia* name *anniha* is a nomen nudum that was never published. The photo of "male *Satyrium calanus falacer*" is a female *S. caryaevorus*. *Plebejus scudderii nabokovi* occurs in Wis. (the photos of *scudderii*) from Oconto, Marinette, Brown, Waupaca Cos. etc., and *P. samuelis* occurs in Wis. in Burnett & Wood Cos. etc. (see *Papilio* [N.S.] #12). The photos look like *Celastrina neglecta*, & Wis. also has *Cel. ladon*, and perhaps also *C. (argentata) serotina*. P. 108 *Junonia coenia* doesn't eat *Sedum*. *Polygonia satyrus* occurs in Vilas & Sawyer Cos. etc. The upper *Phyciodes "tharos"* male is *P. cocyta selenis*, while the female looks like *P. tharos*; both occur in Wis., and *P. diminutor* surely occurs in S Wis. P. 134 *Argynnis (Speyeria) aphrodite* female *uns* is ssp. *alcestis*, which is mentioned on next p. from S Wis. *Dione vanillae incarnata* probably strays to Wis. rather than ssp. *nigrior*. *Lethe appalachia* is also in Wis. Wis. has *Coen. tullia inornata*, & *O. calais strigulosa*, & *O. jutta alaskensis=ascerta* (not *ridingiana*). *Wallengrenia egeremet* & *Polites peckius peckius* occur in Wis. Mature larvae of *Pyrgus communis* hibernate. P. 190 *Erynnis persius* does not eat *Salix* or *Populus* (errors based on *E. icelus*).

MICHIGAN BUTTERFLIES AND SKIPPERS. A Field Guide and Reference.

Mogens C. Nielsen. 1999. Michigan State University Extension, East Lansing, Mich. 248 p.

This is a good book. It has little color photos of adults and shaded-county distribution maps with each species, and the text covers the usual information that most people want. It has some caterpillar photos too. The hostplants are quite accurate, though the common name>scientific name translation on p. 242-8 gives a few more hostplant species than are found in text. There are almost no misidentifications! I would have liked more added comments and local expertise. Many of the adult photos are too orangish.

Detailed comments: *Battus philenor* is not threatened, it is just a rare stray. (Many of the so-called threatened species are really just corner weeds: very common in their whole range, but entering just a little corner of your artificial political unit, thus very limited in your phony biogeographic jurisdiction. Michigan is a totally artificial state, as the "northern peninsula" and "southern peninsula" aren't even connected, and Isle Royale is way off.) *P. polyx. polyxenes* is from Cuba (where it may have never existed as a ssp!). The distribution maps of *P. glaucus* & *P. canadensis* show near-total allopatry,

which is silly as they really intergrade and forms resembling the other are found widely within each other's distribution (even W of Denver in Colo. we have a few stray *glaucus* and some *glaucus**X**rutulus* intergrades). I wonder if the two *Pontia occidentalis* were just misidentified? The *Colias philodice* & *Phoebis sennae* & *Eurema lisa* photos are too orange. *Lycaena phlaeas* was not introduced from Europe. *Satyrium titus watsoni*=*winteri* evidently is the ssp. in Mich. The *Celastrina* seem to be correctly sorted into three sp. *C. ladon*, *C. lucia*, & *C. neglecta*, though the mating pair of *C. ladon* looks like it might be *C. lucia*; a fourth sp. *Cel. (argentata) serotina* probably occurs in N Mich. also, eating *Diervilla* probably. *Podophyllum*=*may-apple* & *Portulaca*=*purslane* are errors for *Euptoieta claudia*. *Argynnis (Speyeria) cybele* oviposits near violets, not on them. *Boloria eunomia* flies more than 2 weeks, and I haven't seen "dogfights". *Bol. freija* doubtfully hibernates as pupa. Most *Phyciodes* "tharos" photos are *P. cocyta selenis* (*selenis* has large uph orange space & orange nudum on elliptical club, & large paler unh marginal patch); only the live male photo and p. 127 bottom left female look like *tharos* (and the latter lacks abd. tips), so *P. tharos* & *P. cocyta* were evidently confused in this book, thus the geographical replacement of them on the maps is surely wrong. Also, *P. diminutor* seems to occur in S Mich. & N Ohio (see *Papilio* [N.S.] #13); it is similar to *P. cocyta* but has two gen. June to Sept. *P. tharos* has 3 gen. in S Minn. and probably in Mich. too. The *P. cocyta selenis* photos are correct. *P. batesii lakota* occurs in Mich.; it has one gen., the larva is not pinkish, & antenna club is almost always blackish. The broods of *Polygonia* & *Nymphalis* aren't stated well, as the 1st gen. overwinters from the previous Sept. The lowest *P. faunus* photo is *f. silvius*. Seasonal forms of *P. progne* should have been mentioned (ups photo is winter form). *Vanessa atalanta* adults hibernate. *P. 149* wild snapdragon does not exist, as *Antirrhinum major* & *A. orontium* are from Europe. *Lethe creola* was an error in Mich. Mich. has *Oeneis calais strigulosa*. The male *Thorybes "bathyllus"* photo looks like *pylades*. Which 2 sp. of *Baptisia* does *E. baptisiae* ovip. on? Mint=*Mentha* is an error for *Pyrgus communis*. *Ancyloxypha numitor* ovip. on various hay grasses. *Oarisma poweshiek* doubtfully eats *Eleocharis*=*spike-rush*. Mich. has the large-patch *P. peckius peckius*. Redtop grass for *Pompeius verna* means *Agrostis gigantea* on p. 246 (even though Gray flora says Redtop is *A. alba*). *Anatrytone logan* & *Poanes hobomok* doubtfully eat *Poa*. *Poanes viator* & *Euphyes dion* larvae are known and feed on *Carex lacustris* in Minn. *Cyperus* was just a lab host for *Euphyes vestris*. *Poa pratensis* is dubious for *Amblyscirtes hegon* & *A. vialis*, as *Amblyscirtes* eat hay-grasses. Some hosts were missed on p. 242-8: Aspen is *Populus tremuloides*, Broccoli & Cauliflower are *Brassica oleracea* var. *botrytis*, Brussels Sprouts is *Brassica oleracea* var. *gemmifera*, cabbage is *Brassica oleracea* var. *capitata*, Kale is *Brassica oler.* var. *acephala*, cherries are *Prunus* spp., Clovers *Trifolium* spp., common mallow=*Malva neglecta*, Dock=*Rumex*, Dogwood=*Cornus*, hard pines=*Pinus* spp., Holly is misalphabetized, Indian Grass=*Sorghastrum nutans*, lawn grasses are *Poa pratensis* etc., Pansy is *Viola tricolor* var. *hortensis*, Pennsylvania sedge is *Carex pensylvanica*, Redtop grass is *Agrostis*, Sumac is *Rhus* spp., sunflower is *Helianthus* spp., Wild pepper-grass is *Lepidium*, wild senna is *Cassia*.

THE BUTTERFLIES OF INDIANA. Ernest M. Shull. 1987.

Indiana University Press, Bloomington, Indiana. 262 p.

This is a good book. The color photos are nice. The county-dot maps are very good. Interestingly, it reports numerous times of observations of mating pairs (all EST), which provides conclusive evidence of mating time (406 copulating pairs of *Pieris rapae* from 9:30-6:30 for instance!), whereas most authors try to ignore mate-locating and mating behavior; his data for *Phyciodes tharos* matings clearly show the genetic ability to mate all day and the temperature-dependent increase in frequency in warm parts of the day.

Detailed comments: *Epargyreus* has two gen. if it flies May-Sept. P. 29 *Vernonia*. *Lespedeza tExana* was lab-only for *Achalarus lyciades*. For *Thorybes bathyllus*, the hosts are *Strophostyles*, *Rhynchosia tomentosa*, *Rhynchosia virginiana*, & *Tephrosia ambigua*. Pl. II all 4 E. "horatius" are *E. juvenalis*. *T. pylades* evidently has one gen. Wild indigo, witch hazel, & oaks are errors for *Erynnis icelus*. P. 41 *Geoffroea*. Pl. III both *E. lucilius* are *E. juvenalis*. *Pyrgus communis* evidently is somewhat migratory. The *Iherminier* are females. *Salix* & *Populus* are errors for *Erynnis persius*. *Digitaria* is an error for *Hesperia sassacus*. P. 71 *Agrostis* not *Agretis*. P. 75 *Oryza* not *Orza*. *Colias eurytheme* hibernates strongly in Utah, mostly as 2nd-3rd-stage larvae. *Trifolium* was used only in lab by *Phoebis sennae* & *Eurema lisa*. Ind. evidently has *Satyrium titus watsoni*=*winteri*. Ind. has *Lycaena dorcas michuron*. *Galium* is an error for *L. helloides*. Pl. XXIX 2nd row 3rd is *S. calanus falacer*. P. 148 *Quercus ilicifolia* does not occur in Ind. P. 150 *Rhus copallina*. *Callophrys niphon niphon* occurs in Ind. Pl. XXX 2nd row 4th is *S. melinus* female. *Crataegus* is an error for *Satyrium favonius ontario*. Pl. XXXI all *Celastrina* are *C. neglecta*. Indiana has *C. neglecta* all over, *C. neglectamajor* in SE Ind., & *C. nigra*. P. 164 Irwin Leeuw. All the hosts of *Plebejus samuelis* are for *P. melissa* except for *Lup. perennis*. Adult *Libythea carinenta bachmanii* evidently hibernates. Pl. XXXV last specimen is *S. aphrodite alcestis*, which is evidently just a form in Ind. Pl. XXXVI last bug is male. *Aster* is not a host of *Chlosyne nycteis*, which has two gen. *Phyciodes tharos* larvae are not green; the photos are *P. tharos*, but *Phyciodes diminutor* probably occurs in N Ind. The photo of *P. batesii* is *P. batesii lakota*, but the 4 dots in text are surely *tharos*. *Wisteria* is an error for *Euphydryas phaeton* (Clark wrote that lab larvae refuse it). P. 195 *Ulmus*. Pl. XLI 5th row left is *P. progne*, so delete *zephyrus* from the book & Ind. P. 200 & 209 *Boehmeria*. *Salix* is error for *Aglais milberti*. *Ludvigia* & *Sedum* are errors for *Junonia coenia*. *Lethe creole* is surely an error in N Ind. *Cynodon dactylon* is lab food only of *Cyllopsis gemma*. Evidently *Cercyonis pegala maritima* occurs in Ind. (R. Gattelle found that topotypical *alope* is a syn. of ssp. *pegala* [in the blend zone *pegala**X**carolina*], thus *maritima* replaces the usual usage of

alope (maritima is older than ochracea which is an infrasubspecific pegalaXcarolina intergrade & carolina is a cream-patch ssp.) thus *maritima* = "alope"; *abbotti* is a syn. of *pegala*). However, *C. p. maritima* intergrades toward uncommon forms like *C. p. nephele* in N Ind. (=olympus). *Danaus plexippus androconia* do not attract females (the male grabs her with his legs for mating).

BUTTERFLIES AND SKIPPERS OF OHIO. David C. Iftner, John A. Shuey, John V. Calhoun. 1992. Ohio State University, Columbus, Ohio. 215 p.

I caught some of my first butterflies in Ohio, near my grandparents' houses near Cleveland and near Green Springs. I caught a Giant Swallowtail on the fragrant soil of a cow pen, and put it into a jar stuffed with grass because I didn't know how to dispatch and preserve it; it's still in my collection. Later I found *Argynnis* (*Speyeria*) *idalia* near Cleveland. This book is good. It has a lot about Ohio's habitats and glaciation. Despite the tiny counties in Ohio, there are a lot of records, from 35-97 species records per county, and the maps show fairly complete coverage of potential ranges. A flight period histogram is given for each species, based on >25,000 records. The usual hostplants are listed plus some new ones, along with an impressive list of nectar plants used by each species. A lot of work was done on sedge-meadow & other wetland species. The book mentioned how *Lycaena epixanthe* might have been widespread before the cranberry bogs were all drained. It must be frustrating living in Ohio: lots of common widespread species, but many species are scarce or nearly-extinct & impossible to find. The book usually uses the annoying "territorial" word to refer to what I used to call perching behavior to await females, which I now term raiting behavior.

Detailed comments: P. 11 says the Xerothermic was 8000-4000 years ago, p. 12 says it ended 5000, and I understood it was even earlier. Oaks are dubious for *Erynnis icelus*. P. 43 Findlay. *Polites mystic* host is usually *Poa pratensis*, though sedges could be eaten also; hosts have to be turflike on non-watery soil as the eggs are laid without glue and drop into the litter. *Huron* is a syn. of *Atalopedes campestris*. *Carex* is dubious for *Anatrytone logan* also (sedge meadows usually have *Poa pratensis* all through them, unless they are the wet ones). I reared *Poanes viator* from *Carex lacustris* in Minn. Pupating *Pap. polyxenes* larvae surely wandered onto *Cosmos*. *Pap. glaucus* balanced polymorphism was disproved. The *Colias eurytheme* records increase until Sept., in contrast to the steady *C. philodice*, evidently because it likes alfalfa more (do some disperse into Ohio later in the summer?; it is considered a resident in contrast to *Junonia* so can't migrate much). *Lycaena phlaeas* was not introduced from Europe. Ohio would have *Lycaena epixanthe phaedra*. Ohio has *Satyrium titus watsoni*=*winteri*. *Satyrium acadica* males rait in late afternoon-dusk, thus are less-active earlier. The book's *ladon* combines three species: Ohio has the spring *Celastrina ladon* over at least the SE ½ of the state, *Cel. serotina* on E edge in Columbiana Co. (pl. 24 row 6), & the June-Sept. *Cel. neglecta* all over. *Cel. neglecta* major lacks hyphen. *Celastrina nigra*=*ebenina*. P. 117 writes that *Speyeria diana* is reestablishing itself in some areas such as W.Va. *Boloria selene nebraskensis* exists even in roadside ditches in Minn., so should be able to survive in Ohio. *B. bellona* has very fast development, so there are 3 gen. Ohio has *Phyciodes tharos* all over the state in 3+ gen. In *Papilio* (N.S.) #13 I showed that *Phyc. diminutor* also occurs across N Ohio (Lucas, Wood, Columbiana Cos., from a study by Porter & Mueller 1998; & Sandusky Co. [this book pl. 28 row 5 middle]) and flies L May-E June & L July-E Sept. in Ohio in several gen.; antenna club is usually orange, but mostly blackish in Columbiana Co. *P. cocyta selenis*=*pascoensis* has one gen. and does not occur in Ohio (the mapped dots from Highland & Hamilton Cos. I think are *P. tharos*, and the 1m1f from Elk Co. Pa. on Pl. 29 also look like *tharos*). We need a thorough study of Ohio *Phyciodes* incl. rearing from eggs/young larvae found on natural hosts. Does *Nymphalis antiopa* have one gen., or one emerging June-E July and a second emerging L July-Aug. that overwinters? The *Aglais milberti* histogram may suggest 3 gen. Note that *Vanessa virginiensis* & *V. atalanta* seem to overwinter (as adults surely). *Cercyonis pegala maritima* replaces the name *alope*. The *Megisto cymela* histogram does show two emergence peaks in E June & E July, the mysterious dual cohorts. P. 153 *Aster undulatus* not *umbellatus* is a *P. batesii* host. P. 165 MOR is used for *Morrow* on pl. 24 at least. Pl. 24 row 5 has 3 *Cel. ladon*, 2 *Cel. lucia*; row 6 is *Cel. serotina* from Columbiana Co. (according to O.D. Pavulaan & Wright 2005 Tax. Rept. 6:7); row 7 is *Cel. neglecta*. Pl. 28 row 5 2nd, is *Phyc. diminutor* from Sandusky Co. (the others are *P. tharos*). Pl. 29 1st row may be *P. cocyta selenis*. Pl. 40 *P. satyrus* were later named *P. s. transcanada*.

BUTTERFLIES OF NEW JERSEY. A Guide to their Status, Distribution, Conservation, and Appreciation. Michael Gochfeld & Joanna Burger. 1997.

Rutgers University Press, New Brunswick, New Jersey. 327 p.

This is a good book, which is quite different from the usual butterfly book. There are only a few color photos of adults, and readers will need other books to see what the butterflies look like. But there is lots of local information specifically pertaining to New Jersey butterflies, especially distribution and flight times, and historical changes in distribution are discussed while citing previous works on N.J. species. *Panhiades* & *Calycopis* are expanding northward for instance. Sometimes the cited references on distribution are all that's known, for instance the authors offer little else on *Erynnis lucilius* occurrence, which remains mysterious. There is good info on habitats & climates in the state. N.J. has totally artificial state boundaries, but the book is a must-have for area butterfly fanciers anyway. Published hostplants, hibernation

stage, and other comments are listed for each species. The book analyzes census techniques & 4th-of-July butterfly counts. It discusses how forest overgrowth can choke out butterflies, and how the spraying for Gypsy Moths damaged lepidoptera (*Amblyscirtes hegon*, *Panoquina panoquin*, etc.) more than that moth. It's interesting that forest succession threatens *Lycaena epixanthe* and its cranberry bogs, and *Calephelis borealis*, etc. Detailed comments:

P. 29, later Scott & others split them into *Argynnis* (*Speyeria*) *atlantis* & *S. hesperis*. P. 43 *Calycopis cecrops* was later found to eat a wide variety of detritus. P. 55 hilltopping is just one type of mate-locating site, and gulching, tree-topping, etc. could also have been discussed. P. 102 *Daucus carota* flowers are not popular for nectar. P. 104 transplantations of butterflies should be done more often, for instance *Melitaea phoebe* should be transplanted from Eurasia to the west (Montana-Colorado etc.) to control weedy *Centaurea*; and lots of endangered/threatened species should be introduced to new suitable sites. P. 123 *Papilio polyxenes* has 1:1 sex ratio. P. 125 *P. glaucus* balanced polymorphism was thoroughly disproved. *Papilio palamedes* is not tropical. *Colias philodice* larvae hibernate, and *C. eurytheme* does too in Utah, even though *S. Ae* claimed it doesn't have much of a diapause in Conn. *Colias eurytheme* gradually colonized the northeast, so it must overwinter there, and is not just a migrant as various other books suggest, though people have seen *eurytheme* migrate in Kansas & Wash. etc. P. 139 *eubule*. Half-grown *Lycaena phlaeas* larvae hibernate. It's silly to want *Lycaena* *thoe* to be declared as threatened, as it is common in most of N.A. and is expanding in W U.S. Christopher Curtis found *Lycaena epixanthe* at a commercial cranberry bog at Whitesbog N.J. in the 1960s. *Callophrys irus* larvae are reported to be unmarked on *Lupinus* & striped on *Baptisia*; are those different taxa? Roy Kendall proved *Diospyros texana* as a host of *Call. henrici*. P. 159 *Pinus banksiana*. The *Celastrina* writeups are good, though p. 172 gives the impossible statement that *Cel. neglectamajor* eggs are found prior to adults even though pupae hibernate with a single gen. (if adults fly M May-E June then look for eggs in E June). P. 175 Scott & Wright 1990 (in Otakar Kudrna's Butt. Europe vol. 2) showed that *Heliconiini* includes *Argynninae*. 2450 eggs for one *Speyeria idalia* female! *S. idalia* may have spread east from the plains when the forests were cleared, as p. 183 notes for *Boloria bellona*, and now it's retreating as the forest regrows. P. 183 *Lysimachia*. *Phyciodes selenis* was named by Kirby in 1837, and *P. cocyta* is the proper name. One specimen is not enough to put *cocyta* in N.J., however P. Opler told me in 1998 that he has seen a series (of *Phyc. diminutor*?) from Sussex Co. in a shale barren with *Calephelis borealis*. *P. batesii* is rather doubtful from N.J. and most or all may have been *P. cocyta*; I have not studied specimens in eastern museums. P. 189 all *Polygonia* & *Nymphalis* & *Aglais* feed regularly on flowers. *P. progne* is generally rare so doubtfully has declined; does fall brood emerge in M Aug. or Oct.? P. 194 top is poorly written on one *N. antiopa* gen., & whole book is written poorly on overwintering adult "broods". Adult *Vanessa atalanta* & *V. virginianensis* hibernate. P. 201 *weidemeyerii*. *Lethe eurydice* & *appalachia* hibernate as larvae. N.J. was later declared to have *Neonympha helicta septentrionalis*. This book cites USFWS reports saying that collecting helped exterminate *Neonympha mitchellii* in N.J., evidently the source of Glassberg's vitriolic comments about collectors in his books; it also blames natural succession to upland forest, which Ray Stanford told me caused the extirpation of one N.J. site. *Megisto cymela*'s hosts have been poorly studied so the host range is undetermined. P. 211 the 2nd gen. is produced by the 1st gen. in Ont., contrary to Brown. The variation in *Cercyonis pegala* is polymorphism. Scalloped Sootywing is the best name for *S. hayhurstii*. P. 226 maybe the identification was wrong of the garden columbine *Erynnis*. Legumes are the only proven valid hosts of *E. persius*. Pesticides are blamed for exterminating *Pyrgus centaureae* *wyandot*. *Ambrosia* is an error for *Pholisora catullus*, & *Poa* an error for *Ancyloxypha numitor*. *Polites peckius* larvae hibernate. *Polites origenes*: Scott 1986 did not mention Mich., & *Andropogon scoparius* was my error for *And. gerardii*. Shapiro only wrote that *P. clandestinum* was "suspected" for *Wallengrenia egeremet*. P. 245 bottom, page priority is not in ICZN Code. P. 248 *Poanes taxiles* is in Rocky Mts. *Poa*, *Avena*, & *Cynodon* were lab food only for *Amblyscirtes vialis*. P. 257, *Cymaenes tripunctus* has ~7 spots. The *Panoquina panoquin* host on *Scirpus* was an error (A. Shapiro pers. comm. to Scott, 1980).

BUTTERFLIES AND MOTHS OF NEWFOUNDLAND AND LABRADOR. Ray F. Morris. 1980. Research Branch, Agriculture Canada, Publ. 1691. Ottawa, Ontario.

This book has good distribution maps for Newfoundland., but it covers Labrador rather weakly. Butt. Canada helps update this book. No records are given for *Papilio brevicauda* in Labrador. Only *P. glaucus canadensis* occurs in Nfld. (the black females were mislabeled, D. Ferguson J. Lepid. Soc. 37:189). Hebron is dubious for *Colias interior*. For *Colias nastes*, fig. 10 is a moth larva with tubercles, which all *Colias* lack. Only *Coenonympha tullia mcisaaci* occurs in Nfld. *Oeneis "chryxus"* does not occur in Nfld. *Erebia mancinus* occurs in Labrador. *Boloria titania arctica* occurs in Lab. & Nfld. *Chlosyne harrisii* does not occur in Georgia. Nfld. has *Phyciodes cocyta cocyta=arctica*. *Betula* & *Salix* are errors for *Polygonia gracilis*. *Polygonia progne* does not occur in Siberia. *Callophrys eryphon* does not occur in Nfld. or Labrador. Newcomb reared many *Lycaena dorcas* to adult stage on "*Potentilla*" *fruticosa*. *Lycaena epixanthe* does not occur in BC. *Plebejus glandon labrador* was later named from Labrador. *Celastrina lucia* occurs in the area. *Pyrgus centaureae* larvae do not eat grasses.

BUTTERFLIES OF DELMARVA. Elton N. Woodbury. 1994. Tidewater Publishers, Centreville, Maryland. 138 p.

A nice little book. Elton was the best butterfly photographer in America; all his photos are magnificent. He used a ~100-mm macro lens and powerful off-camera flash, and used experience to set his f-stop (see p. 104-5). There are photos of most adults, and of many immatures. The book contains all the info included in field guides, but contains little or no information on distribution within Delmarva. Skippers are not included. (Delmarva is a peninsula containing parts of Delaware and Maryland, between Chesapeake Bay & the Atlantic Ocean-Delaware Bay.)

Detailed comments: P. 45 *Colias philodice* does not prefer *Trifolium*. *L. phlaeas* was not introduced from Europe. P. 52 water dock is *Rumex orbiculatus* in glossary but this is an error as Shapiro's Butt. N.Y. "water dock" was *R. verticillatus*. Delmarva evidently has *Celastrina ladon*, *C. neglecta*, & *C. idella*; maybe the photos are *C. ladon*. May apple & purslane are dubious hosts of *Euptoieta claudia*. Dogbane is not a hostplant of *Danaus plexippus*. P. 101 petunias are not visited by butterflies for nectar.

BUTTERFLIES OF GEORGIA. Lucien Harris Jr. 1972.

University of Oklahoma Press, Norman, Oklahoma. 326 p.

This book was good, but is now out of date, so I will make just a few corrections.

Detailed comments: P. 3 Abbot lived 1750-1840. *Monarda* & *Origanum* are errors for *Pholisora catullus*. Pl. 5#21-22 are *Celastrina neglecta*. Pl. 8#9 is *Argynnis* (*Speyeria*) *cybele* female. B/W Pl. 6#1-2 look more like *Euphyes arpa* males as unh lacks pale veins. Pl. 10#12 is male. Pl. 11#9 is probably *Thorybes pylades*. *Magnolia*, *Zanthoxylum* & *Cercis* are errors for *Papilio troilus*, *Magnolia* an error for *P. palamedes*. Georgia has *Celastrina neglecta* all over, *C. ladon* in most of state except the S, *C. idella* on E edge, & *C. neglectamajor* & *C. nigra* in the north. *Sedum* is an error for *Junonia coenia*. *Lysimachia* is an error for *Chlosyne gorgone*.

A FIELD GUIDE TO THE BUTTERFLIES OF TEXAS. Raymond. W. Neck.

1996. Gulf Publishing Co., Houston, Texas. 323 p.

This book is the only one to cover Texas, which has the largest butterfly fauna in the U.S. It has a brief description, some hostplants, brief flight period & brief life history info, general range, & brief comments for each species. There are nearly 200 color photos of living butterflies in nature, which are nice, but many of them are misidentified, and less than half the Texas species are figured; b/w figs. could have been added in text for the others with little expense. It has more than 50 nice color photos of immatures. There is a crude little outline map of Texas next to each species, with the range shaded in black= native or lined= stray or stippled= rare strays, but these maps are not accurate, compared to the county dot maps of Ray Stanford. Some records mentioned in text are not on maps. Another 20+ species are known from Tex. since the book; evidently most of the NE Mex. fauna will sooner or later be found as strays in Tex. (another 50 sp. at least). Overall, the text is too brief. I checked all the hosts, but have not studied much of the text.

Plates: Pl. 21#39 is *Phoebis agarithe* (straight unf line). Pl. 24#56 is *Lycaena dione*. Pl. 26#79 is *Calycopis cecrops*. Pl. 27#87 is *Callophrys irus hadros*. Pl. 28#98 is *Ministrymon clytie*. Pl. 28#100 is *Strymon istapa*. Pl. 31#123 looks like *C. nemesis*. Pl. 33#130 is *A. virgulti duryi*. Pl. 37#161 is *Phyciodes tharos* female. Pl. 43#193 is *Adelpha basiloides*. Pl. 48#230 is *Lethe anhedon* female. Pl. 57#353 is *Hesperia* (*leonardus* *pawnee*?, maybe *attalus*) male. Pl. 59#372 is *Hesperia viridis* male. Pl. 63#428 is *Panoquina panoquinoides*. Pl. 64#441 is *Agathymus mariae*.

Detailed comments: At the top of each species are letters, E=egg, L=larva, C=chrysalis, A=adult, signifying the presence of color photos of these. P. 34, *ilioneus* is a homonym, and *texana* a syn. P. 37 spelled *Viscaceae*. *Pieris napi* does not occur in Tex. *Euchloe hyantis lotta*—often treated as a sp.—is in Tex. P. 46 spelled *clorinde*. *Amelanchier* is dubious for *Satyrium titus*. *Calycopis isobea* has narrow unf red band, & much red on unh thecla-spot, compared to *C. cecrops*. *Brephidium isophthalma* in SE Tex. are not strays, and intermediates between *exilis* & *isophthalma* were reported from Nueces Co. & La. *Euphilotes rita* occurs S of Alpine in Aug.-Sept. (not in Big Bend Park, not June) & eats *Eriog. wrightii*. E Tex. has *Celastrina neglecta*, W Tex. evidently *C. neglecta cinerea*. *Plebejus alupini texanus* occurs in Tex. *Calephelis r. rawsoni* occurs in C Tex., and *C. r. freemani* occurs N of Alpine & in Big Bend Park. (& *C. r. arizonensis* in Ariz.). *Apodemia virgulti duryi* (not *mormo*) occurs S of Alpine on April 18 assoc. *Eriog. wrightii*. It's *Apodemia nais chisosensis*. *Libythea carinenta larvata* has the white upf costal spot separated from the larger white spot. P. 116 *Turnera ulmifolia*. *Poladryas minuta* flies Apr.-Sep. in C & W Tex. (records W Guadeloupe Park, & W of Alpine). *Chlosyne chinatiensis* flies in W Tex. near N.M. and at Terlingua. *C. d. definita* occurs in C & S Tex., *C. d. anastasia* in W Tex.; spelled *Stenandrium*. *Siphonoglossa* is eaten only in lab for *C. rosita*, *Beloperone* in lab for *Phyc. texana*. *Phyc. texana seminole* & *P. frisia tulcis*. *Euphydryas chalcedona* is an error. P. 133 *Artemisia*. *Junonia evarete nigrosuffusa* is a common resident of S Tex., whereas the Black Mangrove Buckeye *P. genoveva* is a stray there. Texas has *Adelpha eulalia*. *Dynamine serina dyonis* is known from near Dallas; *D. tithia* is surely an error. *Hamadryas guatemalena*. *H. iphthime* was recorded from Burnet Co. (not Rio Grande V.), & maybe was mislabeled. *Myriocarpa* was perhaps an error for *Smyrna bomfordia*?; & *Arctocarpus* & *Chlorophora* may be errors for *Marpesia chiron*? P. 155 *monanthogynus*. *Texana* is a syn. of *A. clyton louisia*. Avery Freeman wasn't sure about seeing the *Morpho*. *Neonympha areolata* records are really *N. helicta*?, and *Sorghastrum* was probably just used in lab. *Greta polissena* is surely an error. *Urbanus proteus* larva is the

Bean Leaf Roller. *Stenotaphrum* is a lab host only for *Urbanus procne*. *Gilberti* is a syn. of *Astraptus alector hopfferi*. *Astraptus anaphus* hosts are *Pueraria lobata*, & “creeping wild bean” in S.A. *Lespedeza texana* was eaten only in lab for *Achalarus lyciades*. *Thorybes pylades albosuffusa* is in W Tex. P. 287 *Cologania*. *Dyscophellus euribates* is an error. P. 202 *Acacia angustissima*. All *Pellicia angra* records refer to *P. arina*. P. 204 bottom delete “Texas subspecies *georgina*” which belongs on p. 213. The *Staphylus azteca* was a misid. *S. ceos* female. The correct name for *Gorgythion begga*=*beggina* *vox eludes me*. J. Burns mapped *E. juvenalis juvenalis* from Davis Mts. Texas. *Quercus emoryi* & *Q. grisea* were assoc. records only by J. Burns for *E. tristis*. *Pyrgus oileus* has zigzags on unh submargin. *Hesperopsis alpheus* eats *Atriplex canescens*. *Piruna haferniki* has unh base grayish. *Nastra lherminier* & *N. neamathla* have wider ranges, & *julia* is a ssp. of *neamathla*. The hosts of *Lerema liris* are from R. Kendall, maybe just in lab. *Poa* is an error for *Ancyloxypha numitor*. The *Adopaeoides prittwitzii* record was E of Davis Mts. *Hesperia viridis* unh is not green. *Panicum* is assoc. only for *Hesperia attalus*. *Polites rhesus* flies in May. *Polites subreticulata* is not in U.S. (only *Sinaloa* to Panama, J. Burns 1994 J. Lep. Soc. 48:39). *Stenotaphrum* is lab only for *Polites vibex*; ssp. *vibex* occurs in E Tex., *praeceps* in S Tex; *brettoides* is not in Tex. & is evidently a syn. of *praeceps*. *Wallengrenia egeremet* spelling. P. 253 logan. P. 255 *Dactylis*. *Alabama* is just a form of *Euphyes dion*. SE coastal Tex. also has *E. dion bayensis*. P. 259 ssp. *metacommet* & *turneri* are synonyms. *Atrytonopsis pittacus* flies in L Mar.-July. *A. ovinia edwardsi* eats *Bouteloua curtipendula* in Ariz. *Panicum bulbosum* was only suspected in Ariz. for *Amb. nysa*, *Panicum obtusum* was only probable in Ariz. for *A. eos*, & *Poa* was only a lab food for *Amblyscirtes vialis*. Avery Freeman recorded both *Amb. celia* & *belli* near Dallas, & I found a *celia* in Davis Mts.; the blotch ranges of both are too small. The blotch for *Amb. fimbriata* is misplaced; it surely does not occur in Chinati Mts., and is absent on Ray Stanford’s maps, so does not occur in Texas or anywhere near Tex. *Dysaules* is still a ssp. or form of *Lerodea arabus*. *Agathymus neumoegeni* eats *Agave parryi*=*neomexicana*; *A. lechuguilla* is an error; *chisosensis* is a ssp. *Ag. mariae* eats *Ag. lechuguilla* (not *lecheguilla* on p. 278-9), & *gilberti* is a syn. *Agathymus remingtoni* has ssp. or synonyms *estelleae* & *valverdiensis*. *Coloradensis* is a ssp. of *Meg. yuccae*; *maculosa* a ssp. of *Stallingsia smithi*.

BUTTERFLIES OF HOUSTON & SOUTHEAST TEXAS. John & Gloria Tveten.

1996. University of Texas Press, Austin, Texas. 293 p.

An outstanding book. There are nice color photos from nature for adults of most species, and many for immatures too. Only one misidentified photo! Fine glossy pages. The text is more complete than in most books, with lengthy discussions of interesting stuff and local information; no skimpy formulaic word-processing here. They have done a lot of work on the hostplants and biology of Texas butterflies (they should write *Butterflies of Texas*). There are good insights on the hosts of *Feniseca*, new life history info & larva photos on *Phaeostrymon*, conclusive analysis of *Calpodes* hosts, etc. They clearly seem to demonstrate that *isobea* is a ssp. of *Calycopis cecrops* (they intergrade in Houston & Baytown, and intermediates are illustrated on p. 79 & 81), and cite studies by Mike Rickard that show that *julia* is a ssp. of *Nastra neamathla*. They also show *isophthalma*-like adults from a colony of *Brephidium exilis* near Rockport in Aransas County, suggesting that these are ssp. too.

Detailed comments: *Battus philenor* is illustrated with both black & red larvae. P. 28 has the thoroughly-disproved balanced polymorphism theory again. The discussion of *Pontia protodice* scent glands & uv reflection is incomplete; females have repellent scent glands, and butterfly pheromones are not well studied. P. 82, *gryneus* was in *Mitoura*, then was placed in *Callophrys*, but now the splitters are mounting more zombie attacks (*Loranthomitoura* etc. hypersplitting), but we must resist & keep it in *Callophrys*. Interesting that *Callophrys henrici* larvae were red on seed pods, green on leaves. Very odd that Texas *Strymon melinus* larvae are only green, as they are very polymorphic in Colo. *S. melinus* raits on bushtops even in spring, after 1 p.m. to dusk, so surely they mate then. *Echinargus isola* life history & many hosts were in *Papilio* (N.S.) #6. E. Tex. has *Celastrina neglecta*; the photos seem to be *Cel. lucia* f. *marginata* from Canada. P. 98 Colombia. *Libytheinae* & *Heliconiini* are wrongly raised to family rank, & *Dione vanillae* is wrongly stated to be not closely related to *Argynnis* (*Speyeria*), when actually it is closely related to *Euptoieta*. They write that *Euptoieta* hibernates as adults in Houston. Evidently *Phyciodes t. texana* occurs in Houston. Note that all 1m2f *Phyc. tharos* figured have orange antenna nudum instead of black, a high frequency as in La. Only *Vanessa atalanta* adults hibernate. They note *Vanessa cardui* in midwinter in Houston! Cross out the sentence on intraspecific hybrids & isolating mechanisms on p. 163. P. 171 parag. 5 describes raiting behavior of male *Asterocampa celtis*, not challenges or salt-seeking. SE Tex. has *Neonympha areolata* rather than *N. helicta*, based on the photo. *Megisto cymela* still needs a lot more work, and J. Calhoun found that *viola* intergrades with *cymela* in the Fla. panhandle. P. 206 *Apyrrothrix araxes* is native in Ariz. *Epargyreus clarus* usually ovip. on the host. *Thorybes pylades* courts & mates all day. Intermediates of *Erynnis zarucco* *X* *funeralis* are reported in La. & Fla., which may just represent whiter-fringed *zarucco*, as Houston has only *E. funeralis*. Scott wrote that *Nastra julia* ranges S to Costa Rica; W-Tex.-Ariz.-Calif. records of *Nastra neamathla* are errors for *julia*; this book shows that *julia* is a ssp. of *neamathla*. *Wallengrenia egeremet* occurs at Lake Houston just N of Houston (p. 242) so should have been included in book. P. 244 the antenna base is pale in *Euphyes vestris* too. *Atalopedes* has become common in S Minn. & Denver, so I suspect it now overwinters farther N than we thought. *Poanes yehl* also occurs at Lake Houston (p. 248) & should also have been included. *Euphyes dion bayensis* occurs in extreme SE Tex. Photo of *Amb. vialis* is actually *A.*

alternata female. P. 253, wax is produced to protect the pupa from physical damage in the nest, not to protect a larva outside the nest from predators. Good info on choice of Cannas by Calpodes; the translucent body shows the tracheae—not circulation—and 2 testes (p. 252 larva also shows 2 testes). Index: cauliflower is *B. oleracea botrytis*. Frog-fruit *P. incis*a 141. Nasturtium is *Tropaeolum*. Tube-tongue is *Siphonoglossa*.

BUTTERFLIES OF SOUTHEASTERN ARIZONA. Richard A. Bailowitz & James P. Brock. 1991. Sonoran Arthropod Studies Inc., Tucson, Arizona. 353 p.

This is a good book, which covers the SE 1/6 of Ariz. Much is known but unpublished about Arizona butterflies, which will not be published during the lifetime of the major discoverer, so this publication is welcome. It has b/w photos for each species, along with distributions within the area (mostly the mountain ranges occupied) and flight periods, together with habitat. Life history information is limited to hostplants; it reports some good hostplants for butterflies in the region. Unfortunately, the book often uses the wrong terms for mate-locating behavior, thus for a dozen species it says they “patrol hilltops” when their main behavior is perching to await females on hilltops (if you walk onto a hilltop and disturb him, he will fly around, but will land when you stop disturbing him). And often the book uses the word perch for just sitting on a substrate. The new terminology will help here: raiting behavior consists of males resting (landing) to await females at certain characteristic sites to await females, flaiting behavior consists of males flying around a small area where they await females (example *Papilio eurymedon* flying slowly about tiny clearings among trees on ridgetops/hilltops to await females), and fleeking behavior consists of males flying about a larger area (canyons for *Anthocharis sara*, or the canopy of a lot of host trees for *Hypaurotis*, etc.) to seek females. Various species discussed and labeled as “hypothetical” are in general just errors. Of course new species since have been found in S Ariz. (*Grais stigmaticus*, *Strymon cestri*, *Euchloe [hyantis] guaymasensis*, *Chlosyne marina*, *Phyciodes pallascens*, etc.).

Detailed comments: P. 5 population pressures don't cause migration, it's genetic. *Epargyreus exadeus* records are probably *E. windi* according to A. Warren. P. 22 maybe Osler's May specimen was an error; in Colo. he mislabeled May *Oeneis alberta* as *Sept. Zestusa dorus* rait on hilltops. I haven't had any trouble sorting *Achalarus albociliatus*, *toxeus*, & *jalapus* by their distinctive genitalia. Ach. “jalapus” in Ariz. are really *A. toxeus* (A. Warren, pers. comm.). There must be several gen. of *Thorybes pylades*. *Dobra* is a syn. of *Thorybes mexicana*. *Codatractus valeriana*=*mysis* is the correct name (O. Mielke & A. Warren, Rev. *Brasileira Zool.* 21:307-8), contrary to J. Burns. *Erynnis brizo* & *E. propertius meridianus* & *E. persius* males rait on hilltops. J. Burns reported *Quescus hypoleucoides*, not *Q. grisea*, for *E. juvenalis clitus*. I think there's a possibility that *Pyrgus communis* & *P. albescens* are just dominant/recessive genitalic polymorphs, because of peculiar suggestive variation of mtDNA in Calif. and the costal fold and valva polymorphism in Mex. and the recent replacement of *communis* by *albescens* in Fla., but rearing studies will be necessary to study this. I found *Pyrgus philetas* mostly at wet habitats. *Systasia pulverulenta* was fig. from Baboquivari Mts. in J. Comstock's Butt. Calif. *Hesperopsis alpheus* likes alfalfa flowers in Colo. Cingo should be called *Piruna aea mexicana*. *Nastra julia* occurs from Tex. to Calif., & *neamathla* from Tex. to Fla., and this record is *julia* if valid (Butt. Houston wrote they are conspecific). *Adopaeoides* is common E Aug.; host spelled *Paspalum distichum*. Book probably means *Yvretta carus* rait on hilltops. *Hesperia uncas* & *H. comma susanae* & *H. pahaska* rait on hilltops, & *H. viridis* rait in gulches. I found *H. woodgatei* raiting on Chiricahua Mts. hilltops. *Paratrytone* “*Ochlodes*” *snowi* actually feeds only on *Muhlenbergia montana* (*Blepharoneuron* was my error). *Atrytonopsis hianna deva* (only differs by variable stigma) rait in gulches. *Edwardsi* is a ssp. of *Atrytonopsis ovinia*. I later named *Amb. aenus megamacula* for the region. *A. oslari* males rait in gulches all day. *Amb. tolteca prenda* is in Ariz. *Amb. nysa* rait in gulches incl. low points on roads, all day. P. 107 *Thespis*. *Megathymus yuccae* males do sip mud. *Meg. ursus* females are slower so can sometimes be caught, whereas faster males are nearly impossible to catch. *Battus philenor* larvae are usually dull red in the region. *Papilio polyxenes coloro* surely rait & flait on hilltops; was the coloro record the yellow form? P. 121 top photo is the yellow-form *P. polyxenes f. pseudoamericus* male from Whetstone Mts., and bottom looks like *P. p. asterius* female. *P. crespontes* & *P. thoas* are easy to distinguish using wing pattern and dorsal view of male abdomen tip (see my 1986 book). *P. glaucus rutulus* occurs in Ariz.; *arizonensis* is a syn.; *rutulus* has a partial gen. until Sept. P. 134, the text means “rest” with wings fluttering. P. 141, 146 *Thysanocarpus*. All the *Pontia protodice* photos are spring f. It's *Ascia josephina howarthi*. The *Anth. sara inghami* photos look more like ssp. *thoasa*; mine from Sabino Can. have less black; males mostly fleek in gulches. P. 148 *eriphyle* is a slightly-orangish ssp. limited to BC. *Eurema nise* lacks the unh basal dot of *E. lisa*. *Nathalis iole* always flies low; and dark soil would reflect more heat. *Callophrys spinetorum* has one gen. in Colo. *Satyrrium favonius ilavia* is in Ariz. (*S. polingi* is a separate sp. eastward.) *Strymon melinus* rait on hilltop bush/tree tops. I found *Erora laeta quaderna* in the Patagonia Mts. Ariz. probably has *Celastrina neglecta cinerea*. *Euphilotes bernardino martini*. P. 204 the photos are *Apod. virgulti mejicanus* from (p. 310) Empire Mts.; ssp. *mormo* lacks ups red, and occurs at Portal in Chiricahua Mts. (Gordon Pratt), & I found it at Molino Basin. *Apodemia palmeri* photos are female. *Apodemia phyciodoides* was surely mislabeled from Ariz., as the original collectors were often in Sonora. (*Argynnis*) *Speyeria hesperis nausicaa* is in Ariz. Segregate #1 is *Poladryas minuta arachne=gilensis*; #2 is *P. m. near-nympha*; males rait on hilltops in morning, and may fleek on flowers in afternoon. *Chlosyne leanira fulvia* rait & flait on hilltops. *Chl. acastus sabina* mostly rait in washes. P. 230 *Anisacanthus*. *Phyciodes frisia tulcis*. *P. pallascens* is known from St. David (Sept. 12 1958 Ray Stanford 1f), and other

“campestris” records from SE Ariz. may be *pallescens*. *P. picta canace* & *P. mylitta arizonensis* are in Ariz. *P. mylitta* should be stated to *rait* & *fleek* in washes. *P. 239* surely *Urtica dioica gracilis*. *Nymphalis antiopa* *raits* in valley bottoms & back yard clearings. *Nymphalis milberti* *raits* all day on talus/clearings just off hilltops, & migrates to higher altitude in summer, explaining its presence near summits. *Junonia evarete nigrosuffusa*. *Limenitis weidemeyerii* often visits flowers. *Adelpha eulalia* is a distinct sp. from *A. bredowii* as they are sympatric in S Mex. *P. 267* *Croton texensis*. *Asterocampa celtis montis* occurs in SE Ariz., & compared to *antonia* it has orange up and pale median up band and the spot in fw cell CuA₁ mostly solid black. *Texana* is a syn. of *A. clyton louisa*. *Cyllopsis pyracmon henshawi* has 1st gen. (*henshawi*) & 2nd gen. (*nabokovi*); Douglas Mullins reared one from the other. *Allyni* is a ssp. of *Paramacera xicaque*. The numbers on Appendix B are species nos. not page nos.

BUTTERFLIES OF ANZA-BORREGO DESERT STATE PARK & ENVIRONS.

Lynn & Gene Monroe. 2004. Published by authors.

This is a nice book. The best part is the numerous nice photos of adults and some immatures. The book includes lots of hostplant information from the literature (which are unfortunately not distinguished from local hostplants that the authors or other lepidopterists might have discovered), and a lot of indexes to such information, which seems to be a waste of space to me; I would have preferred that space be used for more local info and local lore.

Looking over the book I found few goofs: *P. 19* lower right photo looks like *P. p. coloro* not *zelicaon*. *P. 26* left middle photo is *P. protodice f. vernalis*, note weak apical markings (and *sisymbrii* has unf cell-end bar broken in middle). *P. 28* upper right looks like *E. ausonides* but is *hyantis* as the book states, because *ausonides* does not occur near there. *P. 41* lower right *L. helloides* is male, not female. *P. 53* it seems that *Everes* is just part of the genus *Cupido*, so *Cupido* is the correct name. *P. 59* the *acmon* up orange band is pink, without a dark inner border except in spring. *P. 83* should be *Prudic et al. 2001*. *P. 102*, *Cynodon dactylon* is a very dubious host of *O. sylvanoides*, as that bug eats only hay grasses mostly on N-facing slopes.

THE BUTTERFLIES OF SOUTHERN CALIFORNIA. Thomas C. Emmel & John F. Emmel.

1973. Natural History Museum Los Angeles Co., Science Series 26. Los Angeles, Calif. 148 p.

This is a good book, and has few errors. It has good color plates of all species, and has lots of b/w drawings (most by Charles Dammers) and some b/w photos of immatures. It has a lot of local lore. *Argynnis* (*Speyeria*) *adiaste atossa* went extinct in 1959 when I first started collecting butterflies, the later-named *Philotes sonorensis extinctis* has been long-extinct, and since then several others such as *Euph. editha* have become nearly extinct in S Calif.; weeds and global warming are diminishing the already sparse (except for host races of *Eriogonum* feeders) fauna. The book is out of date, as much more has been discovered: *vallismortis* has been transferred to *Chlosyne acastus*, *Euphilotes* & *Apodemia* have been expanded greatly and plunged into chaos, *Cercyonis sthenele* near-*sthenele* has been found, many genera have changed, and numerous new ssp. have been named (many mentioned above).

Detailed comments: The text notes that the eyespots on larvae of *Papilio glaucus rutulus*, *P. eurymedon*, & *P. multicaudata* are quite variable in pattern, which is interesting as that trait has been used to distinguish *rutulus* from *P. g. glaucus* which has just the big eyespot. The pupa drawing of *Pontia beckeri* is *Colias* based on wing pattern and lack of hump on A2-3 & lack of bird-dropping appearance. *P. 19* top, *Colias alexandra altamont* & *C. edwardsii* ssp. have two gen. *Colias philodice* hibernate as half-grown larvae. Adults & pupae of *Zerene cesonia* are claimed to hibernate. The *Nathalis* sex-brand is red. *Catalina* is claimed to be a distinct ssp. but the photos look the same as *cethura*. *Gunderi* is a syn. of *thoosa* or is very close. *Andrewsi* is said to have larvae that look like *E. hyantis*, yet adults resemble *ausonides*, & *p. 25* says an *E. hyantis lotta* pop. flies near it near E end of San Bernardino Mts., which provokes questions (and *hyantis* and *lotta* are reported to intergrade on Kern Plateau, why hasn't that been published?). Ken Davenport found *Cercyonis pegala* in S Calif. in Kern Co. *Viridicornis* is now considered a syn. of *S. hydaspes hydaspes*. *E. chalcidona quino* was later named *E. c. hennei*, and *E. editha wrighti* is now *E. editha quino*, which is now nearly extinct. *Microtia* (*Dymasia*) *dymas chara* flies most of the year in Ariz. *P. tharos tharos* occurs in S Calif. (or occurred), while *campestris* is now known as *P. pulchella pulchella*. *Phyc. phaon jalapeno* is in Calif. Fig. 39 is supposed to be from Edwards 1884 but that's not in references, Edwards' Butt. N.A. is 1884 but the fig. isn't in my microfiche copy of BNA, and the fig. didn't come from Can. Ent. 13:226. *P. 49* *Apodemia virgulti deserti* does occur in the W Mojave Desert. *A. dialeuca* (host *E. wrightii*, not *kennedyi*), *A. mormo*, & *A. cythera* are now considered separate species. Still no study of whether *avalona* is just a ssp. of *S. melinus*; they reportedly hybridize on Santa Catalina I. where *melinus* has been introduced. *Ines* is the fall f. of *Ministrymon leda*. *P. 58* notes that *Callophrys gryneus juniperaria* & *C. g. loki* intergrade at NE end of San Bernardino Mts.; *nelsoni* is a ssp. of *C. gryneus* too. *P. 68* the greenish pops from Tehachapi & Tejon Ranges are *Pleb. chlorina chlorina*. (*P. chlorina monticola* & *P. acmon* also occur in S Calif. of course). *Euphilotes bernardino* eats *Eriog. fasciculatum* not *nudum* in El Paso Mts. *Glaucopsyche piasus sagittigera* also occurs in inland mts., rather close enough to ssp. *piasus* that one wonders whether it might be a separate species *G. daunia sagittigera*? *Agathymus stephensi* egg shape is wrong (it is flatter, see Bull. So. Cal. Acad. Sci. 33:84). *Hesperia pahaska martini* was later found to be smaller, so is a

syn. *Hylephila phyleus phyleus* occurs in most of S-Calif., but the later-named *H. p. muertovalle* occurs at Needles, and in diluted form ranges W-ward in Mojave Desert. Fig. 70, the lower left larva & pupa are actually *Hesperopsis libya*, while lower right larva & pupa are *Pholisora catullus*. *Hesperopsis libya* in Inyo Co. are near-*libya* (*lena* is limited to Montana-Colo.). *H. alpheus graciellae* has multiple gen. *Heliopetes domicella* was just a stray at Parker Dam, based on one specimen that I caught (somehow one wing disappeared during capture). *Sidalcea* is an error for *Pyrgus ruralis*, which was later named *P. ruralis lagunae*, now scarce. *E. afranius* as well as other *Erynnis* hibernate as mature larvae. Many more species have been recorded from So. Calif. since the book, including *Dione moneta* from Fullerton Calif. July 20 1929 1 male; *Phyc. texana* from Anza-Borrego SP (Michael Young & Dave Wagner). W. Field's Butt. Kansas p. 108 recorded *Anaea aidea* from Sierra Madre Cal., but maybe he meant Mex. Bull. So. Cal. Acad. Sci. 30:93 mentioned *Urbanus simplicius* from Fertilla, evidently referring to the *U. procne* mentioned on p. 95. P. 99, Evans W. H. 1952 & Evans W. H. 1955 are two different people, the latter Brigadier W. H. Evans of British Museum Nat. History, the former an LA resident. Pl. 4 fig. 36 & 38 are both from Mint Can., so 38 must be *Chl. leanira wrightii* also, the *cerrita* phenotype.

BUTTERFLIES OF CALIFORNIA John A. Comstock. 1927

Published by author. 334 p. & 63 plates.

This book is long out of print, but is of historical importance, and the plates are very nice. So I'll just correct bad misidentifications on the plates, and ignore the endless obvious name changes of species & genera that the reader can easily update. The leather cover has a beautiful embossed butterfly. The book has a lot of drawings of immatures by Charles Dammers and others. The text is interesting reading with lots of lore.

Pl. 1#6 is *Papilio zelicaon* (all LACM specimens from area are *zelicaon*, says K. Davenport). Pl. 5#7 has typical *P. phoebus smintheus sphaeris*. Pl. 6#4&7-8 *P. p. smintheus*. Pl. 8#16 *Pieris marginalis reicheli*. Pl. 9#7-8 *P. marginalis*; 12 *P. marginalis venosa*. Pl. 10#4-5 *Euchloe hyantis lotta*. Pl. 11#1-3 *A. cethura cethura*; 13 *A. julia flora*; 14 *A. julia columbia*; 15 *A. julia near-stella*. Pl. 14#7-12 *Colias philodice*. Pl. 15#11-13 *Colias columbiensis pseudocolumbiensis*. Pl. 16#14 *Eurema daira palmira*; 16 *E. daira daira*. Pl. 18#7-8 *Coenonympha tullia ampelos*; 9-10 *C. tullia swadneri*; 19-21 *C. pegala nephele*. Pl. 19#2-6 *C. pegala nephele*; 7-10&12 *C. pegala near-stephensi*. Pl. 20#15 *C. oetus charon*. Pl. 24#1-3 *Argynnis (Speyeria) egleis utahensis*; 4&6 *S. zerene behrensii*; 5 *S. zerene hippolyta*; 7 *S. hesperis chitone*; 10 *S. zerene conchyliatus*. Pl. 25#3-5 *S. hydaspe near-hydaspe*. Pl. 26#1 *S. zerene zerene*; 2 *S. hesperis cottlei=dodgei*; 4-6 *S. hesperis irene*; 7 *S. zerene zerene*; 8 *S. zerene bremnerii*, mislabeled from Calif. Pl. 27#4-6 *S. zerene gunderi*; 7-8 *S. coronis snyderi*. Pl. 29#4-6 *S. coronis coronis*; 7-8 *S. callippe liliana*. Pl. 30#1-3 *S. callippe juba*; 4-5 *S. callippe rupestris*; 6-9 *S. callippe macaria*; 10-12 *S. egleis oweni*. Pl. 31#1-3 *S. zerene malcolmi*; 4-6 *S. egleis egleis*; 7-9 *S. mormonia mormonia*; 10-12 *S. egleis tehachapina*. Pl. 33#1 *Euph. chalcedona chalcedona*; 4-5&9 *E. "colon" wallacensis*. Pl. 34#11&13&15&17 *E. editha nubigena*. Pl. 35#5 *Euph. chalcedona sierra*. Pl. 36#13-#19 *Chlosyne palla palla*. Pl. 37#1 *Chl. palla altasierra*; 3 *Chl. palla dark-female ssp.*; 19-21 *Chl. leanira obsoleta*. Pl. 38#1 *Chl. leanira wrightii*. Pl. 39#1-3 *Phyc. phaon jalapeno*; 4-6 *Phyc. pulchella near-pulchella*; 16 looks like a mislabeled *Phyc. pallida barnesi*. Pl. 40#2-8&10 *Polygonia satyrus satyrus*. Pl. 43#4&7 *Vanessa atalantaXcarye annabella*; 11 *Junonia evarete nigrosuffusa*. Pl. 44#11 *Junonia evarete nigrosuffusa*. Pl. 45#2 *Lim. weidemeyerii latifascia*; 5-6 *L. lorquini ilgae*. Pl. 46#3 *D. gilippus thersippus*. Pl. 47#1-3 *L. carinenta larvata*; 5 *Apodemia virgulti deserti*; 13-14 *A. virgulti mejicanus*; 15&18-19 *Calephelis nemesis*; 16-17 *Calephelis wrightii*. Pl. 48#7-9 *Strymon istapa*. Pl. 49#6-8 *Satyrium sylvinus desertorum*; 19-23 *Sat. saepium*. Pl. 50#4 *Callophrys gryneus nelsoni*; 16&20 *Callophrys [sheridani] viridis* ["*dumetorum*", a name soon to be banished]; 17-19&21 *Call. affinis perplexa*; 25 *Call. [sheridani] viridis?*; 26&28 *Satyrium fuliginosa "tildeni"*; 27 *Satyrium (fuliginosa) semiluna "maculadistinctum"*. Pl. 52#1-2 *Lycaena xanthoides editha*; 3 *L. xanthoides edithaXxanthoides "pseudonexa"* (unh spots smaller than regular *editha*)(Dunsmuir is a famous site for these intermediates, see J. Scott 1981 J. Res. Lepid. 18:50-59). Pl. 53#10-12 *Hemiargus hanno gyas*; 19 *Cupido amyntula amyntula*; 21-22 *Plebejus atrapraetextus fridayi*; 23-25 *Pleb. melissa paradoxa*; 29-31 *Pleb. podarce*. Pl. 54#1-2&4-5 *Plebejus saepiolus aehaja*; 3 *Pleb. saepiolus gertschi*; 16 *Pleb. saepiolus rufescens*; 17-19 *Pleb. icarioides icarioides*. Pl. 55#7-9 *Plebejus alupini alupini*; 10-12 *Plebejus chlorina monticola*; 13&15 *Pleb. chlorina chlorina*; 14&16 *Pleb. chlorina chlorinaXpleb. neurona*; 27-28&33 *Euphilotes glaucus oregonensis*; 29-32 *Euph. bernardino*. Pl. 56#7-11&14-15 *Philotes sonorensis extinctis*; 20-23 *Glaucopsyche piasus sagittigera*; 24 *Glaucopsyche lygdamus incognitus*; 25 *Glauco. lygdamus columbia*; 30-31 *G. lygdamus xerces*. Pl. 57#1-5 *G. lygdamus xerces polymorphs*. Pl. 58#1-2 *Thorybes pylades*; 19-21 *Hesperopsis alpheus graciellae*. Pl. 59#1 *Erynnis brizo lacustra*; 2-3 surely *Erynnis persius*; 4 *Erynnis afranius*; 5-6 *E. pacuvius pernigra*; 7-8 *E. pacuvius callidus*; 9&11 *Erynnis propertius*. Pl. 60#17-18 *Hesperia comma near-hulbirti*; 19 *H. comma leussleri*; 20 *Hesperia pahaska*; 21 *Hesperia juba*; 22 *H. comma idaho*. Pl. 61#1-3 *Hesperia lindseyi*; 10-12&14-15&19 *Ochlodes agricola nemorum*; 13 *Ochlodes yuma?*; 29 *Polites sabuleti sabuleti* variant like *chusca*. Pl. 62#18-21 *Panoquina panoquinoides errans*.

BUTTERFLIES OF BAJA CALIFORNIA. John W. Brown, H. G. Real, David K. Faulkner. 1992. Lepidoptera Research Foundation, Inc., Beverly Hills Calif. 145 p.

This is a good book, which brought together lots of data on this W Mexico peninsula for the first time. (Since then, there has been little study of the fauna, as collectors are arrested and their bugs seized if they do not have the extortion-priced nearly-\$1000.00 Mexican butterfly-collecting permit.) There are nice dot maps of each species, and 8 nice color plates show all species. N Baja is dry in May-Sept. like Calif., most of Baja is dry all year, and the Cape is moist in Aug.-Oct. Evidently the 17-25 Pleistocene ice advances, along with drought, have kept most of the central mainland Mexican fauna out of Baja Calif.

There aren't many mistakes in this book. P. 18, there are not as many endemics as the book writes, as *festata* is a syn. of *Hypostrymon critola*, & *comstocki*, *dawsoni*, *harbisoni*, *cedrosensis*, *howarthi*, *murphyi*, & *poodiae* are ssp. of other taxa, as noted below. *Codatractus* (not *Thorybes*) *valeriana*. P. 30 line 13 should be Pyle (1981). *Staphylus ceos* isn't especially shade-loving. *Erynnis afranius* *raits* (perches to await females) in gulches/swales in Colo., so I doubt that it mate-locates on hilltops in Baja. I wonder if *Pyrgus communis* vars. could occur in the upper Sierra San Pedro Martir? *Abutilon* & *Sida* are errors for *P. oileus* (belong to *P. communis*). *Hesperopsis alpheus* *gracielae*. Real *neamathla* does not occur W of Texas; Baja has *Nastra neamathla julia* (see Butt. Houston). *Hylephila phyleus* *phyleus* occurs in Baja. *Bromus* is a *Hesperia juba* host in Colo. *Vulpia* is dubious for *Hesperia comma leussleri*. The *Hesperia pahaska* in upper Sierra San Pedro Martir look a little odd as if they might have introgressed with *H. columbia*. (Note that *Polites sabuleti* *norae* of the Sonora coast on *Distichlis spicata* is the same as *margaretae* but male lacks stigma & sterigma has a few fewer folds.) *Poanes melane melane*. *Amblyscirtes tolteca* *prenda* is in Baja as the figs. are *prenda*. *Dysaules* is a syn. or form of *Lerodea arabus*. *Panoquina panoquinoides* *errans*. *Comstocki*, *dawsoni*, & the later-named *harbisoni* are ssp. of *Agathymus stephensi*, & *coloradensis* a ssp. of *Megathymus yuccae*. *Papilio glaucus rutulus*. It's *Ascia josephina howarthi*. *Lycaena arota* host is *Ribes quercetorum*. *Habrodais grunus poodiae* will be a distinct species when poodles sprout wings & fly. P. 119 says 1 *Hypaurotis crysalus* specimen is known from Sierra San Pedro Martir. Baja has *Callophrys affinis perplexa*, & *dumetorum* should not be used as it surely will be suppressed by ICZN. *Nelsoni* & *loki* & *cedrosensis* are ssp. of *Callophrys gryneus*; *cedrosensis* is related to *C. g. thornei*. *Festata* is a syn. of *Hypostrymon critola*. *Strymon istapa* not *columella*. *Hemiargus hanno gyas*, *Echinargus isola*, *Cupido amyntula*, & *Celastrina neglecta* echo occur in Baja. 126b is *Euphilotes mojave*; the host is probably *Eriog. davidsonii*, because *E. pusillum* does not occur in Baja Calif. The blue of *Glaucopsyche lygdamus maritima* was no doubt just darker due to fading; maybe mislabeled? *Plebejus chlorina monticola*. *Apodemia virgulti maxima* was recorded from Jan. 2, Mar. 23-25, May 26, Sept. 29-Oct. 1, Nov. 9-13, Dec. 28 by Opler & Powell. *Deserti* also seems to be a ssp. of *Apod. virgulti*, while *Apod. dialeuca* is evidently a different sp. *Murphyi* is an obvious ssp. of *Apod. palmerii*. *Libythea carinenta larvata* is in Baja. C Baja *Chlosyne leanira* were later named *austrima* Austin & Smith 1998. *Phyc. phaon jalapeno* is in Baja. P. 103 the O.D. named it *Euphydryas chalcedona hennei* (so no parentheses). P. 158½ says *Aglais milberti* was seen in Sierra de Juarez. The common *Junonia* from mangrove tidal flats at Puerto Lopez Matteos at Magdalena Bay has host and narrow upf band like true Fla. *J. genoveva*, but the band is a little whiter and the upf is blacker with narrower orange submarginal band, like fig. 18, thus is evidently *J. genoveva* new ssp. (the sp. name has been switched back to *genoveva* from *evarete*). Fig. 19 is *J. evarete nigrosuffusa* from mainland Sonora. *Mestra dorcas amymone*. P. 124 Gorelick. Most of the butterflies on the plates should have been illustrated on uns, not ups (& most butterflies throughout the world should be mounted to show uns, because the identification traits are usually on uns, not ups). Numbers were trimmed off the bottom of several plates. Pl. 1#34 is *Nastra neamathla julia*.

THE BUTTERFLIES OF WEST VIRGINIA AND THEIR CATERPILLARS.

Thomas J. Allen. 1997. University of Pittsburg Press, Pittsburg, Pennsylvania. 388 p.

This is a very good book. The author seems to have good biological knowledge of butterflies & plants, the text is mostly accurate, there are shaded-county maps placed with the species, and best of all, there are nearly 150 color photos of larvae & pupae. The color plates are nice, though many are a little overexposed.

Detailed comments: Most of the hostplants were copied from Opler & Krizek etc., and evidently few hostplants were found in W.Va. Oviposition times based on small sample sizes in Opler & Krizek are repeated with erroneous claims that oviposition occurs only during that part of the day. The book uses the word "perch" for resting, so mate-locating behavior is seldom described. P. xii & 4 has a superfluous line dividing Kanawha Co. P. 32, *Lonicera* & *Syringa* & *Phlox* & *Vicia* are not very popular nectar sources for butterflies, & *Viola* is shunned. P. 50 some of the *Papilio "canadensis"* in this book later proved to be *Papilio appalachiensis* (add 8 co. records for the new *appalachiensis* in E W.Va.), which has the unf marginal stripe like *canadensis* but is claimed by Pavulaan & Wright to be usually larger than *glaucus* & *canadensis* (a bit smaller than *glaucus* in Scriber's studies below). This book claims that *canadensis* is smaller so most of the specimens discussed in the text may represent small spring-form *glaucus* that look like *canadensis*. *P. appalachiensis* has been treated as a distinct sp. with different mtDNA (but later work showed that mtDNA is not very useful for phylogeny), whereas *canadensis* is just a ssp. of *glaucus* tending toward *P. glaucus rutulus* (since the O.D., D. Wright wrote me that his gels of W.Va. *canadensis* represent small *glaucus*, and thus there is no sympatry between *canadensis* & *appalachiensis*). Actually (J. M. Scriber & G. Ordning, Ent. Exp. Appl. 115:247-263; & Scriber, Ordning, & R. Mercader 2008, chap. 6 Introgression & Parapatric Speciation in a Hybrid Zone, p. 69-87 in book *Evol. Of Pops. & Species*), *appalachiensis* would seem to be just a

ssp. of *P. glaucus*, as it is a “hybrid swarm” between *glaucus* & *canadensis* that is obligate-diapause univoltine between the first two flights of *glaucus* thus is allochronic with *glaucus*; it has intermediate degree-day habitat & hw black band width & tails, its larvae can detoxify both *Liriodendron* & aspen, it has the Ldh allozymes of *canadensis* but the Pgd allozymes of *glaucus*, & mostly lacks black females; thus it is basically a hybrid swarm with a few odd enzyme recombinations. Interestingly, Scriber etc. found a similar hybrid zone in C N.Y. & N.Y.-Vermont that is almost the same as *appalachiensis* but a bit smaller in size and has all the Pgd allozymes of both *glaucus* & *canadensis*, & flies after the local *canadensis*; this hybrid zone may be recent as it increased its ability to detoxify *Liriodendron* from 10% to 35% since the 1980s. They note (on p. 71) that these could all be considered ssp. of one species, as the *appalachiensis* & N.Y.-Vt. hybrid swarms are allochronic and look like lab hybrids. P. 54, *Pontia protodice* mates all day. P. 62 *Colias philodice* mates all day long. Larvae of *philodice* & *C. eurytheme*—normally not pupae—hibernate (though Ward Watt states that *eurytheme* lacks organized diapause). P. 74 *Crataegus* is an error for *Feniseca* by C. Abbot. P. 76 half-grown *L. phlaeas* larvae hibernate. P. 77 *L. thoe* oviposits on detritus at hostplant base like most *Lycaena*. Evidently *S. titus watsoni*=*winteri*=*campus* occurs in W.Va. P. 92 *Callophrys polios* ovip. on stems below the flowers, and if *Arc. uvaursi* is absent in W.Va. then the *polios* record must have been an error. P. 99 *humuli* is a syn. of *S. melinus*. P. 104, *Celastrina neglecta* occurs throughout W.Va. in summer, *C. ladon* in E counties in spring, & *C. serotina* “type II” in E counties later in spring. P. 120 *Podophyllum* & *Desmodium* are dubious hosts of *C. Abbot*. P. 122 *S. diana* eggs surely do not take 4 weeks to hatch. Evidently there are no demonstrated *Viola* hosts for *Argynnis* (*Speyeria*) in W.Va. P. 135 *C. nycteis* eggs are not white. P. 137 the unh marginal brown patch in *tharos* only sometimes contains a pale crescent. P. 138 all *Phyciodes* hibernate as unfed 4th-stage larvae. The male on pl. 15 row 2 left has uph like *cocyta*, but it may be *tharos* as flight time & small black antenna club suggest *tharos*. P. 139 W. Edwards’ nearly 100 specimens from Coalburgh Kanawha Co. were mostly *P. cocyta* *selenis*=*marcia*, with some *P. tharos*, according to R. Gatrell; *P. c. selenis* has one generation, while the similar-appearing large *P. diminutor incognitus* Gatrell has two gen. and ranges evidently from NE Ga. and SE Kentucky to E Penn., so should be looked for in W.Va. too. P. 140-1 it would be nice to reexamine W. Edwards’ supposed specimens of *batesii*; the pupa is not more pointed, & most of Shapiro’s “*batesii*” specimens were evidently *P. cocyta*. P. 142 *E. phaeton* eggs surely take less than 3 weeks. P. 147 *P. faunus* ovip. on stems. P. 151 maybe *N. antiopa* has 2 gen. sometimes, as in Colo. P. 156 *V. cardui kershawi* is in Australia. P. 156 *Cirsium* is Asteraceae. P. 157 *V. atalanta* ovip. all day. P. 158, *atalanta* adults hibernate. P. 160 *arthemis*. P. 162 *archippus* mates all day. P. 164 *A. celtis* may ovip. singly or in small clusters. P. 171 *Lethe appalachia* ovip. all day. P. 172 *Cynodon dactylon* is lab host only for *C. gemma*. P. 177 *nephele* is surely just a form in W.Va. P. 180 *plexippus* mates at overwintering sites. P. 195 *E. brizo* egg turns red, & is laid on twigs (the plate “*brizo*” larva is evidently *E. horatius*). P. 199 *E. martialis* eggs turn light orange. P. 205 *P. centaureae* ovip. all day; mature larvae surely turn redder and then diapause (not pupae). P. 207 *P. communis* males mate-locate by resting sometimes also (the behavior described is flaiting, which is patrolling a small area to await females). P. 218 *H. leonardus* larvae don’t have green highlights. P. 220 mature larvae of *H. metea* hibernate. P. 223 *P. themistocles* ovip. all day. P. 225 *P. origenes* mating & ovip. occur all day. P. 225 *Andropogon scoparius*; *Carex scrabrata* is evidently only a lab food. P. 227 Opler & Krizek didn’t say *Paspalum setaceum* was favored (it surely isn’t). P. 228 *P. mystic* mates all day. P. 229 the *W. egeremet* immatures description may refer to *W. otho*. p. 230 *P. verna* probably mates all day. P. 231 *A. campestris* ovip. all day. P. 233 *Anatrytone logan* mates all day. P. 234 *Poa* is dubious for *Poanes hobomok*. P. 239 *E. vestris* mates and ovip. all day; the egg develops a red ring & red top; the error *Cyperus* is a lab host only. P. 240 *A. hianna* surely ovip. all day. P. 243 *A. vialis* mature larvae hibernate. P. 244 sugar cane is a grass.

Plates: Localities are later listed for adults on these plates, but not immatures! According to Pavulaan & Wright, Pl. 3 row 1 left is *appalachiensis* male Rt. 1, Prince, Fayette Co. WV May 22 1987; row 2 right is *appalachiensis* black female East Fork Glady Creek, Randolph Co. WV June 18 1984; row 3 left is *appalachiensis* dusted female E fk Glady Crk. June 18 1984. Pl. 7 should show *E. nicippe* as orange. Pl. 10 “type I violacea” is *C. ladon*, “*C. l. ladon*” is *C. neglecta*; 4th row right is probably true *C. ladon* rather than *C. serotina* “type II”. Pl. 15 4th row are *P. batesii* near-*batesii* from Ont. Pl. 20 top right is *appalachiensis* male from Lanesville, Tucker Co. WV June 18 1973. Pl. 35 upper right is *Celastrina* sp. unknown, probably true *C. ladon* as it is shown on fruits of *Prunus serotina*; 2nd row left is evidently true *C. ladon* as host is *Cornus florida* flower; middle is maybe *Cel. neglectamajor* as claimed but may be *C. neglecta* (see next); right is evidently *C. neglecta* even though it is from *Cimicifuga*, as the photo is the reverse of the *C. neglecta* on p. 69#6 in Allen et al.’s *Caterpillars in the Field & Garden* book, which claims that both species eat *Cimicifuga*; row 3 left is on a leaf of some plant. Pl. 40 “*brizo*” looks like *E. horatius* larva. P. 47 “*tharos*” pupa has small cones (& pl. 37 larva is pale too though a bad photo mostly of uns) so maybe it is *P. cocyta selenis*?

**CATERPILLARS IN THE FIELD AND GARDEN. A FIELD GUIDE
TO THE BUTTERFLY CATERPILLARS OF NORTH AMERICA. Thomas J. Allen,
Jim P. Brock, J. Glassberg. 2005. Oxford Univ. Press, Oxford, England. 232 p.**

This is a very nice book, with lots of great photos of caterpillars of most butterfly species in North America. The book makes no scientific use of the photos, but enterprising scientists can now make use of them to improve classification and phylogeny (a few phylogenetic changes are suggested below). Most of the photos were taken with quality macro lens &

ring flash, so we get good lighting all over the larvae (the only drawback of flash is a tiny ring reflection on shiny heads such as *Chlosyne nycteis* on p. 87 & the two reflecting spots on *Megathymus yuccae* on p. 151 etc., a minor problem compared to the great views of all sides of the larvae [a single flash off to the side produces bad shadows on part of the larva]).

Since not many people like caterpillars, the book will doubtfully make money; it was supported financially by the “binoculars” books of Glassberg. I have thousands of photos of eggs/1st-stage larvae/older larvae/pupae of esp.-Colorado butterflies, but wonder where the \$ can be found to publish those photos, since I doubt that a book on immature butterflies would sell enough to recoup publication LO\$\$E\$, as state books and caterpillar books are vanity-press projects like the usual scientific papers (in which authors must pay page charges & reprint charges etc. and get zero \$ in return).

I’m puzzled: where are the eggs, 1st-stage larvae, and pupae??, don’t they deserve study too? We have the cult of the binoculars, in which people worship butterfly watching and want to throw collectors in jail. Now do we have the cult of the caterpillar, in which books are published with caterpillar photos but no mention is made of egg 1st-stage larva or pupa? Why?? Just another inexplicable hula-hoop craze??

In the text, many related species without photos are still listed beside photographed species, and the text is a hodgepodge of all those species, which only clutters the text and causes confusion. The book uses the NABA checklist for common names on the photos, which causes confusion in identifying some of the photos of difficult groups “stenchospecies”, so one must consult the useful Photo Locations and Credits (p.177-192) to clarify which bookkeeping species they are, as noted below. Many color pages have missing numbers in the sequence (for example p. 45 has photos 1,2,3,4,5,6,8,9, with 7 missing), and in many of those cases the sequence differs in Photo Locations; I have tried to make sense of this below. The missing numbers were evidently lower-quality photos that were deleted at the last minute, throwing the sequences into chaos. They should have simply placed text and photo and photo location and credit in one spot for each species; why scatter them in three places to deliberately produce confusion and errors? Compounding the confusion, the text pages facing the photo pages have numbers on some species, which refer to numbers on the maps, not numbers on the photos. And then where the text got too long, they just oozed out the excess text onto pp. 155-176.

I haven’t studied the text of this book much, as it has mostly just a few hosts and simple stuff like all the Butterfly Watching books, which is often wrong (for instance p. 161 says it is best to search for *Phyciodes* larvae prior to flights, which is actually the worst time as they are solitary in litter during the day then). To find caterpillars, one must know the hibernation stage, which isn’t often given. I would rather have those missing photos included, and eliminate the text.

Detailed comments: P. 5 bottom, if you want to get anything done studying bugs in nature, you should go alone; if you go with other lepidopterists you will waste time talking instead of working. P. 12 says that *Wallengrenia otho* larva carries its shelter with it!, is that true?? (very dubious, it isn’t a caddisfly). P. 16 claims that *Dryas iulia* & *Siproeta exude* acidic fluid from their scoli that causes a skin rash—I wonder if that’s true. P. 20 *Eriogonum* & *Ceanothus* are poor garden plants as they are difficult to grow. P. 28 *polydamas*. P. 35 photo 4 is labeled 35.5 in Photo Locations, evidently just change it to 35.4. P. 37#5 is P. “*marginalis*” *venosa*. P. 43 *Colias meadii johanseni* larva is identical to *meadii*. P. 39 #8 is *E. hyantis lotta*. P. 41#1 is *A. cethura pima*; #2 is *A. sara sara*; #3 has no location but is the *A. sara* (not *julia/stella*) phenotype. P. 43#4 is *Col. occidentalis wasatchia*; #5 is *C. harfordii* (not *alexandra*). P. 49#4 “*dorcas*” is *L. florus dospassosi*. The map should show where *L. xanthoides editha* intergrades with *xanthoides* in SW Ore. & N-C Calif. P. 53 my *Hypaurotis* larvae weren’t pink, is the photo too red? P. 55#4 & p. 57 #7 are both *Satyrrium favonius ilavia* from Hualapai Mts., text p. 54 calls it *Satyrrium favonius* and text p. 56 calls it *Satyrrium ilavia*. P. 57 evidently change photo 9 to 8 to match Photo Locations. P. 58 the *dumetorum* name is horrible here, as it properly applies only to the coastal Calif. bug *viridis* (map shows *C. affinis perplexa*, *C. a. affinis*, & *C. a. apama*), and now *sheridanii* is the same species as *viridis* (the name *dumetorum* should be suppressed by ICZN, and *sheridanii* given priority over *viridis*); p. 59# 1 is Kern Co. *C. affinis perplexa*, #3 is Mendocino Co. *C.* (ignoring evil name *dumetorum*) *sheridanii viridis*, #4 is *C. sheridanii [viridis] neoperplexa*. P. 61 evidently change #9 to 8 like Photo Locations. P. 63 has *C. gryneus gryneus* on photos #1-2, *C. gryneus siva* #3. P. 65#3 looks like *S. istapa* larva, but Photo Locations says it’s from Gainesville Fla., yet bug does not occur in N Fla., so this location must be wrong or it’s the butterfly greenhouse; also, data for inset photo is missing on p. 181. P. 68-69 wrongly lumps various *Celastrina* into *C. ladon* based on NABA dogma: based on photo locations/text, p. 69 photo 4 is *Celastrina lucia* Auct., 5-6 are *Cel. neglecta*, 7 is *Cel. idella*, 8 is *Cel. serotina*. P. 71 1 & 3 are evidently *E. “battoides” ellisii* Kane Co. Utah; 9 is *Plebejus podarce* (not *glandon*) from Tioga Pass Cal. (actually *cassiope*—not *podarce*--is closely-related to “*glandon*” *megalo* and would be considered a ssp. except that *S. Kohler* has found *cassiope* in Mont. maybe near *megalo*). P. 73 photo 4 is *Plebejus scudderi empetri* from N.B. P. 73 photo 5 has no inset, yet p. 182 says it does. P. 75, change #7,8,9 to 6,7,8 to match Photo Locations. P. 77#5 is *A. mormo mormo*. P. 79#1 is *Libythea carinenta bachmanii*. P. 81 photo #7 looks like *callippe*, but there’s an error somewhere, because p. 183 says this photo is from Hondo, Medina Co. Tex., yet *S. callippe* does not occur in Tex. P. 83 #8 is *B. titania grandis*. P. 87 photo 2 is *Chlosyne acastus neumoegei*, 5 is *C. palla calydon*; 8 is *C. acastus sabina*. P. 89 #1 is *C. leanira leanira*, #3 is *C. 1. coronado*. P. 91 #1 is *Euph. anicia maria* from Weber Co. Utah (the photographer Todd Stout told me that *maria* has the same whiter-hairy larva as *E. colon nevadensis* [though another photo on a paler background shows less-noticeable white hairs on the scoli], so perhaps *maria* hybridized with *E. colon* to give them similar larvae, as the adults are also similar; one

might even think that *maria* is a ssp. of *colon*, as its larva is like *colon*, adults are mostly blackish like *colon*, and Todd Stout finds that there is no blend zone with *E. anicia wheeleri*; but the valvae are like *anicia*, so perhaps these similarities are due to tremendous hybridization/introgression with *colon* “*nevadensis*”, as such hybridization has been proven to occur in NE Nevada); #3 is *E. anicia wheeleri* from Tooele Co. Utah; #4 is *E. anicia anicia*Xeurytion (*windi*) from Yellowstone Nat. Park; #5 is *E. chalcedona* near-*olancha* from Kern Can., Kern Co. Calif. P. 93#1 is bad (overexposed, bad view). P. 93, the Photo Locations on p. 184 lists THREE explanations for 93.6, even though there are only two photos; evidently the third from N.B. should be deleted. P. 95 the insert on photo 10 looks like *Hemileuca?* moth larvae, & has no data except photo JG=J. Glassberg. P. 101#1 is *Lim. arthemis astyanax*; #3 *L. a. arizonensis*. P. 103#7 is evidently some kind of moth, as it lacks the known long head horns of *Eunica* and lacks scoli present on *Nymphalinae*. P. 109#1 is evidently *N. areolata* rather than *N. helicta*; #5 has an error, as Photo Locations says this is from W.Va. yet *C. tullia* does not occur in or near W.Va., so maybe photo is *C. tullia* from somewhere else. P. 117 *Codatractus valeriana*=*mysie* does look like it’s related to *Zestusa* & *Cod. arizonensis*, and not to *Thorybes* where it was placed. But why are *Urbanus proteus* & *dorantes* in the same genus? P. 122 *juvenalis*. P. 123, #123.8 & 123.17 are missing on Photo Locations. P. 124 *Pyrgus scriptura* always has several generations. P. 127#11 is obviously *P. alpheus gracielae*. P. 129, why is *Nastra lherminier* so much like *Atrytone arogos*? P. 131 the *Copaeodes minima* looks exactly like *Oarisma garita* top, so either they are extremely similar and it is *Oarisma minima*, or this photo is mislabeled (the *Copaeodes aurantiaca* photo differs greatly). P. 135, #5 missing in Photo Locations. P. 139 #10 is too pink. P. 143 note narrower evil Cyclops eye on *E. pilatka*. P.145 has error, photo 11 *Asbolus capucinus* is only in Fla., yet Photo Location says Arizona, maybe photo is ok and it really came from Fla.? P. 146 says that the spectacular *dracula* fangs on *Amblyscirtes* heads are not used to whack predators as I thought, but are used to drag the leaf nest across the ground—I wonder if that’s true?, it seems very dubious as these aren’t caddisflies either (Butt. Houston describes *A. aesculapius* pupating on the host leaf beyond a stretch chewed down to the midrib, and others who have reared *Amblyscirtes* incl. Richard Heitzman and myself have not noticed caddisfly behavior). *Amblyscirtes* have good traits of head pattern to identify some species/groups. P. 214 Heather Blue does not eat *Dodecatheon*.

FLORIDA BUTTERFLY CATERPILLARS, AND THEIR HOST PLANTS. Marc C. Minno, J. Butler, D. Hall. 2005. University Press of Florida, Gainesville, Florida. 342 p.

This book is amazing. Half the book is on caterpillars with color photos, and the other half is color photos of their hostplants! I assume the book will lose a lot of money due to weak sales, but it’s nice to have the book anyway.

Just a few gripes: Where are the photos of eggs and pupae?, who says that people care about caterpillars but not eggs & pupae?? Technically, the authors should buy a nice ring flash for their macro lens: Canon, Pentax, Sony, Olympus, Sunpak, & Nikon all have nice guide #14 in meters ring flashes. Or at least a dual-flash attached to the lens tip. These are important for butterfly and especially larva photography, because they get closer to the subject when the lens focuses closer, so that more light is placed on the subject and depth of field is greater and necessary f-stop adjustment is much less. Flash provides extra light so that small f-stop can be used to get more of the bug in focus, and it freezes (1/500th sec) the action of a fast-crawling larva etc. And ring-flash provides even illumination all around the subject to put the whole caterpillar in view. Look at *Erynnis baptisiae* larva p. 201 for instance, the authors used sunlight or lamp to light the larva and the underside and the whole front of head is black because of poor illumination; with ring flash this would be a beautiful photo, because a side view of larva with head turned toward the camera is the ideal position to get a great single shot of nearly all of a larva’s parts. One can see with half/most of the book’s photos that they have not used ring flash, as many larvae are too dark or parts are in shadow etc., so the photos in general are not as good as those in Allen’s *Caterpillars in the Field & Garden*. (Those zealots who claim that ring-flash provides unnatural flat illumination compared to one light source at the side, should retire to the art museum where they belong, and never come out again; we need to light up all the parts, not create shadows for art-lovers.)

Details: P. 5 some structures on the larval head are wrong. The worst goof is p. 167 *Nathalis iole* photo, which is actually *Phoebis agarithe* larva from p. 171; and that *agarithe* photo differs from *agarithe* in *Caterp. in Field & Garden*, so is it misidentified too? P. 171 why is *Feniseca* larva whiter than the better photo on p. 31? P. 185 has interesting *Junonia 3* sp. larva photos (the *evarete* & *genoveva* names should be switched back the way Scott book treated them, based on Neild’s Butt. Venez.)(Turner & Parnell [1985 J. Res. Lepid. 24:142-153] reported Mangrove Buckeye to have iridescent turquoise bases of the upper rows of scoli, whereas Tropical Buckeye has iridescent purple bases). P. 193 *M. cymela* photo is too dark. P. 211 *Polites origenes* larva too dark. P. 207 note *Euphyes pilatka* has a narrower “evil Cyclops eye” on top of front of head, which is wider in all other *Euphyes*. P. 215 *Poanes aaroni* & *yehl* photo are too red. P. 296 top doesn’t look like *Colo. Fraxinus pennsylvanica*.

CATERPILLARS OF EASTERN NORTH AMERICA. A Guide to Identification and Natural History. David. L. Wagner. 2005. Princeton University Press, Princeton New Jersey. 512 p.

This great book is mostly about moths, but 51 pages are butterflies. The photos are very nice—very few eggs or pupae again—and seem to be correctly identified. The text below each photo gives identification tips (not very useful as few of a

genus's species are included), plus life history notes, and some hostplants. The book gives only common names of hostplants, a mistake as the people who study moths—and most of these studying butterflies—are going to want to use just latin names of the plants; and there is no list translating these common names into scientific names. It gives some new hosts (for *Erynnis* for instance).

Detailed comments: *Euphyes vestris* chooses dryland sedges in the west, esp. *Carex pensylvanica* which is common in the east too. Hedylidae larva & pupa appear in the Pieridae introduction, which is odd as the larva looks more like Satyrinae. *Pontia protodice* was probably never fully native in the NE, just a stray. *Colias philodice* is polyphagous on legumes and does not prefer *Trifolium* in Colo. *Satyrrium calanus* has amazing larval polymorphism. The book does not give localities of photos, and I had thought that the *Phyciodes phaon* photo was *P. cocyta*, but the same photo reversed is in the Caterp. in Field & Garden book where it is from Collier Co. Fla. Dave Wagner, so it is *P. phaon*, & the white head reported in lit. for phaon evidently may just not be visible on these side views, as the larva looks like Charles Dammers' drawing of phaon. The adult male Phyc. "tharos" has a *P. cocyta* uph, but a tharos antenna club, so may be tharos; young larvae are gregarious, and the first two stages sometimes feed on top of a silk web, never under a silk nest; *P. cocyta* larvae are not pinkish; unfed 4th stage larvae hibernate in all *Phyciodes*. P. 131 has *Asterocampa clyton* photos above *A. celtis* photos evidently.

THE AUDUBON SOCIETY FIELD GUIDE TO NORTH AMERICAN BUTTERFLIES.

Robert M. Pyle. 1981. Alfred A. Knopf, N.Y. 916 p.

I already published a rather complete corrections/review of this book (*J. Res. Lepid.* 20:55-58) so here I give only complete revised photo corrections: Photo 17 is *Phyciodes tharos* (not *mylitta*) larva/pupa, 31 is aberrant *Adelpha*, 89 is *Colias interior*, 108 is *Phoebis agarithe*, 127 is *Kricogonia lyside* female, 165 right is *Ochlodes sylvanoides*, 203 is a reduced-spotting aberrant if it is *Paratrytone snowi* female, 326 & 337 are *Papilio polyxenes* (note black tegulae), 354 & 357 are European *machaon*, 365 is *Libythea carinenta* larvata, 368 left is *Polygonia gracilis zephyrus*, 368 right is *Nymphalis l-album j-album*, 396 is *Satyrrium calanus*, 430 is *Callophrys apama homoperplexa*, 457 is *Callophrys niphon*, 501 is evidently *Plebejus (atrapraetextus?) samuelis* female, 509 is *Celastrina* sp. like *ladon*, 516 is *Lycaena rubidus* female, 543 is *Calephelis nemesi*, 553 is *Emesis ares* female, 555 is *E. zela* female, 557 is *E. chalcedona sierra?* or *E. anicia*, 575 ups is *Phyciodes cocyta selenis* female, 575 uns may be *P. batesii lakota* or *P. c. selenis* male, 590 is *P. pulchella near-pulchella* female, 607 left is *Argynnis (Speyeria) coronis halcyone*, 610 left is *S. mormonia*, 610 right is probably *S. zerene picta*, 613 left is *Boloria bellona*, 630 male is *P. cocyta selenis*, 654 is *Adelpha iphicla*, 662 right is *Asterocampa celtis*, 663 right is *Asterocampa leilia*, 686 is *Smyrna blomfieldia*, 687 is *Diaethria anna*, 732 is *Oeneis taygete*, 737 looks like *O. calais stanislaus* or *O. calais altacordillera*, 743 is evidently *O. calais altacordillera*, 758 is *Hamadryas februa*, 759 is *Hamadryas guatemalena*.

WESTERN BUTTERFLIES . PETERSON FIELD GUIDES. Paul A. Opler.

Illustrated by Amy B. Wright. 1999. Houghton Mifflin Co. New York, N.Y. 540 p.

This book is replete with errors. The paintings are bad (they are wiggly as if viewed under water), many are misidentified, about 50% of the black identification lines seem to be placed at random and not on identification features, and there are errors aplenty on about every page. The 1983 Tilden & Smith field guide that preceded it was a little dated, but was more accurate. The book was produced rapidly on a word processor and little editing on text or plates was done thereafter. (The Brock & Kaufman book has replaced the Peterson Field Guide books for amateurs.)

Plates: Fig. 6, W BC is really in the northwest coast biome, and N Tex. is part of the Great Plains biome (which isn't much of a biome it's so depauperate). P. 32, wyomingo is not known to be reproductively isolated. Middle of page meant *Euph. chalcedona*, not *anicia*. Pl. 4, it's the space not the notch that is viewed from above. Pl. 5, *glaucus* & *canadensis* interbreed extensively. Pl. 8 the words Great Southern White are placed on the wrong bug. Pl. 9, lotta fig. has white scales in bar, legend says it doesn't. Pl. 10, *chrysomelas* border too narrow. Pl. 11, *hecla* fig. looks like *canadensis*, & *canadensis* looks like a narrow-bordered *hecla*. Pl. 14, what paler band inside margin? Pl. 17, text says *longula* has tail. The *comstocki* looks misidentified as band should be sinuous. Pl. 19, the *cassius* unf is not visible. *C. humulus* does not ovip. on leaves. The *amyntula* is too spotted. *Podarce uns* looks surreal. Pl. 24, *atlantis uns* looks like *S. hesperis beani*. Pl. 25 should have shown *uns*, only experts can identify most *Boloria* from ups. Pl. 27, *vesta* unf not shown! Pl. 28, *mylitta* crescent words too far from bug. Pl. 29, *oreas* is really *silenus*. Pl. 30, *astyanax* is p. 332 not TTK. Pl. 31, arrows point to wrong *Asterocampa* spots. Pl. 32, "formosa" is next to wrong bug and is on plate but not on legend. *E. disa* & *mancinus* are transposed. Pl. 33-34 the *Oeneis unf* is not shown, when it should be on *O. alberta* etc. Pl. 34, the *rosovi* upperside has the tiny yellow points. Pl. 36, *Urbanus* arrows misplaced. Pl. 38, *juvenalis* looks like *zarucco*. *Baptisiae* misspelled. *Persius* doesn't look like it. Pl. 39, *mejicana unh* is blue-black. Pl. 40, upper numitor appears to be labeled *Chisos Skipperling*. Pl. 41, *viridis* arrow misplaced. *Origenes* spots too big. Pl. 43, *elissa* & *hegon* have no legs, *elissa* head upside down, *vialis* too brown on unh. Toltec Roadside Skipper is *Amb. tolteca* not *aea*. *Arabus* arrow misplaced. *Panoquina leucas* & *ocola* transposed.

I haven't read most of the text, so a careful study of the text would find a lot more errors. A lot of species are split following tradition, even when there's no data. *Joanae* is missing (it is usually treated as a species but is a ssp. of *Papilio machaon*). *Papilio ornithion* larva can't be smaller as adults are the same size. *P. glaucus alexiades* is in NE Mex. *Rutulus* is not closer to *P. eurymedon*. The phony *Pieris marginalis* range stops at Yukon. P. 157 says *lotta* & *hyantis* are intermediate on Kern Plateau but keeps them as separate sp. anyway! The *Anthocharis "sara"* are split into 4 species, when there can't be more than 2; *stella* doesn't patrol ridgetops. If *Colias eurytheme* were a migrant into Colo., I would have seen individuals migrating, and I haven't. *Colias occidentalis* & *christina* & *alexandra* text & maps are all confused, the *occidentalis* map includes "astraea"/wasatchia etc., *christina* map includes "astraea", *alexandra* map includes *columbiensis*. *C. meadii* is in C Ida. too. P. 173, *Zerene eurydice* lacks a black border only in female. *L. phlaeas* is just a stray in Colo. & Neb. & Dakotas, isn't in Ida. on map, & wasn't introduced from Europe. P. 188, *L. cupreus artemisia* is rather intermediate between Cal. & Rocky Mts. pops. *L. editha* should be *L. xanthoides editha*. *L. dorcas* map is wrong in W Canada. *L. helloides* in Alaska in text, not on map. *L. hermes* host is buckthorn family. *Parrhasius m-album* Colo. record was error. If *Calycopis cecrops* & *isobea* intergrade, lump them. *Fixsenia polingi* map misses Chisos Mts. *Satyrium acadica* was mislabeled W to Ida. *Satyrium californica* host in Ore. is most often *Purshia* (A. Warren Butt. Ore.). The *Call. johnsoni* on plate has black edging of postmedian white line. The *Callophrys gryneus* group bugs are sorted incomprehensibly into species. *Callophrys polios* rait in gulch bottoms, and on host. If *Strymon melinus* genetically swamps *avalona*, then they are ssp. *Brephidium exilis* map is wrong, it's year-round in Mex. and C Valley of Calif. *Leptotes marina* overwinters farther north too. *Zizula cyna* is resident in W Tex. in text, not on map. "Everes" *comyntas* is BC in text, not on map; it occurs in Panama. P. 232, *Gaspé* not Gasp. P. 233 echo probably belongs in *C. neglecta*, certainly not in *C. ladon*; all year in SE Ariz. is a lot more than one flight; *C. ladon* does not range southward as bad map shows. *C. neglecta* has 2-4 gen. The current dogma host-races rather than real species are used in *Euphilotes*. *Baueri* is claimed to have weak fringe checkering, but it is strong on plate. *E. pallescens* and *E. rita* are said to blend on p. 241, so why aren't they lumped? *Philotiella speciosa* is Ore. in text, not on map. Cranberry Blue map has missing gaps. P. 246 *Kamehameha*. *Plebejus acmon* has pink uph band not edged by dark (except in spring), & ranges N to Chelan Wash. That idiotic name *Lupine Blue* for *P. alupini* again. The *Apodemia* treatment is bad: *duryi* is not yellowish & is little different from *mejicanus* except on unh; the lump of *mejicanus* with *deserti* doesn't match the other splitting; *davenporti* is missing; *dialeuca* doesn't belong in this *virgulti* concept; *virgulti* has several gen.; larva is not purplish-blue; too many cooks here. *Apodemia palmeri* has ssp. in Baja Cal. *A. phyciodoides* was mislabeled from Ariz. P. 265, submarginal black dots are not useful to identify *chisosensis*. P. 267 the long palpi mimics a petiole. P. 266 & 268 *Heliconius* coil their pollen-covered proboscis, exude a drop of fluid into the coil, wait for pollen proteins to dissolve into the drop, then suck up the drop with proboscis tip; proteins are NOT absorbed through impermeable-chitin proboscis walls. *Hel. charithonius* range has a silly chicken-head. Winter range of *Euptoieta claudia* is surely underestimated. *Argynnis (Speyeria) cybele* is not in C Colo., and *S. idalia* only strays there. P. 275 photo flower is probably *Machaeranthera bigelovii*. *S. edwardsii* is in E portions of the Rockies. *S. callippe* doesn't occur in S-C Colo. *S. atlantis* in Mont. in text, not on map. *S. hesperis* snaky range in NWT is silly. *Boloria bellona* in Ga. in text, not on map. *B. freija* isn't on alpine tundra. *B. a. distincta* isn't greenish. The larval BODY is orange in *Poladryas minuta minuta*, white in *arachne*. *Chlosyne californica* has orange upf basal spots; text does not distinguish it from *lacinia*. *C. definita* is in Lower Sonoran Zone also. *C. marina* is not on pl. 26. *C. gorgone* larva is black, or varying degrees of red; hole in range map is silly; occurs in coastal S.C.; it was just a stray in Utah-Ida. etc. *Phyciodes texana* crab-claw stray range is silly. P. 304, all the *Phyciodes* have spring f. *marcia*. P. 305 photo, it's the unf pattern of *P. phaon* that is highly contrasting; it doesn't colonize Colo. or Neb. White crescent is variable in the unh marginal patch of *P. tharos* & *cocyta* & *batesii*, but the brown patch is usually narrower & darker in *tharos*, broader & oranger in *cocyta*. (The book doesn't even mention the antenna club, which in *tharos* is usually black in the east and orange in Great Plains, vs. orange in most *cocyta*.) The *cocyta* larva is NOT pinkish as C. Oliver miswrote. The TL of *P. (cocyta) diminutor* was S Minn. not shown on map. Only some *cocyta* hosts (and most eastern *batesii*) have large stem-clasping leaves. *P. batesii* doesn't occur in E S.D. The antenna club of *Phyc. pulchella=pratensis* is orange from Ore.-Mont. to Alaska; the key ID feature of *pulchella* is the orange postmedian-subapical costal unf spot (which has black in *tharos-cocyta-batesii*); the bug on gumweed is a female; *pulchella* does not eat fleabane=*Erigeron*. P. 310 photo is *P. pulchella montana* (uph postmedian band is yellow only at rear). *Phyc. pallida* has one flight. *Phyciodes mylitta*: spelled *Silybum*; Kans. not on map. P. 317 photo legend, replace hindwing by forewing. *Polygonia gracilis* often has two gen. *Nymphalis californica* map: it surely overwinters more places than that (A. Shapiro claims that they only occupy the Cascades to breed in summer, then fly back to lowland Calif. for the winter). Text says *N. vaualbum* breeds in Alaska & NWT, not shown on map. *Aglais milberti* males actually rait (perch) on cliffs/talus just below hilltops. *Vanessa virginiensis* map has silly northern space (unless Scotty on the Space Ship Enterprise beamed it up, if the bug flew north to Alberta, it surely must have gone through Montana on the way), and overwinters far to the north of range shown (Butt. Ohio says it is "only occasionally migratory" and "is one of the first butterflies to appear each spring", based on a lot of data). *Vanessa tameamea*: spelled *Pipturus*. *V. carye annabella* winter range is surely wrong (text says all year in S Ariz.). The *lorquiniXweidemeyerii* hybrid zone is 200 miles wide. Text wrongly claims that *lorquini* occurs in S Ariz.! Map & text disagree on W & S Tex. for *Myscelia ethusa*. *Hamadryas glauconome* orange scales are basal to black crescent. *Marpesia*

chiron map has silly club. *Anaea andria* is native in S Colo., and Butt. & Moths of Missouri writes seeing them fly around in winter there. P. 344 top photo has two white-centered black rings. *Asterocampa clyton* map is wrong, it's native in SE S.D. etc. *Morpho polyphemus* often flies fairly low in my experience. *Lethe eurydice* mate-locates all day. *Cyllopsis pertepida* doesn't occur in W Utah desert. *Megisto rubricata* is bivoltine in Tex. *Cercyonis meadii* isn't in N-C Colo. & S-C Wyo. *Erebia vidleri* map missed Wash. *Erebia stubbendorffii theano* isn't on Vancouver I.; it occurs on grassy places just below or just above timberline. *Mckinleyensis* is an obvious ssp. of *E. magdalena*. *Neominois ridingsii* surely occurs all over E Montana; *wyomingo* is a ssp. *Oeneis (nevadensis) macounii* is on Isle Royale Mich. *O. uhleri* is in S Utah. *Oeneis melissa* text, replace Mt. Katahdin, Me., with Mt. Washington, N.H. *O. philipi "rosovi"* text should have mentioned the strong unh median band. *Danaus plexippus* larva has only 2 pairs of black filaments. P. 377-8, spelled *Ichthyomethia*. *Codatractus valeriana* ranges S to Oaxaca (mysie is a syn.). *Achalarus albociliatus* does not have contrasting pale margin. *Thorybes pylades* range has silly holes from Neb.-Sask. *T. mexicana* LACKS costal fold; *T. drusus* has it. *Cogia outis* is a breeding resident in SW Missouri, and does not stray. *Erynnis brizo* larvae aren't purplish, except probably turning redder in diapause. The unh pale apical spots are greatly reduced in size in *E. propertius*, and reduced or absent in *E. propertius meridianus*, contrary to p. 398. *E. zarucco* in N Ill. in text, not on map. *Erynnis afranius* doesn't occur in Alaska-Yukon, and doesn't nectar on Golden Banner. *E. persius* is not on tundra. *Pyrgus centaureae*, E U.S. range missing on map. *Pyrgus scriptura* is surely all over E Mont. *Pyrgus oileus*, hollyhock host refers to *P. communis*. *Heliopetes domicella* isn't resident in Calif. *H. ericetorum* is native in Wash. *H. laviana* in N.M. & N Tex. in text but not on map. *Pholisora mejicanus* C Tex. in text, not on map. *Hesperopsis alpheus* in Coahuila in text, not on map; occurs 50 miles from *H. a. graciellae*. *Piruna pirus* W Tex. in text, not on map. *Nastra neamathla julia* are ssp. reportedly integrading in Houston; *julia* is native along Colorado R. and in C Tex. *Lerema accius* resident in Ariz. in text, not on map. *Ancyloxypha arene* is native in S Tex. *Oarisma garita* eats numerous grasses & sedges. *Thymelicus lineola* is in Colo. (missed on map) and S Minn. *Hylephila phyleus* another silly map gap in N Tex. Silly amoeba-like gap in N range of *Hesperia uncas* & *H. ottoe* & *Polites themistocles* & *origenes*. *H. juba* hw is not translucent; larvae overwinter, not adults. *H. leonardus montana* has yellow-to-dark brown unh. *H. pahaska* yellow felt does not protrude from stigma (one must use a needle to see the color). *H. attalus* is on mid-tallgrass prairie. *H. dacotae* is W of sassacus. *H. juba* fw is not more pointed than nevada. *Atalopedes* in Man. in text, not on map. *Polites rhesus*, Tex. in text, not on map. *Polites peckius peckius* is in Gunnison/Saguache Cos. Colo. *Polites sabuleti norae* is in Sonora; NW Mont. in text, SW Mont. on map. *P. themistocles* is in N Calif. & Ore. Silly hollows in taiga range of *Polites mystic*. *Polites sonora*: replace *sonorensis* with *sonora* & *sirius* with *siris*; *sonora* & *mystic* seldom fly in same habitat. *P. vibex* in Ohio in text, not on map. *Pompieus verna* no map. *Atrytone arogos*, Ill. in text, not on map. Mex.-El Salvador *Anatrytone* is *A. mazai*, and *logan* isn't in Ariz. *Ochlodes yuma* is in C Calif.; *Ida*. not on map. *Poanes hobomok* is in Capitan Mts. of C N.M.; *Poa* is an error. *Poanes zabulon* is not in N.M. (& not on map). *P. melane*, N.M. not on map. *Partrytone snowi*, pine dropseed is an error. *Euphyes bimacula*, S Va. not on map. *E. vestris* lacks blacker veins, & its head is not very orange; Alta. not on map. *Atrytonopsis hianna*, Ont. not on map (*deva* is a ssp. of *hianna*). *A. vierecki* is also in grassland. *A. python* does have hourglass spot. Pyle's rotten Sheep Skipper name again! for *A. ovinia edwardsii* (the sheep Bob referred to are only in N Ariz.). P. 458, replace TK by 426. *Amblyscirtes aenus* map, with its silly amoeba-like protrusions, looks like Bullwinkle the Moose! *A. tolteca f. prenda* is wet-season form. *A. nereus* unh is not green. *A. eos* is just a rare stray in S Colo.; more odd gaps in map in C Tex. *A. vialis* silly gaps in N-C Alta. & N plains. *A. phylace* fw is not squared, and isn't in C Mex. P. 466, roadside skippers don't have stubbier wings. *Calpodus ethlius* occurs in N Fla.-S Ala. *Panoquina panoquinoides errans* does not wander. *Nyctelius*, Calif. not on map. P. 470, only *Megathymus* larvae move up & down in their burrow. *Agathymus neumoegeni* isn't in S Ariz. as shown; scabrous *Agave* is an error. *A. aryxna baueri*, Calif. not on map. P. 537, *idalia* not *dalia*.

A FIELD GUIDE TO WESTERN BUTTERFLIES. The Peterson Field Guide Series. J. W. Tilden & Arthur C. Smith. 1986. Houghton Mifflin Co., Boston, Massachusetts. 370 p.

This is a better book than the above, despite getting my name wrong on p. viii. The plates are nice photos, though some are b/w. The book is old and out of print, so I won't make many comments, and will ignore various minor misspellings and taxonomic progress. I wrote to Tilden about new host records in book, and he replied & clarified some*.

Plates: Pl. 1 fig. 4 is toothed-line *C. pyracmon* spring f. *nabokovi*. Pl. 11 fig. 6 is brownish on uns as is *rusticus*. Pl. 13 fig. 6 should be green. Pl. 16 fig. 1 is *Phyciodes cocyta*. Pl. 26 fig. 3 female is *C. columbiensis*. Pl. 28 fig. 1 is *Colias columbiensis pseudocolumbiensis* male. Pl. 28 fig. 6 female is *E. salome*. Pl. 30 fig. 8 is *Satyrrium sylvinus nootka*. Pl. 42 fig. 3, *Ochlodes sylvanoides*. Pl. 44 fig. 13 is *Nastra julia*. Pl. 45 fig. 6 is *Pyrgus scriptura* spring f. *pseudoxanthus*.

P. 44, *damei* is a form of *C. sthenele*, not *pegala*. P. 53, *excubitor* was named by 4 authors. *O. alberta* is an error from Mt. Graham. P. 64, *Vaccinium staminium* was a dubious Abbot & Smith host. *Crataegus* is an error for *Nymphalis antiopa* as Tilden (*pers. comm. 1986) could not find data. *Jatropha* is an error for *Anartia jatrophae*. *Boloria selene* has 3 gen., *nebraskensis* isn't in N.D., & *sabulocollis* is spelling. P. 89 *Uncompahgre*. *Phyciodes texana*: *Beloperone* used only in lab. *P. picta* & *Microtia (Texola) elada*, *Siphonoglossa* eaten only in lab. *Papilio eurymedon* is recorded from *Ceanothus cordulatus* & *thyriflorus*, dubious? *Isomeris arborea* is an error* for *Pontia protodice*. *Colias behrii*, *Gentiana newberryi* is error. *Colias nastes*, *Trifolium repens* is lab only. *Eurema lisa*, *Trifolium* is lab only. *Anthocharis cethura*, the *Sisymbrium*

host is an error for *Descurainia pinnata*. *Satyrium liparops*, *Crataegus* & *Acer negundo* are both errors. *Callophrys comstocki* doesn't eat *Eriog. umbellatum*. *Callophrys gryneus* assoc. *Juniperus scopulorum* in W Texas*. *Euristrymon polingi*, *Quercus grisea* has no data*. E. ontario, *Crataegus* is error, *Quercus turbinella* is assoc. only. *Strymon melinus*, *Yucca* is new*. *Lycaena cupreus*: spelled *Rumex paucifolius*. *Echinargus isola* assoc. *Prosopis glandulosa* var. *torreyana**=mesquite & *P. pubescens**=screwbean. *Plebejus acmon*, *Trifolium* & *Medicago sativa* are errors. P. 204, replace midtibia by hind tibia. *Agathymus neumogeni chisosensis* is not on *Agave scabra*. *A. aryxna baueri* eats *A. chrysantha*, not *parryi*. *Lerodea eufala* & *Polites vibex*, *Stenotaphrum* eaten only lab. All the *Amblyscirtes vialus* hosts are lab only. *Atrytonopsis hianna deva* isn't in Colo. or Utah. *Euphyes vestris*, *Cyperus* is lab only. *O. pratincola* is evidently *O. sylvanoides*. *Ancyloxypha arene*, *Polygogon semiverticillata* is assoc.* only. *Celotes limpia* is not recorded on *Sida*. *Heliopetes laviana*, *Sphaeralcea* is dubious as source unknown*. *Pyrgus ruralis*, *Horkelia tenuiloba* is based on Lember's Sierra Nevada *Horkelia fusca*, & *tenuiloba* is not in Sierra Nevada anyway; *H. bolanderi clevelandii* is assoc. only. *Erynnis afranius*, *Lotus scoparius* is new.

A FIELD GUIDE TO EASTERN BUTTERFLIES. Peterson Field Guides. Paul A. Opler. Illustrated by Vichai Malikul. 1992. Houghton Mifflin Co., Boston, Massachusetts. 396 p.

This is a better book than the western field guide.

The paintings on the plates are good, though usually printed too small; sometimes wing shapes are a little off. Pl. 18 *Colias palaeno* has a red ring misplaced on wing (real *palaeno* lacks ring, it looks like *C. pelidne*, and if it is *pelidne* it can't be from Churchill). Pl. 18 *C. occidentalis* from Fort Calgary BC is *C. occid. christina* from Calgary Alta. Pl. 22, *C. hesseli* has unf line broken. Pl. 25, *Celastrina "ebenina"* female looks like *C. ladon* female. Pl. 27, *Libytheana* is *carinenta* larvata. Pl. 28, *fulvia* female not labeled on plate. Pl. 30, *Chlosyne marina* painting & text is not the *Chlosyne melitaeoides* that occurs in Tex. Pl. 32 uses Tropical Buckeye under bug for *genoveva*. Pl. 38, "chryxus" is *O. calais strigulosa*. Pl. 43, arrow wrong on *poweshiek*. Pl. 48, *Panoquina "fusina"* doesn't look like the weak-banded Mex. ssp. *evansi*. *P. panoquin* has no dash, so looks like *P. ocola*.

Only common names are given for hosts (for instance, the host of *Lycaena dione* Broad Dock, is what?, presumably *Rumex obtusifolius*, which is Broad-leaved Dock in Britton/Brown flora. Front of *Papilio machaon joanae* head is black, not yellow. P. 56, E Mex. has *Papilio glaucus alexiars*. P. 71, *Coliadinae* females are usually larger than males. P. 73, only *Colias alexandra* has 2 flights (not "broods") in Sask. Map missed *Colias occidentalis krauthi* in S.D. P. 93, *Lycaena phlaeas* overwinters as half-grown larva (& it was not introduced from Europe). *Satyrium titus* host is chokecherry not chokeberry. P. 121, no Balloon-Vine Hairstreak in book. *Heliconius* don't absorb pollen proteins through proboscis wall! P. 156, *Boloria eunomia* has silver unh spots. P. 162, *leanira* has priority over *fulvia*. P. 170, *Phyciodes cocyta* (*selenis*) larva & spines are not pinkish. *Phyciodes picta*, *Siphonoglossa* is lab host only. *Polygonia satyrus* has two flights. *Polygonia progne* uph is black on outer half only in summer gen.; elm is dubious host. Nfld. missing from *Nymphalis vau-album* map. *N. californica* map has silly stray hook amid distribution dots. *Nymphalis urticae* is an artificial import. *Vanessa cardui kershawi* occurs on Australia. *Chrysopelea* is a ssp. of *Anartia lytrea*. P. 191 *Adelpha* map has silly dip in E N.M. P. 203, *A. celtis* oviposits singly. There is no blend zone of *Lethe eurydice fumosus* & *eurydice* in Neb. *Neonympha mitchellii* was in NW, not NE, Ohio. There are not two *Megisto* sp. in Fla. as shown. *Erebia stubbendorffii* *theano* is also in subalpine zone. P. 218, *Oeneis* overwinter as young & old larvae. *Danaus plexippus* larva has tubercles only on T1 & A8. P. 244, SE Kansas is native on map, stray in text. P. 259, it's E. *persius* with numerous long white hairs on upf. Hollyhock & mallows hosts belong to *Pyrgus communis*, not *P. oileus*. *Pholisora catullus*, mints are errors. *Nastra neamathla* is native in La. & Miss. *Hesperia uncas* is native NW of Minneapolis. *H. pahaska* is absent in E Tex.-La. Silly space in *Polites mystic* map in Dakotas. *Poanes taxiles* larva head is uniformly red-brown. *Bayensis* is just a ssp. of *Euphyes dion*. P. 303 all *Amblyscirtes* like flowers. P. 335, *hutchinsi* is a ssp. of *Euphydryas editha*. The "life list" wastes 31 pages!

BUTTERFLIES OF NORTH AMERICA. Jim Brock & Kenn Kaufman, 2003.

Kaufman Focus Guides. Houghton Mifflin Co. New York, New York. 384 p.

This is a good little book, a pocket "field guide", which has taken over the function of the Petersen Field Guides. The book looks similar to the nice Golden Field Guide to Birds. The photos are very nice, and distracting background greenery is removed. There isn't much in the book about the butterflies' biology or behavior, mostly just a brief mention of hostplants, as the book concentrates on identification. Most of the photos are correctly identified. The book lumps some of the more closely-related species and often confuses the result (there isn't room to explain difficult bugs), and splits the usual species--I won't comment on most of the taxonomic controversies. If there are very distinctive-looking ssp., the book often illustrates some as "variations" without naming them. Most maps have just one color, but many maps that have both light-green and dark-green are confusing, as p. 11 says light green means uncommon vs. dark-green common; most such maps show dark-green vs. dark-green as roughly the overwintering vs. summer migrant range, but not quite, and that distinction is contrary to p. 11 that describes both colors as "summer, or more than one season"; and *Junonia coenia* uses

the colors to say that it is not common in the west, which is quite wrong; *Danaus plexippus* is less common all over the Great Basin; *Neominois* and most *Erebia* and *Oeneis jutta* are mapped wholly in light-green implying wrongly that they are uncommon; *Pyrgus ruralis* is shown in error to be common in the U.S. and uncommon in Canada; *Cercyonis meadii* & many *Hesperiinae* etc. are wrongly shown to be uncommon in the north or west; these colors in *Amblyscirtes* make no sense at all. Thus these map colors are very poorly chosen and are usually defective and should be ignored. Details: P. 98, the orange cap is nearly always present in *S. californica*. P. 21 *P. canadensis* photos are close to *P. glaucus rutulus*. P. 24 *P. eurymedon* flaits (patrols a small area) in tiny clearings in ridgetop trees all day to await females. P. 33 middle photo is yellow form Bruce's Swt. which extends into Canada. P. 34 *P. indra* raits (perches to await females) on cliffs below hilltops. P. 55 "widespread form" is C-N Calif. ssp. *hyantis*. *Anthocharis sara* lumps *A. julia*. P. 62 confuses *Colias alexandra/C. christina/C. occidentalis*, each one an artificial mixture of 2-3 species; the orange male on bottom is *christina*. The *C. occidentalis* photos are a mixture of *occidentalis?* and *occid. sacajawea* or *occid. wasatchia*. The two *pelidne* uns look more like *scudderii*. P. 65 male uns looks like *gigantea*. P. 84 the Rocky Mts. Canadian/Alaska *L. florus* are wrongly lumped into *L. dorcas*; the salt marsh bug is in N.B. too. P. 112, the *Callophrys dumetorum/perplexa* species is ridiculously confused, as *C. viridis* "dumetorum" is a different species from *C. affinis perplexa* (*affinis* name is older than *perplexa*, and *viridis* is same bug as older-but-soon-to-be-banished *dumetorum*, and Coastal is *viridis* "dumetorum", and the name *sheridanii* should be protected from *viridis* by ICZN so that the coastal bug should be called *C. sheridanii viridis*). P. 126 the pale variant of *Leptotes marina* is *L. cassius*. P. 130 *C. humulus* is similar to *C. neglecta*, not spring *C. ladon*. Cherry Gall Azure missed. *C. nigra* has larger unh submarginal black dots. P. 132 *P. acmon* lumps *acmon* & *P. alupini* (top middle male is evidently *P. alupini lutzii*). The female *P. "lupinus"* is *P. chlorina monticola*, the male looks like *P. chl. monticola*, & the uns may be *P. alupini alupini*. P. 136 *podarce* is a distinct species. P. 144 *arizonensis* is ssp. of *C. rawsoni*. P. 156 *Dryas iulia* misspelled. P. 161 the upper left photo of Great Basin Frit. is the Montana ssp. *albrighti* female. P. 163 the middle left pale male of Callippe Frit. is *Argynnis (Speyeria) coronis* male. P. 164 *nokomis* map missed N Ariz., C Ariz., S Ariz., NE N.M., & SE N.M. populations. P. 167 the upper left photo of "Northwestern" Atlantis Fritillary is obviously *S. hydasphe rhodope* (top middle and top right are ssp. *nausicaa*, bottom left and bottom right ssp. *ratonensis*, bottom middle ssp. *irene* or its clones. P. 174 *montinus* & *chariclea* are ssp. of *B. titania*. *Natazhati* is the Pleistocene Fritillary. P. 176 *P. batesii* map missed Wyo., Neb., N Ariz. P. 178 *P. pulchella=campestris* has orange antenna club from Ore.-Mont. to Alaska. P. 179 the female *orseis* is *P. pulchella montana*. P. 183 the Black Crescent uns is male *P. frisia tulcis*. P. 184 *C. whitneyi damoetas* is darker in Colo. P. 187 *Chlosyne gabbii* photos say Utah where *gabbii* does not occur (*acastus* is in Utah). P. 187 ssp. *hoffmanni* (middle photo) does not occur in Ore. P. 192 *minuta* map missed C Tex. & S-C Tex. P. 196 *P. satyrus* averages a little paler in July gen. P. 198 *progne* uph is mostly black only in summer. *P. oreas* map missed C Wyo. & Utah. P. 202 *N. antiopa* raits (males perch to await females) in gulches, not patrols. P. 204 *Vanessa cardui* doesn't overwinter in north as map shows. P. 207 the upper right tropical buckeye is a mangrove buckeye; reverse the names *evarete* & *genoveva* again. The upper right *P. genoveva* may be *evarete*. *Nigrosuffusa* is a ssp. P. 225 Cream Banded missing on plate. P. 230 *do rubricata* males really hilltop in midmorning?? (dubious). P. 232 *P. xicaque allyni*; *Paramacera* misspelled. P. 237 the middle *C. sthenele* is *C. oetus oetus*, and the left is probably *oetus* too. P. 242 *E. disa* has more obvious white band. *E. callias* & *E. stubbendorffii theano* etc. hosts see Scott *Papilio* (N.S.) #6. P. 244 *E. fasciata* host cottonsedge. P. 248 *O. alpina* occurs near Great Slave Lake. P. 260 *T. pylades* raits just off the hilltop—not on top—all day. P. 265 Coyote Cl. photo too red. P. 281 both *zarucco* ups may be *E. baptisiae?*, *valva* is often needed to separate these. P. 282 *E. propertius meridianus*. P. 292 *H. ericetorum* is common native in Wash. too. *Heliopyrgus* would seem to be a subgenus of *Heliopetes*, as the genitalia of *Heliopetes purgia* is similar. P. 295 *Xenophanes* photo has weird gray areas, evidently a photoshop error. Hoary Skipper photo is odd too. P. 297 male ups "hayhurstii" is *mazans* (hw margin more scalloped in *hayhurstii*). P. 306 *prittwitzii* dot occurs farther E in W Tex. In *O. poweshiek* the black unh anal margin is diagnostic. P. 313 right male "juba" I think is *H. comma*; lower right *H. "attalus"* looks like *H. ottoe*. P. 315 right uns male "pahaska" looks more like *H. viridis*, & the left ups female & right uns female "Apache" look like *H. pahaska*. P. 317 "pawnee" uns female looks like *H. l. montana* or *pawneeXleonardus*. P. 321 "macswaini" is *H. juba*. P. 329 Woodland Sk. should have a yellow-unh variation photo too. P. 334 "berryi" female ups is *Problema byssus*. P. 338 *At. hianna* occurs in E Colo. & NE N.M.; *loammi* is ssp. of *A. hianna* (as is *deva*). P. 342 *linda* is ssp. of *A. aenus*. P. 344 *reversa* unh is not reversed (background is just darker). P. 346 *Notamblyscirtes simius* males rait (perch to await females) on hilltops, they don't "patrol below hill summits" (Brock has unfortunately confused the mate-locating behavior of quite a number of species, see Butt. Ariz.). P. 352 *dysaules* is a variety of *arabus*. P. 355 upper right "Ocola" looks like *P. lucas=sylvicola*. P. 371 Jeffrey R. Slotten.

BUTTERFLIES THROUGH BINOCULARS. A Field Guide to Butterflies in the Boston-New York-Washington Region. Jeffrey Glassberg. 1993. Oxford Univ. Press, N.Y., N.Y. 245 p.

This book was the start of his series of "binoculars" books, using nice photos from nature. The text of the "Binoculars" books doesn't have much scientific content, so I mostly just correct the photos of these books.

P. 34, Christopher Curtis once told me that *Lycaena epixanthe* occurred in the commercial cranberry bog at Whitesbog N.J. P. 55, the *P. batesii* at Cresheim Pa. may have been *Phyciodes cocyta*; all the N.J. & Philadelphia area records are

dubious, and there were no 20th century records until Shapiro's dubious records which were not verified. P. 57, P. progne overwinters Oct.-May. Plate 5 fig. 7 is spring form male protodice. Pl. 15#4 is Phyciodes phaon from way south, with orange antennae, not batesii. Pl. 17#4 is Polygonia faunus faunus. Pl. 19#3 line positioned wrong.

BUTTERFLIES THROUGH BINOCULARS. THE EAST. Jeffrey Glassberg. 1999.

Oxford University Press, New York, New York. 246 p.

The photos are nice in this book (only the blurred one on plate 42#4 is bad), which is intended for butterfly identification for butterfly watchers. P. 29 & 183 note that the U.S. Forest Service exterminated a colony of *Atrytone arogos* in Fla. by burning it. P. 32 blames the demise of *Neonympha mitchellii* on one collector, then p. 136 blames its demise on a group of greedy immoral collectors (I wrote to ask for details but got no response), but Ray Stanford told me it was killed by successional overgrowth of vegetation along a power line due to cessation of mowing following the abandonment of the power line. Despite the title, this book does not cover most of N Canada.

P. 124, *lupini* misspelled *lupinus*. P. 131, *Asterocampa celtis* males usually land on people's heads to mate-locate (raiting behavior). *Asclepias* is an error for *Danaus eresimus*.

The plate numbers should have been printed above the photos, and not just on the side of the page to the left. Plate 23#12 is *Glaucopsyche lygdamus* male, not *Celastrina*. Plate 24#7 is *Plebejus anna* (or maybe *Pleb. atrapraetextus fridayi*) from Calif. Plate 32#9 is not *cocyta* but is *P. batesii batesii* from N.C. (and note the background leaf compared to photo 7). On plate 32 reverse the numbers #9 & 10 below the photos & leave the other words the same, as 9 is *batesii* & 10 is *P. cocyta* (note that the leaf on #7 & 9 is the same). Plate 34 fig. 8 looks like *Polygonia faunus*. Plate 44#3 is *Oeneis calais stanislaus*, not the very different *O. calais strigulosa* occurring in the east. Plate 55#4 is *Lerodea eufala* female (*neamathla* has upf spots weak and only in male). Plate 55#6 is maybe *Nastra julia* but the spots are too white. P. 222 missed Scott's book.

BUTTERFLIES THROUGH BINOCULARS. THE WEST. Jeffrey Glassberg. 2001.

Oxford University Press. New York, New York. 374 p.

The title adds "A field guide to the butterflies of Western North America", but the book actually only covers western U.S. The photos are very nice, and this is a good book for butterfly watchers. P. 49 photo #3 is *Papilio zelicaon* female. P. 57#5 is *Euchloe ausonides* I think. P. 59 fig. 5 is *Anth. julia browningi*. P. 67#3 is *Colias cesonia*. P. 83#8 is *Lycaena florus near-florus* male, #2 is *L. florus near-megaloceras* female. P. 97 photos 3 & 4 are reversed? as 4 is greener. P. 123 #3 is *Plebejus atrapraetextus*, #4 has too much orange to be *anna* & may be *Pleb. atrapraetextus fridayi*, #5 male & 6 inset female are *Pleb. atrapraetextus fridayi* from Sonora Pass. P. 129#5 is probably *Calephelis nemesis*, & #4 may be *nemesis* also. P. 133#3 is *Apodemia virgulti deserti*. P. 141 fig. 5 is *S. atlantis sorocko*, #3-4 & #6-8 are *S. hesperis*. P. 149#3 is *Argynnis (Speyeria) zerene platina*. P. 153#2 is *Boloria freija* female. P. 167#5-6 are *Chlosyne harrisii hanhami*. P. 169#3 inset looks more like *C. palla*. P. 174 *P. pulchella "campestris"* antenna clubs are orange from Ore.-Mont. northward, as on fig. 1 on p. 175. P. 177, *P. tharos* has mostly black antenna clubs from E S.D. S to S Kans. & Tex. eastward, W to Ariz.-W Colo. *P. batesii* also has orange clubs in Utah-Colo.-Wyo. P. 177 photo 7 is *Phyciodes batesii batesii* from N.C., 8 is evidently *batesii lakota* from McNair Minn. (the same female as fig. 10 of the EAST book) or could be *P. cocyta*. P. 240 *P. xanthus* has a strong X on upf, and I've never seen a postbasal white uph dot on *P. scriptura*. P. 244 the *Pholisora catullus* map is wrong and the unh is never bluish. P. 253#1 is too red on unh & #2 has the spots too whitish, but maybe these are *Nastra julia*. P. 259#5 is *Ochlodes sylvanoides*. P. 260 *H. attalus* occurs on tall/midgrass--not shortgrass--prairie. P. 263#11 is what? P. 264#10-12 are ssp. *peckius* from Ariz., while the Great Plains has ssp. *surllano* with narrow unh patch. P. 267#8 is ssp. *sabuleti* not *chusca*. P. 276, an opera singer is a diva, not a deva. P. 286, *Notamblyscirtes simius* is a hilltopper, not a roadside skipper. P. 289#4 is *Panoquina lucas=sylvicola*. P. 308 I haven't seen any valid legume host records for *lutzi* or *texana*.

BUTTERFLIES THROUGH BINOCULARS. A Field, Finding, and Gardening Guide to Butterflies in FLORIDA. Jeffrey Glassberg, Marc C. Minno, John V. Calhoun. 2000.

Oxford University Press, New York, New York. 242 p.

The photos in this book are very nice, and all or nearly all are properly identified! The photo of *Celastrina "ladon"* on pl. 16 is odd and almost looks like *C. nigra*, but the date is more like *C. neglecta*, is it *ladon*? The *Polites origenes* female on pl. 34 may be *P. themistocles*. The *Poanes yehl* female on pl. 38 looks like *P. viator*, but maybe it is *yehl*. The little identification bars are good, showing nice little identification spots on the uns of *Eurema* for instance.

BUTTERFLIES of the CAROLINAS FIELD GUIDE (2003, 414 p.), BUTTERFLIES of OHIO FIELD GUIDE (2004, 344 p.), BUTTERFLIES of MICHIGAN FIELD GUIDE, (2005, 376 p.),

(not seen: Butterflies of Georgia Field Guide, & Butterflies of Florida Field Guide),
all by **Jaret C. Daniels**, Adventure Publications Inc., Cambridge, Minnesota.

Here's a new strategy: mass production of state field guides for butterflies. The text and photos are prepared for each species in a master file, and then a state book is assembled by just using pages from the master book file for the species occurring in that state (each page does include a little map of the state with the range shown). On the positive side, Daniels is a professional nature photographer at Univ. of Florida, and the main photos are nice and are big (3-4"), and he evidently used the butterfly experts there because nearly all the photos are properly identified. Little photos are often included of the other sex and undersides and larvae, and the larva photos are nice, though small. The text for each species includes the usual stuff plus a short description of larva and hostplants. There is a good conversion list of plant common names to scientific names. Mass production of field guides would be a good idea, if the master file contained expert information, and a local expert added local color for each state. The only problem is that lepidopterists are poor, and state butterfly books are money losers, so how can this series succeed even with those efficiencies of mass production?, will it continue? My main gripe is that the species are arranged in each book in what appears to be random/haphazard fashion. The species are intended to be arranged by a conspicuous color, such as black, blue, brown, white, yellow, etc., and the color is shown on the edge of the page. But the result appears to be a haphazard arrangement of species in each book that I don't like, and often the same species will be repeated! just 20+/- pages onward, if one sex differs from the other, and both copies have identical text with just the big & little adult photos switched between male and female, which wastes a lot of space.

If identification were the goal, these books should have just reproduced tiny photos of each species all ganged by similar appearance in a 20-page section in the front of each book, with brief name & page # below each tiny photo, directing the reader to the full-page treatment later on.

I haven't checked most of the hosts and other detailed info in these books, as it is the usual info. Below are a few corrections on the photos, etc.

Butterflies of the Carolinas: *Pyrgus centaureae* is not a stray. *Celastrina ladon* isn't everywhere in the Carolinas. P. 113 mistakenly prints the name *Callophrys irus* for *niphon*. P. 150 photo looks like *Wallengrenia egeremet* rather than otho to me. I wonder if the larvae of *Poanes aaroni* p. 302 & *P. viator* p. 326 are really that pink? *Euphyes berryi* is misspelled *beryi*. *Phyciodes selenis* is properly called *P. cocyta*, which ranges S to W.Va., but the bug in N.C. should be called *P. diminutor incognitus*, which is not a stray. The *Euphyes pilatka* photo is too orange.

Butterflies of Ohio: The photo of *Amblyscirtes hegon* looks strange, as if the bottom of the hw were replaced by the abdomen of a blue. The photo of *Wallengrenia egeremet* looks like *W. otho vesuria* from Jamaica. *Phyciodes selenis* is properly called *P. cocyta*, which p. 269 says is not present in Ohio; actually *P. diminutor diminutor* occurs all across northern Ohio, at least in Lucas, Wood, Sandusky, & Columbiana Cos., as I noted in *Papilio* (New Series) #13; it resembles *P. cocyta* but has two gen. The small ventral photo of *Argynnis* (*Speyeria*) *aphrodite* looks like *S. atlantis* to me. I found *Speyeria idalia* common in the Cleveland suburb of North Royalton about 1961. Text lists *Pieris virginienensis* in NW Ohio, but map shows it in NE.

Butterflies of Michigan: The photo of *Oarisma poweshiek* looks like *O. garita*, it's too tawny for *poweshiek*. The photo of *Wallengrenia egeremet* looks like *W. otho vesuria* from Jamaica. The *Erynnis horatius* larva is probably *E. juvenalis*. The photo of *Oeneis chryxus* is real *O. chryxus* which occurs in the Rocky Mts., and is quite unlike Mich. *O. calais strigulosa*. The *Asterocampa celtis* photo is evidently the Florida ssp. The photo of *Phyciodes batesii* is ssp. *batesii* from N.C., it's not ssp. *lakota* from Mich., which has a large brown unshaded marginal patch. The photo of *Polygonia gracilis* is *P. g. zephyrus* from W N.A. not Mich. ssp. *gracilis*. I haven't seen any *Oeneis macounii* females with that much brown on upf base, but I have one *O. nevadensis* female with almost that much brown, so I assume the little female *O. macounii* photo is *nevadensis*. The ventral *Speyeria aphrodite* photo is *S. atlantis*. The *Euchloe olympia ups* is aberrant.

SWALLOWTAIL BUTTERFLIES OF THE AMERICAS. A Study in Biological Dynamics, Ecological Diversity, Biosystematics, and Conservation. Hamilton Tyler, Keith S. Brown Jr., Kent Wilson, 1994. Scientific Publishers, Gainesville, Florida. 377 p.

Keith Brown is one of the foremost experts on latin american butterflies, and Kent Wilson has studied *Papilionidae* for more than 50 years.

This book reminds me of a "Dilbert" cartoon by Scott Adams, showing Dilbert giving a presentation to an audience. Panel one: "This next transparency is an incomprehensible jumble of complexity and undefined acronyms." Panel two: "You might wonder why I'm going to show it to you since the only possible result is to lower your opinion of my communication skills." Panel three, as Dilbert points to the tortuous mess on the wall screen: "Frankly, it's because I like making complex pictures more than I like you." Please Keith, do not write any more complex scientific papers, just give your info to someone who knows how to effectively communicate information. The goal is to publish data accurately so that the original data can be reconstructed, and present it so that users can easily find what they want. I gave my *Papilionidae* hostplant card file info to Keith, and I see in this book that little of it was used, and one cannot look up the hosts and reconstruct the original bug-plant hostplant records, as the host data has been simplified/changed so much. It

takes quite a while to figure out all the complexity in this book, the jumble of numbers and letters in the plates, the positioning of photos of a taxon in multiple places, the oozing of plate legends from one plate to the next, the overlapping map ranges, the abbreviations for everything, lines & arrows going everywhere, everything that possibly could be made confusingly complicated has been thus mesmerized. Instead of one easy index, there are three, so one must first search for the right index. Every table and fig. and box is a new puzzle for deciphering. This book is a perfect example of how it would be easiest to just put the bug's name under each photo, to aid readers and reduce errors (then put the text on the internet so that it could be fixed and continually updated). The deliberate attempt to complicate everything (notable in Keith's prior papers on *Heliconius* etc.) has introduced numerous errors, which would take many months and a lot of expertise to fully decipher; it would be easier to start over with a new book. And a lot of the book concerns general principles of mimicry and other biology that Keith evidently teaches in his classes, but frankly, *Papilio* lovers don't really want a book loaded down with that general stuff, which forces the *Papilio* treatments to be compressed (folded, spindled, & mutilated like the original IBM data cards) so much that data retrieval is difficult. And I doubt that many college biology students really care about *Papilios* anyway. Of course there is a lot of good stuff in this book that will be of use to people, though it will take some study to extract, and verification of extracted detailed information will be desirable. So, we need a new book.

The book lumps previously-oversplit species using the biological species concept, which is good, because in popular (and some nonpopular) groups of butterflies, every new variant was named as a new species. But in North America, I still treat *Papilio zelicaon* & *P. polyxenes* as distinct species. *P. zelicaon* & *P. p. coloro* overlap a little in range at the E side of the mts. from San Diego Co. to Kern Co. Calif., where they don't interbreed. True, in C Colo. you find a lot of intermediates between *zelicaon* & *polyxenes*, but there are also a lot of intermediates between *P. machaon bairdii/brucei* & *polyxenes*, & between *P. m. bairdii/brucei* and *zelicaon* (I have many drawers full of the various intermediates and forms, as there is great variation within each, for instance each critter varies completely from wide-banded to narrow-banded, etc.), yet the three species still exist and have not merged, so I keep them all as three species (they are bookkeeping species within the stenochospecies *P. machaon*). In Alta. *machaon* & *zelicaon* have merged in some areas and are distinct in others, according to F. Sperling. I do treat *joanae* as *P. machaon joanae* though as the book does, as it has various eyespot etc. traits & mtDNA of *bairdii* (though mtDNA has proven to be comparatively useless for the study of phylogeny), and is reproductively compatible with *P. machaon*, even though introgression has made it look more like *polyxenes*. The book treats *brevicauda* as a ssp. of *P. machaon* also, which is reasonable as the few crossing experiments showed compatibility; *brevicauda* looks different but its major characteristics (lack of sexual dimorphism, black form) also occur in other *machaon* ssp., and the amount of orange varies within *brevicauda* (*gaspeensis* has little ups orange, *bretonensis* is intermediate between *gaspeensis* & the orangish *brevicauda* thus invalid). *P. machaon brucei=dodi* is a valid yellow-form ssp. Pl. 91: S is *brucei* not *gaspeensis*; R is *kahli* from Riding Mts. Man.; H is aberrant *nitra* female from Little Belt Mts. Mont.; 8 is really *nitra* from Pine Creek Calgary Alta.; V is *polyxenes*; 2 is aberrant. Pl. 92: S is female, probably of *zelicaon* (listed also for *coloro*); Q is not named but is looks like *pseudoamericanus*, maybe from Mex. In *Papilio indra*, *phyllisae* is a syn. of *indra*, & *nevadensis* is just a variable intergrade. Pl. 90, *kaibabensis* is on S Rim of Grand Can. also. The book correctly treats *rutulus* as a ssp. of *Pap. glaucus*, but *maynardi* is a syn. of *glaucus*. Note how *P. glaucus alexiares* has the mtDNA & allozymes of *glaucus* (p. 147-148) and has some black females like *glaucus*, yet has the wing pattern & valva of *rutulus* (p. 321); so why do misguided people continue to treat *rutulus* as a distinct sp. from *glaucus* (and treat *canadensis* as a distinct sp. from *glaucus*, despite extensive intergradation)? *Arcticus* is a form of *canadensis* resembling *rutulus*. *Texanus* & *ilioneus* are syns. of *Papilio troilus*, and *fakahatcheensis* Gatrell was later named for the S Fla. ssp. *Magnus* & *xanthus* & *sayii* & *montanulus* & *yukonensis* are syns. of *Parnassius phoebus smintheus*, *maximus* is a valid ssp. in C Mont. with blackish females, & *golovinus*, *alaskensis*, & *elias* are syns. of *apricatus*. The "unnamed ssp." of *P. phoebus* from "Region of Ciudad Victoria, Mexico" fig. on pl. 50 is obviously just mislabeled Rocky Mts. *smintheus*, maybe from Colo. where Arthur Moeck also collected. In *P. clodius*, *strohbeeni* is a distinct ssp., but the others are basically synonyms in a variable species in which few bugs in any locality look different (a ridiculous farce of naming ssp. only by locality without actual distinguishing features). Pl. 52, 18 is not indexed & B is given two names; there are too many names for Antillean *Battus polydamas* ssp. P. 61, males emerge a day or two earlier than females ("protandry") in discrete-generation insects in order to place the maximum population of sexually-active males during the time when most females are emerging, for three reasons: in order to maximize the number of matings for males, & minimize the time required for females to mate, & maximize the number of offspring (J. Scott, J. Anim. Ecol. 46:909-924, 1977). P. 114, *Papilio troilus* larvae look like snake mimics. P. 127, fig. 5.6 does not exist. P. 192, the several dozen rain forest refugia Brown concludes have not fared so well in studies of other creatures. There are one too many veins on Fig. 11.2 #B. The *Papilio homothoas* that I caught in Colombia surely is a ssp. of *P. cresphontes*; *melonius* probably is also. Pl. 64, S on photo has "sera" above it so must be *serapis*. P. 27 *Baronia* is not *transandean*.

The book gives "Fast Keys" to adults and larvae, and maybe they do mostly work. Unfortunately keys are bad for identification purposes, because just one mistake in the sequential chain of a dozen comparisons will lead one to a grossly wrong identification. Perfection at every couplet is required to successfully identify something with a key, while butterflies are variable and are far from perfect. Tables are the proper way to identify butterflies, for many reasons: one can compare

all the traits to the bug to see if there is a genuine best match, one can quickly read the most distinctive traits that are the easiest to use, variation can be included, the tables double as a data repository, adding a new sp. to a table is easy but very difficult to a key, etc. etc. The human brain excels at pattern recognition, which is closest to what is done by simultaneously considering all the traits in a table. It's difficult and time-consuming to construct keys, and considering their large failure rate during use, one should spend the time making tables instead, which have actual lasting value.

DISCUSSION

How to write butterfly books. Here are some general thoughts for those people who contemplate writing butterfly books. If you are writing a book on some state or province, you will not make any money. When your manuscript is finished, send it to some real butterfly experts to try to get the errors corrected. Show it to some ordinary people to get their opinion as to its organization, ease of use, quality of illustrations, etc. Put the map and ideally the photos on the same page as the text. Don't use crummy paintings of butterflies. Do not illustrate flying butterflies in the mounted position (with the rear edge of both forewings parallel) as butterflies don't fly that way, and lepidopterists will know you are an idiot if you position them this way. If you don't put the photos with the text, group them in one section of plates, and save money by showing only left half ups next to right half uns of each individual. Don't waste space repeating info (don't describe the bug if there's a picture [most books have useless descriptions of the adult butterfly which nobody ever reads] and limit descriptions of photographed bugs to distinguishing features; don't describe distribution if there's a map; etc.). Get rid of "checklists", they are useless waste of space, and nobody looks at them either. Combine all indexes into one. Do not use just common names of plants, because those common names are difficult to convert to scientific names (I have all the big floras, and looking up the common names in those always leaves one with a residue of common names that cannot be deciphered without uncertainty or error). (Most amateurs know very few plants, so they will know very few of the common names of plants anyway, so you might as well just give the scientific names of the plants. And as a comparative "expert" on plants myself, I try not to learn the stupid common names of plants. The few people who are experts on plants and are interested in butterflies, want to see plant scientific names, not common names.) I know you will copy much of the information from other books, but try to copy from books that are mostly accurate, and if you write something very different or unusual, try to document it at least with a slight reference as to its origin in your state or whatever. (Book authors tend to copy each other a lot, and most of the hosts listed are generally copied from other states without a statement that few or none are from the current state, so lots of old mistakes such as confusing the hostplants of *Pyrgus oileus* & *communis*=*syrichtus*, and stating that *Phyciodes cocyta* has pinkish spines, are later repeated in dozens of books, each book copying the one before.)

Necessary butterfly books. The current system of writing books produces too much repetition of past errors and not enough presentation of current better knowledge. What we really need, is a Wikipedia-style internet encyclopedia of all butterfly species in the world, the text for each prepared by expert(s) on that group and corrected continuously as new knowledge and new experts arise, with synonymy, information on hostplants and biology and behavior with references, and low-res jpg color photos of the egg larva pupa & adult. Science publishing has become so fragmented and expensive that it is now impossible to keep up with the literature, and even large libraries can obtain only a very small fraction of publications, because their money goes to a few "must have" \$2000./year big journals, so a wikipedia-style butterfly reference authored by thousands of people is necessary. There is controversy of taxonomic treatment in many groups, and many taxonomists are vehemently opposed to the other view, so maybe the encyclopedia would have to accept several different treatments for a group for a few decades until the controversy is settled, or an editor would have to enforce the stenospecies =superspecies & bookkeeping species =semispecies organization upon the taxa. We need a simple moth book covering all the moths of North America, which would in vol. 1 simply illustrate every species, 100 per page, one half ups next to other half uns, with just one species or ssp. name & sex below each, so that 300 pages could illustrate two each of 15,000 species. Then vol. 2 would be the text identifying each illustration, and providing information on the species & drawings etc., which would be wikipedia-style coauthored by hundreds of people and updated as information expands. We need a Butterflies of Latin America with the same format. Ditto for Africa, tropical Asia, etc. Can we make these books without \$support from a super-rich person or government? Maybe not. (A total reform of science publishing is also necessary. Each scientific paper would have to be submitted to one of three repositories, and if it passes minimum standards, it would be duplicated to the other two repositories, and only the abstract would be published (online). Journal & magazine & book publishers such as *Oecologia* & *Science* etc. would have to bid for the right to publish articles, and at last authors would receive a fair portion of that money; few would get bids of course, so the majority of articles would be "published" only by copies ordered by purchasers. Individuals could subscribe to all the papers with a given subject matter, at the lowest-possible cost of transmission [such as Adobe Acrobat] consistent with financially maintaining the system.)

ACKNOWLEDGEMENTS

I thank John Rawlins of Carnegie Museum for helping with some matters regarding William Henry Edwards' types. Virginia Scott of Univ. of Colorado Museum helped me research butterflies there. Many lepidopterists offered opinions as to identification of photos and other matters in these books over the decades, which often found their way into these corrections. The authors of many of these books helped decipher some of the questions I had with their books, as noted

above, and those clarifications are listed herein. Norbert G. Kondla & Crispin Guppy contributed some corrections to the Pelham Catalogue, and Kondla contributed corrections to other books. Jonathan Pelham kindly sent some old papers and advice.

PAPILIO BONUS

Bad Taxonomist's Credo. If I named it, it is a species. If you named it, it is a subspecies. If you named it and I don't like you, it is a synonym.

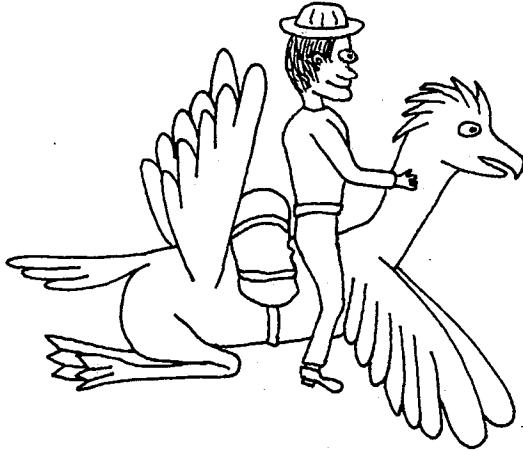
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1. New Papilionoidea and Hesperioidea from North America. James A. Scott, 1981, 1-12, \$2.00
2. The life history and ecology of an alpine relict, *Boloria improba acrocneuma* (Lepidoptera: Nymphalidae), illustrating a new mathematical population census method. James A. Scott, 1982, 1-12, \$2.00
3. Distribution of Caribbean Butterflies. James A. Scott, 1986, 1-26, \$2.50
4. Larval hostplant records for butterflies and skippers (mainly from western U.S.), with notes on their natural history. James A. Scott, 1986, 1-37, \$3.00
5. The courtship of *Phyciodes*, and the relationship between *Phyciodes tharos tharos* and *Phyciodes tharos morpheus* (= *pascoensis*) in Colorado. James A. Scott, 1986, 1-8, \$1.00
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7. Biology and systematics of *Phyciodes* (*Phyciodes*). James A. Scott, 1994, 1-120, \$9.00
8. *Speyeria hesperis* and *Speyeria atlantis* are separate species. James A. Scott, Norbert G. Kondla, and Stephen M. Spomer, 1998, 1-31, \$3.00
9. A new *Celastrina* from the eastern slope of Colorado. James A. Scott & David M. Wright, 1998, 1-15, \$2.00
10. *Phyciodes* (*Phyciodes*): new discoveries, new subspecies, and convergence. James A. Scott, 1998, 1-42, \$4.00
11. New western North American butterflies. James A. Scott & Michael S. Fisher, 1998, 1-12, \$1.00
12. Taxonomic Studies and New Taxa of North American butterflies. James A. Scott, Michael S. Fisher, Norbert G. Kondla, Steve Kohler, Crispin S. Guppy, Stephen M. Spomer, and B. Chris Schmidt, 2006. 74 p. & 6 color pl., \$14.00
13. *Phyciodes* (*Phyciodes*): More Progress. James A. Scott, 2006, 38 p., \$7.00
14. Butterfly Hostplant Records 1992-2005, with a treatise on the evolution of *Erynnis*, and a note on new terminology for mate-locating behavior. James A. Scott, 2006, 74 p., \$10.00
15. Building the California Academy Drawer. James A. Scott, 2006, 40 p., \$6.00
16. Portable (Six Drawer) Cabinets for California Academy Drawers. James A. Scott, 2006, 10 p., \$1.50
17. Proposals for a new insect study, commerce, and conservation law that deregulates dead insects, and proposals for fixing the endangered species act as applied to insects. James A. Scott, 2006, 17 p., \$3.50
18. Geographic variation and new taxa of western North American butterflies, especially from Colorado. James A. Scott & Michael S. Fisher, with some parts by David M. Wright, Stephen M. Spomer, Norbert G. Kondla, Todd Stout, Matthew C. Garhart, & Gary M. Marrone, 2008, 84 p., 10 figs., 5 color plates, \$9.00
19. Corrections/reviews of 58 North American butterfly books. James A. Scott, 2008, 129 p., \$8.00
20. Biological Catalogue of North American butterflies. James A. Scott, 2008, 51 p., \$5.00

NOTE: PAPILIO (NEW SERIES) appears irregularly. It is mailed free to the British Museum (Natural History); others must pay. There is no subscription. Instead of subscription charges, persons desiring reprints should request them from authors, enclosing the advertised price. Any new name or nomenclatural act in this publication is intended for permanent, public, scientific record. Manuscripts must be scientifically sound and readable, but are not edited for format or style or length. To eliminate page charges and reprint charges (all charges demanded by the traditional vanity press scientific journals), publication delays, correcting proofs, and printer's errors, accepted papers are reproduced by modern quality photo/print methods by the author(s), dated, and mailed by the author(s). Mss. should be sent to Dr. James A. Scott, 60 Estes Street, Lakewood, Colorado 80226 U.S.A. "Papilio Bonus" parts are diversions from the regular scientific content—political or sarcastic commentaries or purely humorous cartoons or writings—concerning some aspect of entomology.

PAPILIO BONUS: DR. BOB'S SECOND EXPEDITION TO THE LAND OF HUMONG

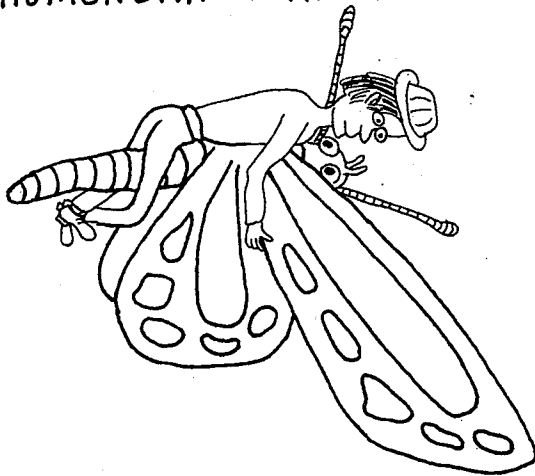
DR. BOB'S SECOND TRIP TO HUMONG BEGAN WITH A SPEEDY FLIGHT ON THE QUADDUCKOSAUR



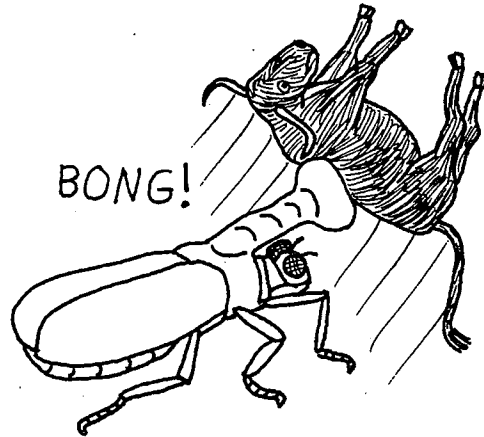
THEN HE RODE A WINGLESS HORKEY TO PACK MORE STUFF



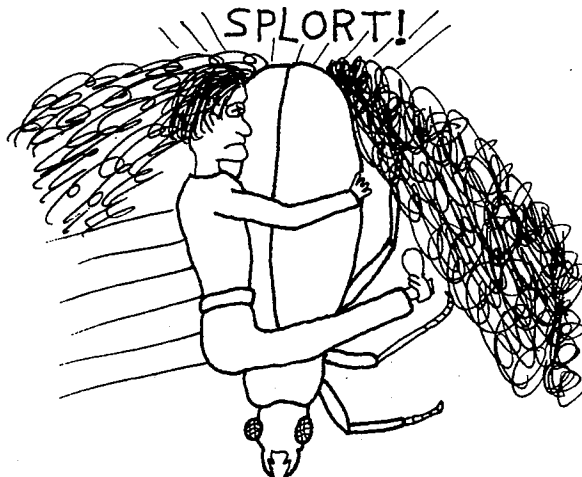
DR. BOB CAPTURES THE HUMONGAN BIRDWING



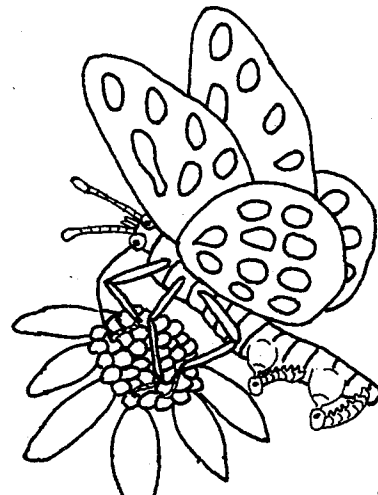
THE COWABONGA BEETLE DOES HIS THING



WHILE PURSUING A HUMONGAN FRITITILLARY, DR. BOB DAGWOODED A BOMBARDIER BEETLE



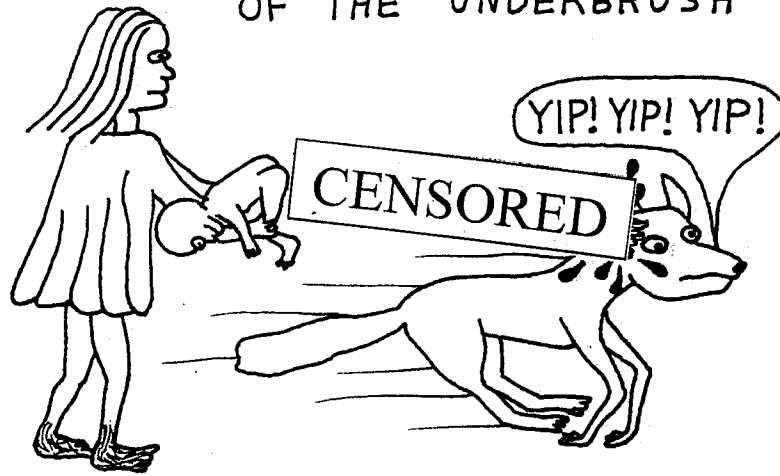
THE LADY FRITITILLARY NURSES HER YOUNG FRITITILLARIPILLARS



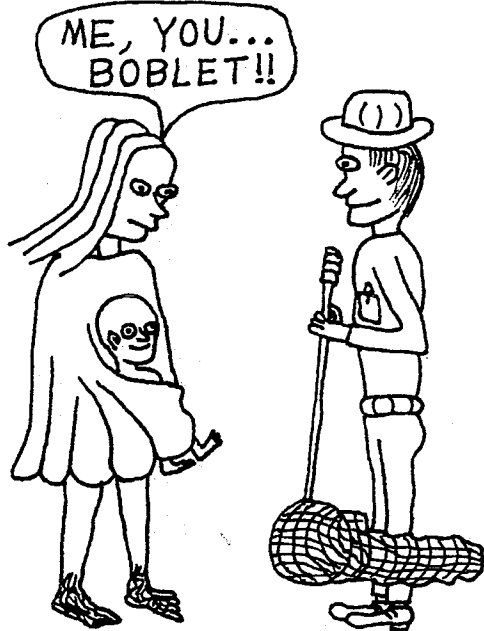
DR. BOB IS PURSUED BY THE VICIOUS HUMONGAN FOX. IS THIS THE END FOR DR. BOB?



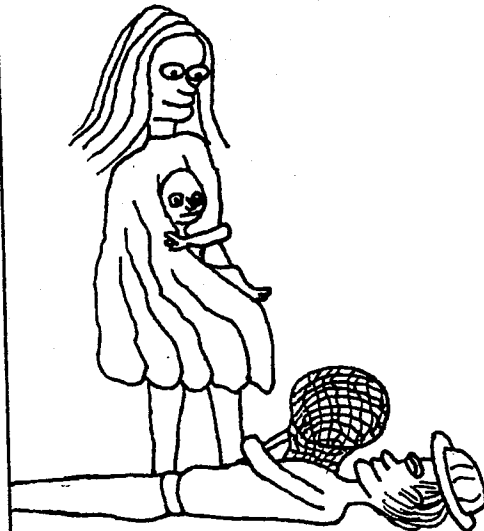
THEN WABOOJI WOMAN POPS OUT OF THE UNDERBRUSH

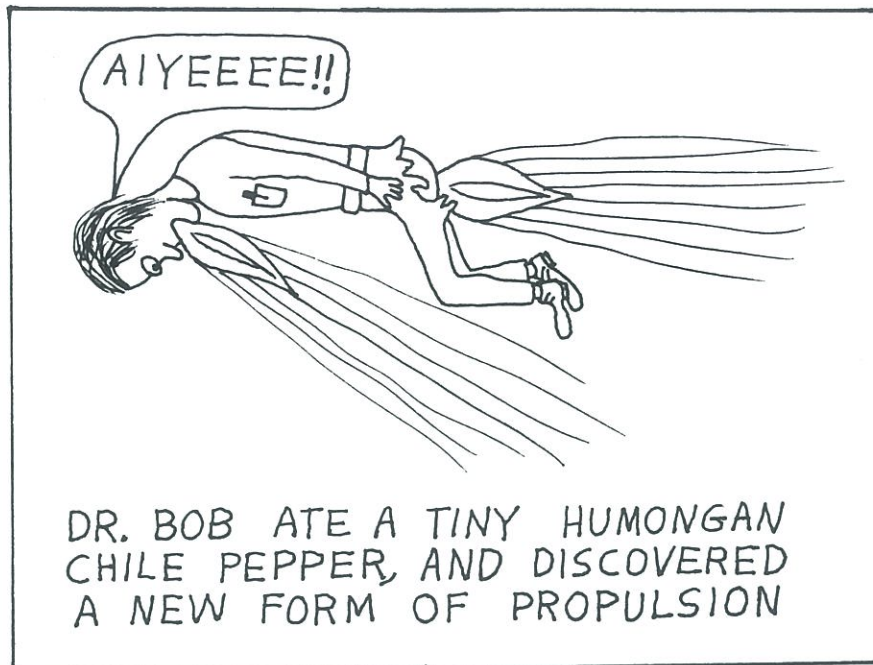


DR. BOB AND WABOOJI WOMAN STARE INTENTLY



DR. BOB WAS FEARLESS, BUT FATHERHOOD WAS DIFFERENT





DR. BOB ATE A TINY HUMONGAN CHILE PEPPER, AND DISCOVERED A NEW FORM OF PROPULSION

BOBLET CAPTURES HIS FIRST BUTTERFLY

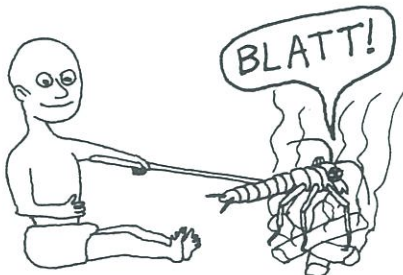


SO YOUNG, YET ALREADY AN EXPERT BONER

LIFE WAS GREAT IN HUMONG: FAMILY, WABOOJI JUICE, AND ROASTED HUMONGAHOPPER FEMURS

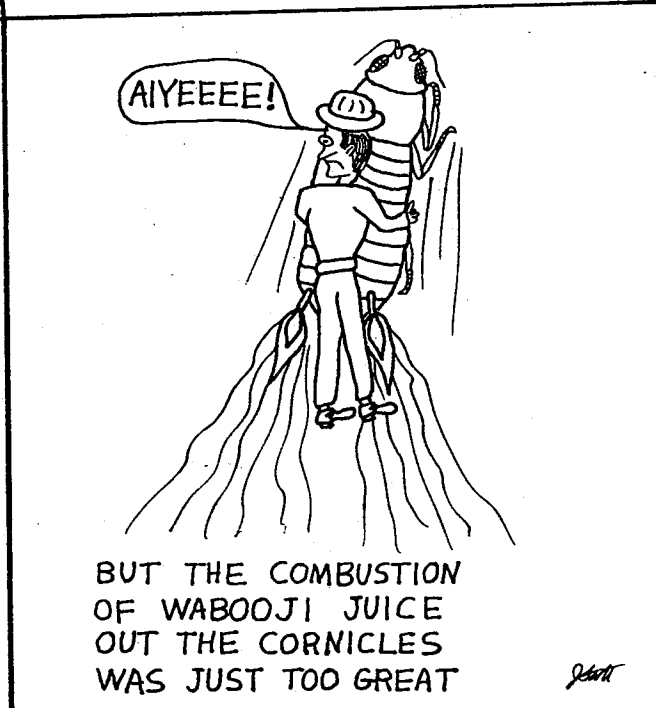
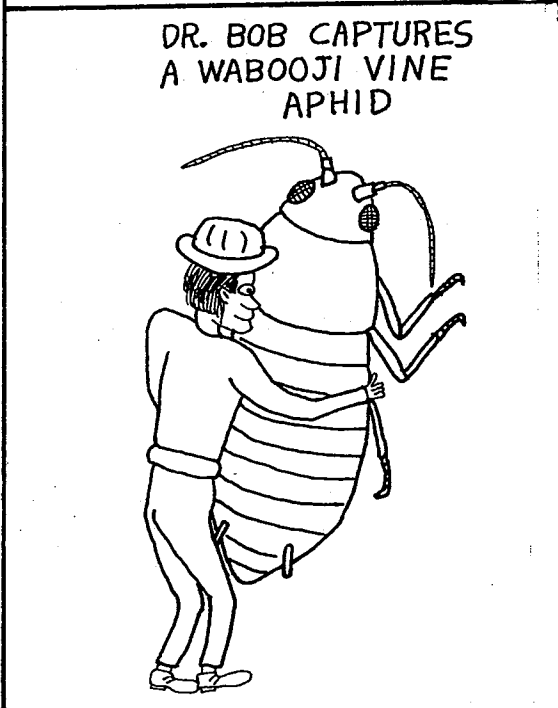
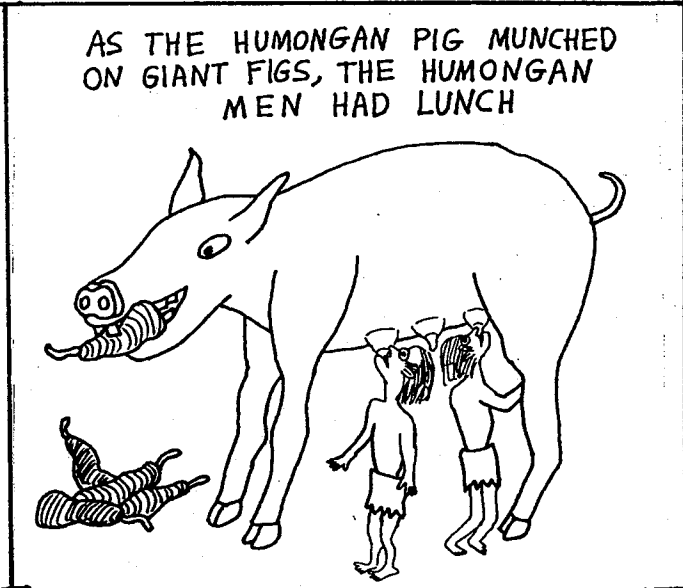
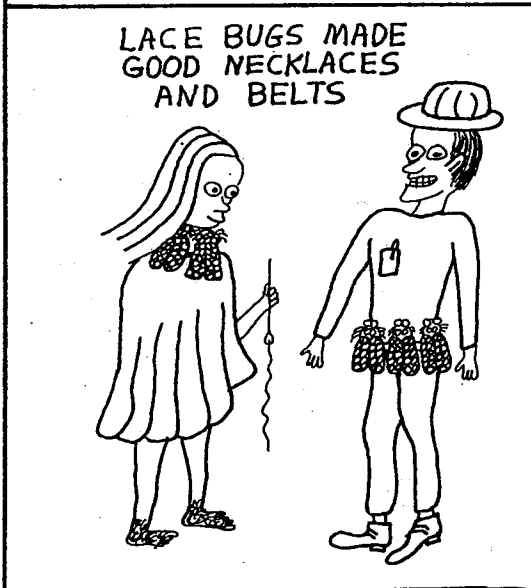
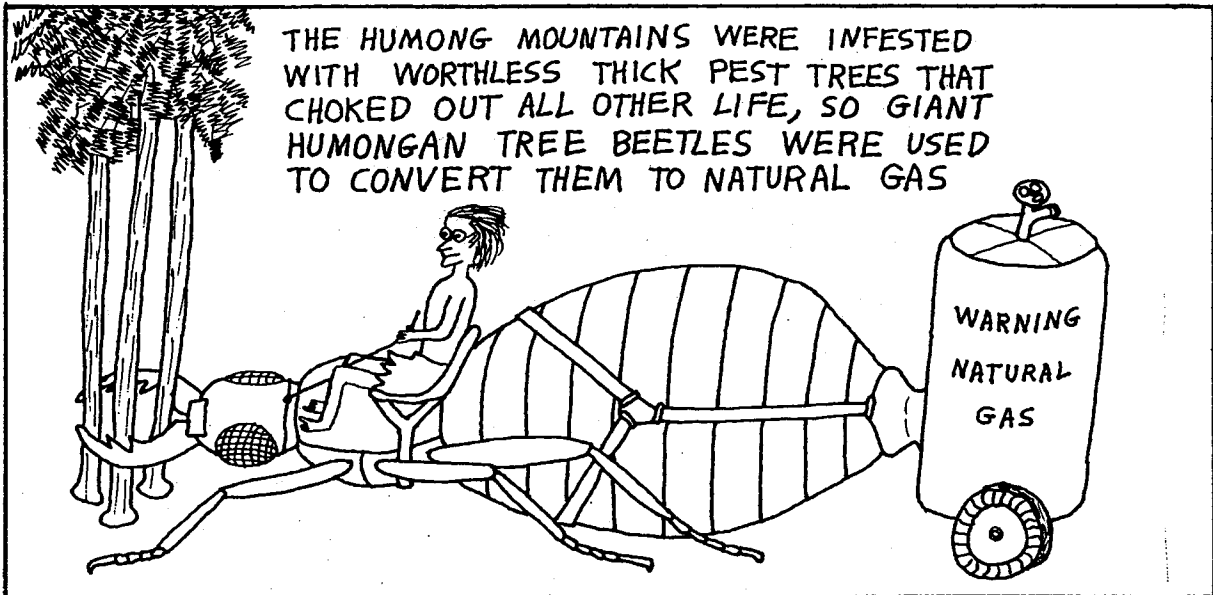


BOBLET LIKED THE GRILLOBLATTIDS THE BEST, BECAUSE WHEN GRILLED THEY MADE A FUNNY SOUND



UH-OH, BOBLET JUST ATE THE HOLOTYPE

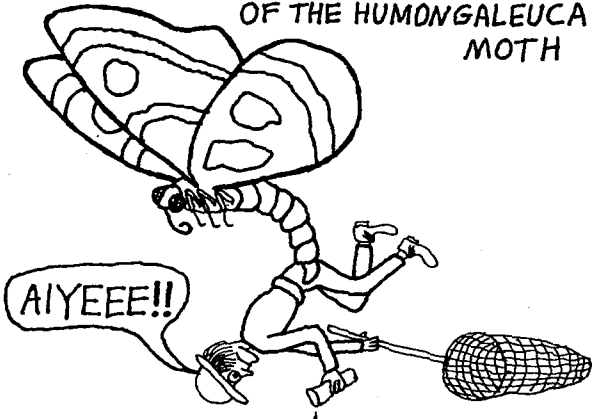




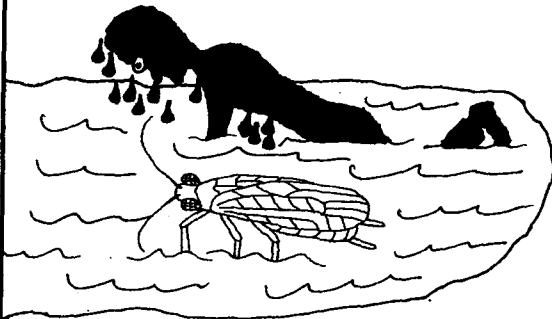
STONED STONEFLIES
WERE CAPTURED WITH
WABOOJI JUICE



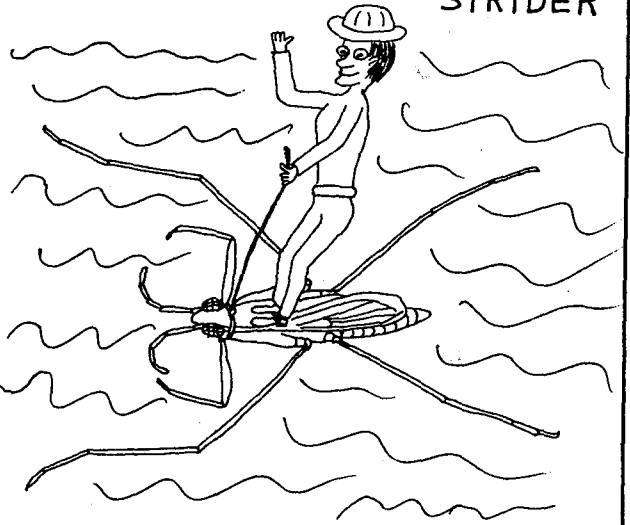
DR. BOB'S WABOOJI JUICE WAS JUST
TOO SIMILAR TO THE FEMALE PHEROMONE
OF THE HUMONGALEUCA
MOTH



THE MUDFLY WAS JUST
TOO QUICK FOR DR. BOB



DR. BOB TAKES A FAST RIDE
ON THE HUMONGOUS WATER
STRIDER



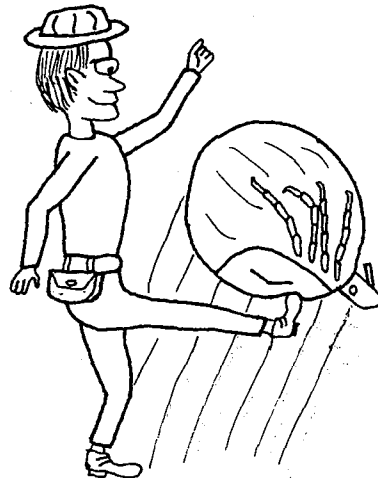
DR BOB WAS
ATTACKED BY
THE HUMONGAN
TICK



LUCKILY, DR. BOB
HAD A FULL CAN
OF CHEESE FIZZ



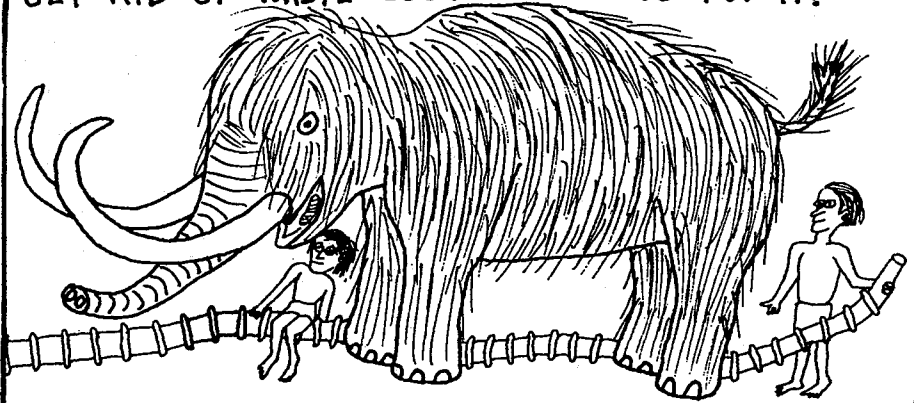
NOW THE HUMONGANS
HAD A NEW GAME
—TICK KICK



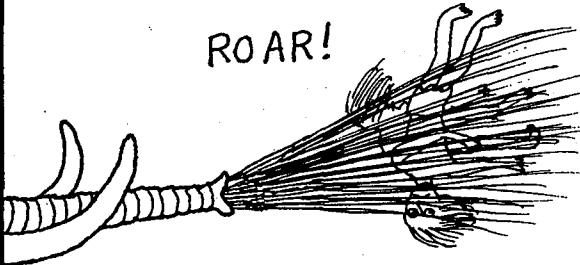
TO FERTILIZE THEIR GARDENS & WABOOJI VINES, THE HUMONGANS COPIED DUNG BEETLES AND USED WOOLLY MAMMOTH POO



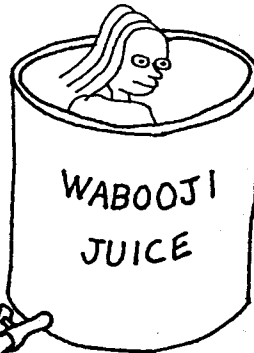
THEY USED MAMMOTH POO TO GENERATE NATURAL GAS ALSO. BUT AFTER LISTENING TO DR. BOB WARN OF GLOBAL WARMING, THEY DECIDED THEY MUST GET RID OF WASTE CO2. WHERE TO PUT IT?



ROAR!

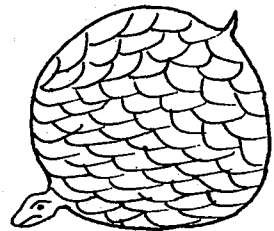


THAT'S NOT THE PLACE TO PUT CO2!

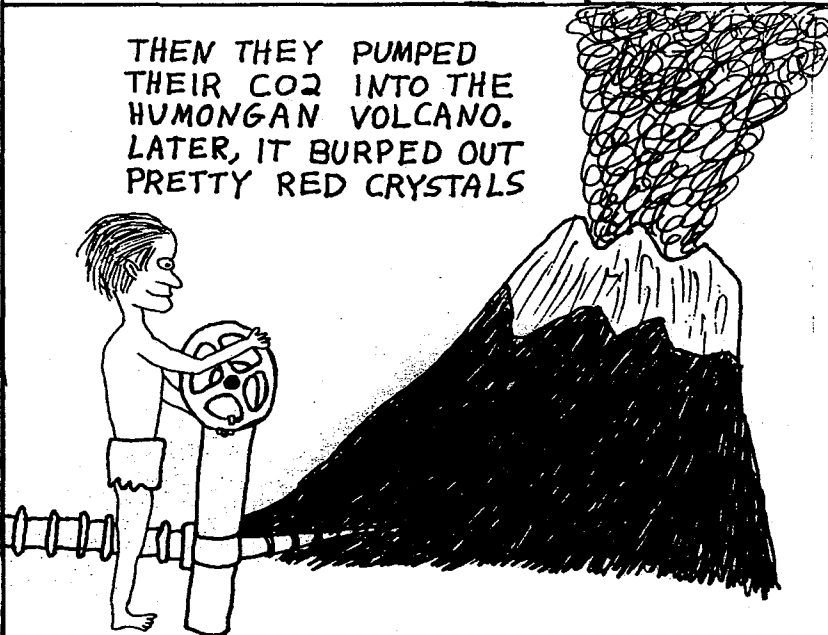


THEY PUT CO2 IN WABOOJI JUICE, AND MADE WABOOJAMPAGNE

THEN THEY MADE PYTHON BALLOONS

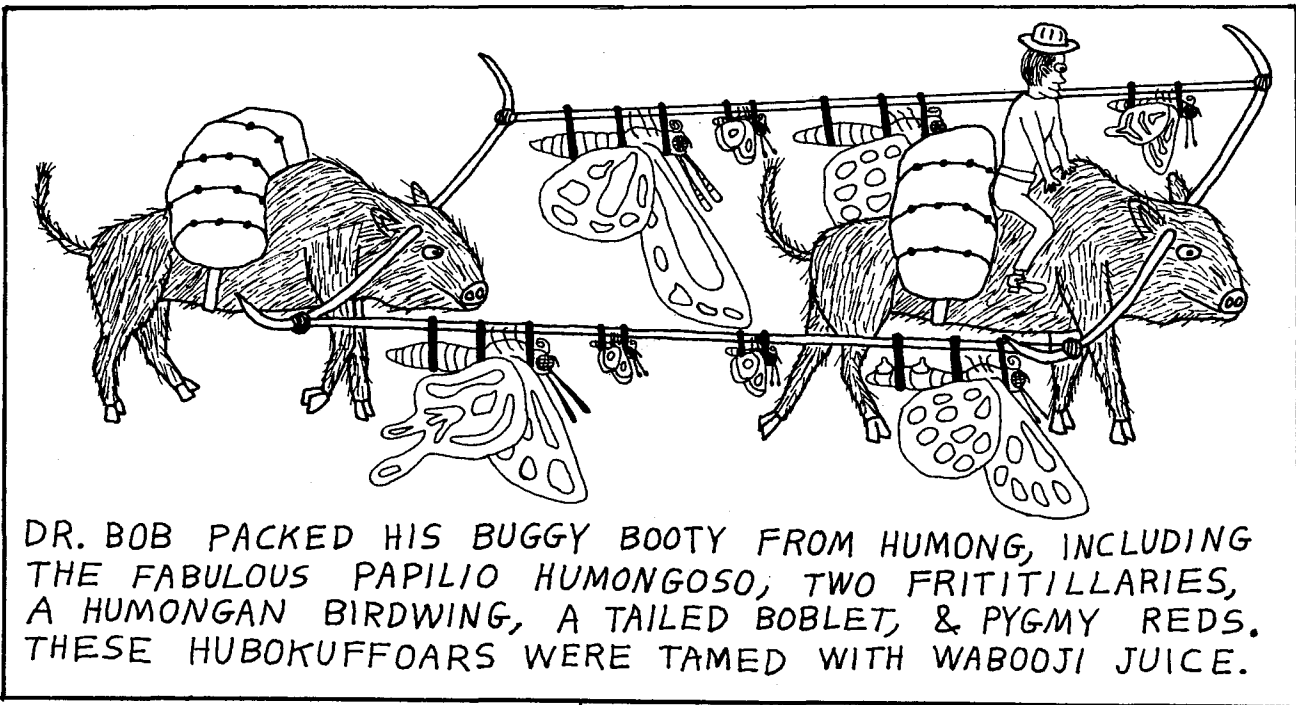


THEN THEY PUMPED THEIR CO2 INTO THE HUMONGAN VOLCANO. LATER, IT BURPED OUT PRETTY RED CRYSTALS

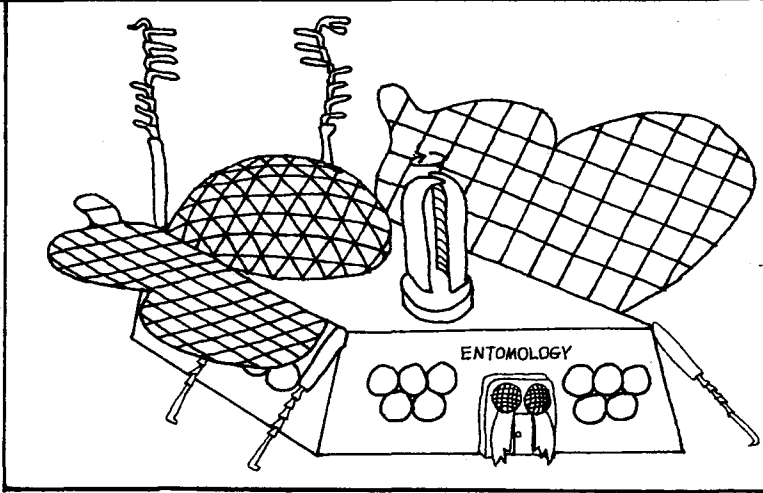
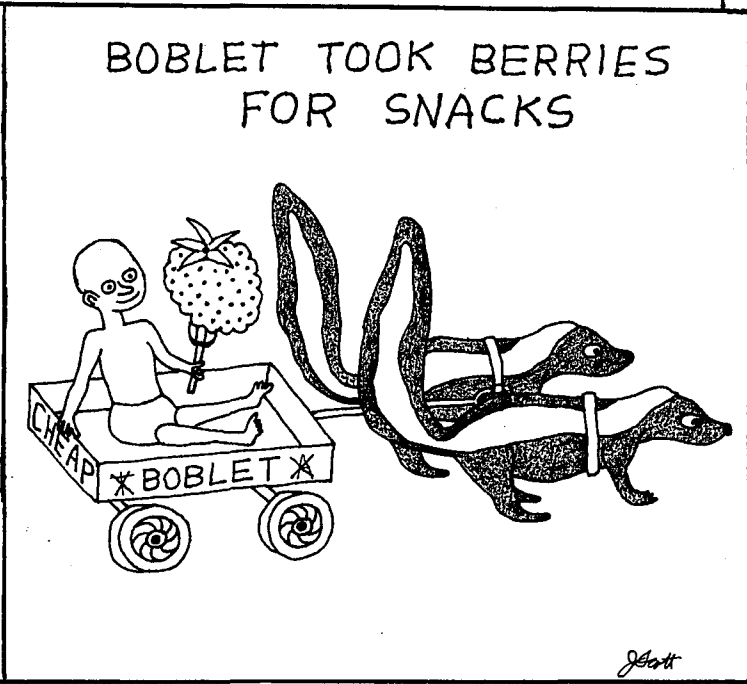
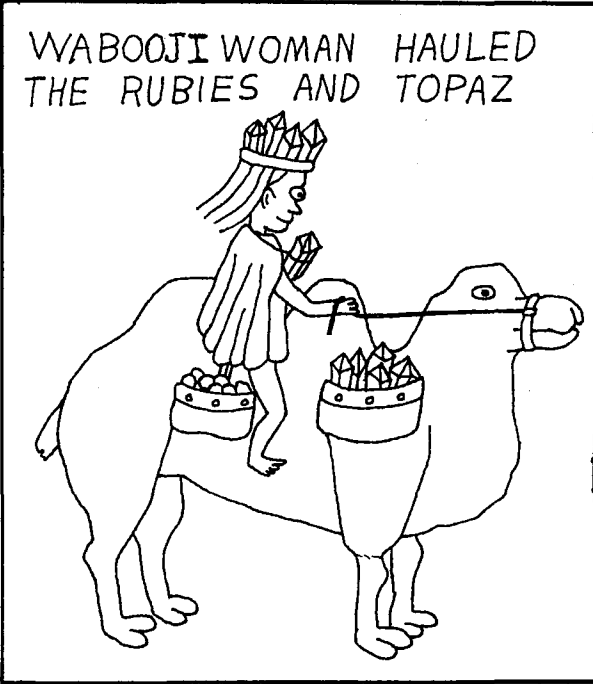


WOW! RUBIES?





DR. BOB PACKED HIS BUGGY BOOTY FROM HUMONG, INCLUDING THE FABULOUS PAPILIO HUMONGOSO, TWO FRITITILLARIES, A HUMONGAN BIRDWING, A TAILED BOBLET, & PYGMY REDS. THESE HUBOKUFFOARS WERE TAMED WITH WABOOJI JUICE.



DR. BOB NOW HAD ENOUGH MONEY FROM WABOOJAMPAGNE AND GEMS AND HDTV AND WIND-VANE POWER, TO PUT SOLAR CELL WINGS ON HIS BUG MUSEUM, & PLAN A THIRD EXPEDITION TO THE LAND OF HUMONG. GLORY BE, THOUGHT DR. BOB.