abundant in some specimens. An investigation of how sexual differences in diet influence parasite load is nearing completion.

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Molecular Systematics of Emydine Turtles. Linnaeus Fund Research Report

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The emydid turtles (Emydinae: genera *Clemmys*, *Emydoidea*, *Emys*, and *Terrapene*) are among the most familiar and well-studied chelonians in the world. This small turtle subfamily contains only ten species, yet exhibits greater ecological and morphological diversity than its more speciose sister group, the Deirochelyinae. Some species are fully aquatic (e.g., *Clemmys marmorata*) while others are almost entirely terrestrial (e.g., *Terrapene ornata*). In addition, species in the genera *Emydoidea*, *Emys*, and *Terrapene* possess shells with a movable plastron (plastral kinesis) while members of the genus *Clemmys* lack this trait.

Although emydines are extensively studied, popular, and of recent conservation concern, they lack a robust phylogeny. Morphological treatments of the Emydinae (Bramble, 1974; Gaffney and Meylan, 1988) hypothesized that the box turtles and other hinged genera form a monophyletic group (Fig. 1A). By default, the species without plastral kinesis were lumped into the genus *Clemmys*. Mitochondrial sequence data from the 16S ribosomal gene (Bickham et al., 1996) suggested that the genus *Clemmys* is not monophyletic (Fig. 1B). An attempt to combine these data with ecological, behavioral, biochemical, and additional morphological characters did not fully resolve the conflict between the morphological and molecular phylogenies (Burke et al., 1996). Despite the fact that some consensus has emerged from these studies, most hypothesized arrangements could be clarified and strengthened with additional molecular data.

Our objective was to shed light on the evolutionary history of emydines using all ten extant species, suitable sister taxa, and appropriately evolving molecular markers.

Materials and Methods. — We obtained liver tissue from museum specimens and blood samples from living zoo specimens for all 10 extant emydid species and 2 deirochelyine outgroup species (Appendix 1). We isolated genomic DNA from liver tissue and blood samples by standard proteinase K digestion and phenol/chloroform purification (Maniatis et al., 1982). We amplified a 1200 bp region of the mitochondrial genome encoding the entire cytochrome b gene and part of the adjacent transfer ribonucleic acid, threonine (tRNA1hr) via polymerase chain reac-
tion (PCR; Saiki et al., 1988) using the primers GLUDGL (Palumbi et al., 1991) and M (Shaffer et al., 1997) (Table 1). We amplified an additional 900 bp region of mtDNA encoding a portion of the nicotinamide adenine dinucleotide dehydrogenase subunit four gene (ND4) and flanking tRNA histidine (tRNAhistor), serine (tRNAser), and leucine (tRNaleu), using the primers ND4 and Leu (Arevalo et al., 1994) (Table 1). We used the following thermal cycle parameters for 50 µl amplification reactions: 35 cycles of 1 min denature at 94°C, 1 min anneal at 50–52°C, and 2 min extension at 72°C. We purified PCR products using the Wizard Prep Mini Column Purification Kit (Promega, Inc.) and used purified template in 10 µl dideoxy chain-termination reactions (Sanger et al., 1977) using ABI Big Dye chemistry (Perkin-Elmer Applied Biosystems, Inc.) and the primers listed in Table 1. We ran cycle-sequenced products on a 4.8% Page Plus (Ameresco) acrylamide gel using an ABI 377 automated sequencer (Perkin-Elmer Applied Biosystems, Inc.). We sequenced all samples in both directions.

We aligned DNA sequences with the sequence analysis program Sequencher™ 3.0 (Gene Codes Corp.). We translated protein coding nucleotide sequences into amino acid sequences using MacClade 3.06 (Maddison and Maddison, 1992). We identified tRNA genes by manually reconstructing their secondary structures using the criteria of Kumazawa and Nishida (1993). We deposited all mitochondrial DNA sequences in GenBank (Appendix 1).

We used maximum parsimony (MP; Swofford et al., 1996) and maximum likelihood (ML; Felsenstein, 1981) phylogenetic methods to infer the evolutionary relationships of emydine species. We conducted all phylogenetic analyses in PAUP 4.0b4a* (Swofford, 1998). We combined the cyt b and ND4 data sets and analyzed them together on the basis of total evidence (Eernisse and Kluge, 1993). We polarized the phylogeny via outgroup comparison (Maddison et al., 1984) using the chicken turtle, Deirochelys reticularia, and the painted turtle, Chrysemys picta.

We executed MP analyses with the branch-and-bound search algorithm (Hendy and Penny, 1982) using unordered characters. To assess the robustness of individual nodes, we used the bootstrap resampling method (Felsenstein, 1985) by employing 1000 replicates of closest searches in PAUP*. Additionally, we calculated branch support (Bremer, 1994) for internal nodes using the program TreeRot 2 (Sorenson, 1999).

To determine the most appropriate model of DNA substitution for reconstructing emydine relationships under ML, we executed a hierarchical likelihood ratio test (LRT; Felsenstein, 1993; Goldman, 1993; Yang, 1996) in the program Modeltest 3.0 (Posada and Crandall, 1998). The
A) Single most parsimonious tree (L =1088; CI= 0.642; RI = 0.466). Numbers above the nodes indicate bootstrap support while those below the nodes represent decay indices. B. Maximum likelihood estimate of emydine phylogeny (LnL = -7798.4184; \( \alpha = 0.2766 \)). Branch lengths are drawn proportional to the maximum likelihood estimates of genetic divergence.

Figure 2. Phylogenetic trees for emydine and outgroup mtDNA lineages. A. Single most parsimonious tree (L =1088; CI= 0.642; RI = 0.466). Numbers above the nodes indicate bootstrap support while those below the nodes represent decay indices. B. Maximum likelihood estimate of emydine phylogeny (LnL = -7798.4184; \( \alpha = 0.2766 \)). Branch lengths are drawn proportional to the maximum likelihood estimates of genetic divergence.

model of DNA evolution that best fit our sequence data was the general time reversible model (GTR; Rodriguez et al., 1990) of nucleotide substitution in conjunction with gamma (\( \Gamma; \) Yang, 1994a,b). The GTR + \( \Gamma \) model accommodates unequal base composition by using the empirical base frequencies, estimates the uneven ratio of each type of nucleotide substitution, and accounts for the heterogeneous rates of nucleotide substitutions across all sites.

Results. — Of the 2092 aligned base pairs, 609 were variable and 339 were parsimony informative. Among the ingroup taxa, 461 base pairs were variable and 251 were parsimony informative.

The branch-and-bound, equally weighted MP analyses produced a single most parsimonious tree (Fig. 2A), 1088 steps in length (CI = 0.642; RI = 0.466). The ML GTR + \( \Gamma \) reconstruction also yielded one tree (LnL = -7798.4184; \( \alpha = 0.2766 \)) nearly identical to the most parsimonious tree (Fig. 2B). In all analyses the emyridine turtles group to the exclusion of the deirochelyines (100% bootstrap; 37 decay index) and phylogenetic relationships were well resolved and well supported for most nodes of the tree.

The four North American box turtles (genus Terrapene) form a monophyletic group (100% bootstrap; 16 decay index) in which the two western species, T. ornata and T. nelsoni, form one clade (99% bootstrap; 10 decay index) and the aquatic T. coahuila and widespread T. carolina form another (94% bootstrap; 5 decay index). The genus Clemmys is not monophyletic. Instead, C. marmorata belongs to a clade containing Emydoidae blandingii and Emys orbicularis (98% bootstrap; 10 decay index). However, the relationships among these three taxa are not well resolved, as indicated by the conflict between the MP and ML reconstructions; the MP tree connects Emys orbicularis to C. marmorata (56% bootstrap; 1 decay index) while the ML tree links Emys orbicularis to Emydoidae blandingii. Additionally, the spotted turtle (C. guttata) does not group with the other eastern US Clemmys, C. insculpta and C. muhlenbergii. Instead, C. guttata gains some support as the sister taxon to the box turtles (70% bootstrap; 3 decay index) in both the MP and ML reconstructions. Finally, Clemmys insculpta and C. muhlenbergii form a robust monophyletic group (100% bootstrap; 10 decay index). Both MP and ML phylogenetic methods suggest that C. insculpta and C. muhlenbergii are the sister clade to all other emydines. This phylogenetic hypothesis, however, is poorly supported (63% bootstrap; 2 decay index).

In summary, the Emydinae can be divided into four well-supported clades: 1) Terrapene; 2) Clemmys guttata; 3) C. insculpta and C. muhlenbergii and; 4) C. marmorata, Emys orbicularis, and Emydoidae blandingii. Unfortunately, relationships between these emyridine clades remain enigmatic. Both MP and ML phylogenetic analyses yield the same topology, placing C. insculpta and C. muhlenbergii as the sister group to a monophyletic clade containing the rest of the emyridine turtles.

Discussion. — Our molecular phylogeny is both congruent and incongruent with previous estimates of emyridine relationships (Gaffney and Meylan, 1988; Bickham et al., 1996; Burke et al., 1996). Importantly, the large number of informative characters (339) in our multi-gene data set
allows us to address various hypotheses of emydid taxonomy and evolution. We discuss these taxonomic and evolutionary questions at length elsewhere (Feldman and Parham, in press), but briefly highlight two important points of our proposed phylogeny here: the paraphyly of *Clemmys*; and the paraphyly of hinged emydines.

Our mtDNA data explicitly show that the genus *Clemmys* is paraphyletic (Fig. 2). We propose that the spotted turtle, *C. guttata*, is the closest living relative to the North American box turtles. Our data also suggest that *C. marmorata* is not closely related to other *Clemmys*, but shares a more recent common ancestor with *Emys orbicularis* and *Emydoidea blandingi* (lastly, our phylogeny indicates that *C. muhlenbergii* and *C. insculpta* form a monophyletic group exclusive of, and sister to, all other emydine turtles.

A paraphyletic *Clemmys* stands in contrast to both the accepted taxonomy of the Emydinae (Collins, 1997; Crotzer, 2000) and the morphological phylogeny of the group (Gaffney and Meylan, 1988; Fig. 1A). A non-monophyletic *Clemmys*, however, is not an entirely original concept. Previous molecular (Bickham et al., 1996; Fig. 1B) and combined treatments (Burke et al., 1996) of the Emydinae have suggested a paraphyletic *Clemmys*. Thus, we propose a new taxonomy for the Emydinae in a more thorough summary (Feldman and Parham, in press). Our taxonomy is consistent with the Linnaean system of ranks as well as the informative scheme of phylogenetic taxonomy (de Queiroz and Gauthier, 1992).

The most notable result of our mtDNA phylogeny is the paraphyly of the hinged emydines. Emydine shell kinesis involves several morphological specializations: 1) an alignment of scales with plastral sutures and a reduction of sutural connections to form a hinge; 2) segmented scapulae that facilitate head and limb retraction; and 3) a closing mechanism modified from cervical musculature (Bramble, 1974). This particular combination of traits is unique among living chelonians and is thought to have evolved only once (Bramble, 1974; Gaffney and Meylan, 1988). However, our data suggest that shell kinesis evolved either twice (once in *Terrapene* and once in the *C. marmorata* + *Emys* + *Emydoidea* clade) or evolved once and was lost twice (in *C. marmorata* and *C. guttata*). Using information from the fossil record and data on the independent derivation of plastral kinesis in other living turtles (e.g., various batagurid genera), we hypothesize that plastral hinging evolved twice in parallel in the Emydinae (Feldman and Parham, in press).

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### *Clemmys guttata* MVZ 175961, AF258885, AF258870; *Clemmys insculpta* ROM 1523, AF258864, AF258876; *Clemmys muhlenbergii* zoo specimen, Wildlife Conservation Society, Bronx Zoo, New York, New York, AF258863, AF258875; *Emydoidea blandingi* ROM 20922, AF258857, AF258869; *Clemmys marmorata* MVZ 164994, AF258855, AF258867; *Emys orbicularis* CAS 182905, AF258856, AF258868; *Terrapene carolina* MVZ 137441, AF258859, AF258871; *Terrapene coahuila* zoo specimen (T00228), Gladys Porter Zoo, Brownsville, Texas, AF258860, AF258872; *Terrapene nelsoni* zoo specimen, Arizona Sonoran Desert Museum, Tucson, Arizona, AF258861, AF258873; *Terrapene ornata* MVZ 137743, AF258862, AF258874; *Chrysemys picta* MVZ 230532, AF258866, AF258878; *Deirochelys reticularia* MVZ 209023, AF258865, AF258877.

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The striped mud turtle, Kinosternon baurii, is a small, aquatic turtle that ranges from the lower Florida Keys north to Virginia (Iverson, 1992). Throughout most of their range, striped mud turtles are found in freshwater habitats, however, in the lower Florida Keys they can be found in both freshwater and brackish water habitats (Dunson, 1981). The lower Florida Keys populations are listed as endangered by the state of Florida and may warrant special attention.

A potential exists for restricted gene flow between populations in the lower Florida Keys and those in the remainder of the species’ range because they are separated by a seven-mile expanse of open ocean. Historically, the lower Florida Keys populations were considered a separate subspecies (K. baurii baurii) distinct from the mainland K. baurii palmatum (Stejneger, 1925). Both morphological and physiological differences have been proposed as supporting this subspecific designation (Uzzell and Schwartz, 1955; Dunson, 1981). Although no subspecies currently are recognized, taxonomic controversy continues regarding the genetic subdivision among striped mud turtle populations (see Dunson, 1981; Lazell, 1989).

The objective of this study was to assess the degree of genetic isolation that exists between mainland, upper, and lower Florida Keys populations of the striped mud turtle. Mitochondrial control region DNA sequence data were examined to determine levels and patterns of variation in 36 Kinosternon baurii individuals from 10 different geographic locations from throughout the species’ range.

Methods. — Each captured turtle was marked and measured, and a small blood sample was withdrawn before the turtle was released at its site of capture. Sampling locations are illustrated in Fig. 1 and are: Stock Island, Dade Co., Florida (n = 7), Summerland Key, Dade Co., Florida (3), Big Pine Key, Dade Co., Florida (2), Grassy Key, Dade Co., Florida (3), Snapper Creek Canal, Miami, Dade Co., Florida (3), Nine Mile Pond and Royal Palm Hammock, Everglades National Park, Dade Co., Florida