PROC. BIOL. SOC. WASH. 95(3), 1982, pp. 423-427

SYSTEMATIC RESOLUTION OF THE GENERA OF THE *CRINIA* COMPLEX (AMPHIBIA: ANURA: MYOBATRACHIDAE)

W. Ronald Heyer, Charles H. Daugherty, and Linda R. Maxson

Abstract.—Re-evaluation of morphological analyses and generic taxonomy of Australian myobatrachine frogs in the *Crinia* complex is undertaken in light of new genetic data. All species currently in the genera *Ranidella* and *Crinia* are assigned to *Crinia*. The genera *Paracrinia* and *Geocrinia* are retained.

Four Australian genera, *Ranidella, Crinia, Paracrinia*, and *Geocrinia* (Myobatrachidae: Myobatrachinae), collectively designated the "*Crinia* complex," comprise a group of small, rather nondescript frogs (Heyer & Liem 1976). The species of this complex have been partitioned into genera by recent authors with differing results. All recent authors agree concerning the smallest groups of species clusters in this complex, but considerable disagreement exists regarding the assignment of these groups at the generic level. Currently hypothesized groups and classifications are summarized in Table 1.

The following morphological and behavioral characters differentiate the species clusters: vomer and vomerine teeth, omohyoideus muscle, outer metatarsal tubercle, belly texture, egg placement, and mating call. However, clustering algorithms provide no unequivocal pattern of relationships among these species: "There is no way to group the . . . taxa so that two of the derived states of these characters define the same assemblage. Rather, a grouping which results in a cluster having all the taxa with the same derived state of one character leads to convergence of states in the other characters" (Heyer and Liem 1976:9). Thus, the data presented by Heyer and Liem (1976) are certainly open to alternate clustering interpretations than theirs. The phenetic analyses of Blake (1973) and Thompson (1981) recognize the same basic species groups, but the clustering pattern of the groups was highly variable dependent on data scoring and the algorithm used. Within the Crinia complex, morphological variation is so limited that it has been impossible to achieve a stable clustering scheme and, hence, taxonomic consensus. For these cases where the nature of the morphological data preclude a definitive analysis of relationships, use of a different data base is required for analysis of relationships.

Daugherty and Maxson (in press) recently estimated genetic relationships among species of the *Crinia* complex based on MC'F (micro-complement fixation) data from the serum protein albumin. These genetic data, in concert with the morphological data, provide a new basis for determining evolutionary lineages within the complex. The major lineages are herein proposed as generic units in order to provide a stable classification for this complex.

The Genetic Data

Immunological distances derived from comparisons of serum albumins provide both cladistic information and a time framework for interpreting evolutionary

	E Contraction of the second			
	Blake, 1973	Heyer & Liem, 1976	Thompson, 1981	This study
georgiana	Together with haswelli, Cri- nia	Crinia	Not studied	Crinia
haswelli	Together with georgiana, Cri- nia	Paracrinia	Not studied	Paracrinia
laevis cluster	Geocrinia	Geocrinia	Not studied	Geocrinia
signifera cluster	Species group of <i>Ranidella</i>	Ranidella	Distinct species group of Ran- idella	Crinia
riparia	Together with tasmaniensis, second species group of Rani- della	Together with tasmaniensis, Australocrinia	Second distinct species group of <i>Ranidella</i>	Crinia
tasmaniensis	Together with <i>ri- paria</i> , second species group of <i>Ranidella</i>	Together with ri- paria, Austral- ocrinia	Third distinct species group of <i>Ranidella</i>	Crinia

Table 1.—Species clusters and generic assignment of the Crinia complex.

relationships. In the past decade, such protein data have been used extensively in phylogenetic studies of diverse amphibian taxa (e.g., Heyer and Maxson 1982; Maxson 1981). Albumin immunological distances (ID) have been shown to estimate sequence differences in albumins between species (Maxson and Wilson 1974) and to accumulate measurable sequence differences at an approximate rate of one substitution per lineage per million years (Wilson *et al.* 1977). Daugherty and Maxson (in press) have measured a series of immunological distances among the albumins of many members of the *Crinia* complex. The data consist of oneway comparisons to *signifera*, currently assigned to the genus *Ranidella* (Table 2). The pattern of divergence from *signifera* reveals the major genetic lineages within this complex.

Members of the *signifera* cluster (Table 2) exhibit ID values ranging from 24 to 40. The distance to *riparia* is 15 units and to *tasmaniensis* is 53 units. Clearly, *riparia* belongs to the same genetic lineage as other members of the *signifera* cluster. The ID value for *tasmaniensis* is somewhat higher than values measured to members of the *signifera* group, but not as large as values to other lineages (*haswelli* and *laevis*; see below) within the *Crinia* complex. Furthermore, an ID value of around 50 is often seen between species within other frog genera (e.g., Maxson and Wilson 1975; Heyer and Maxson 1981). The immunological evidence thus supports Thompson's (1981) proposal that *Australocrinia* (i.e., *riparia* and *tasmaniensis*) be synonymized with *Ranidella* (i.e., the *signifera* cluster).

The ID to georgiana is 29 units, suggesting that georgiana is part of the same genetic lineage as the frogs in the signifera cluster. For both georgiana and riparia, the ID values to signifera are smaller than most ID values measured between signifera and other members of the signifera group. The taxonomic

2111. UDAREALDINA DI	Species compared	ID to signifera	Thursday.
signifera cluster:			
	signifera	0	
	glauerti	24	
	parinsignifera	24	
	bilingua	30	
	sp. nov.	31	
	remota	31	
	deserticola	40	
	riparia	15	
	tasmaniensis	53	
	georgiana	29	
	victoriana	133	
	haswelli	140	

Table 2.—Albumin immunological distances between *signifera* and other species of the *Crinia* complex.*

* Data from Daugherty and Maxson (in press).

conclusions are that *Ranidella* (including *Australocrinia*) and *Crinia* are congeneric and that *Ranidella* is a synonym of *Crinia*.

On the other hand, the ID value of *signifera* to *victoriana*, the only member of the *laevis* group tested to date, is 133. This very large value is concordant with recognition of the genus *Geocrinia*. The ID of *signifera* to *haswelli* is similarly high, 140. The similar, but high, ID values of *haswelli* and *victoriana* to *signifera* indicate a distant relationship of *haswelli* and *victoriana* to *signifera*. The values do not indicate what the relationship of *haswelli* is to *victoriana*; it could be close or distant. In fact, preliminary data (Maxson and Daugherty, unpublished) indicate a distant relationship between those taxa (ID value between 90 and 100). These data are consistent with recognition of the genera *Geocrinia* (including *victoriana*) and *Paracrinia* (including *haswelli*).

Discussion

Several conclusions regarding evolution of the *Crinia* complex logically follow from recognition of the genera *Crinia*, *Geocrinia*, and *Paracrinia*.

Several of the character states that differentiate among the species clusters have apparently evolved independently several times. Loss of the vomer and vomerine teeth has occurred within both *Crinia* and *Geocrinia*. Both smooth and granular bellied frogs occur in the genus *Crinia*. All major variation in mating call occurs within the genus *Crinia*. Perhaps most notable is that variations in life history occur within, rather than among lineages. The change from a lotic to lentic egg placement and larval morphology has taken place entirely within the genus *Crinia*, and these life history differences cannot be used to define generic units. A similar situation occurs within the genus *Geocrinia*, with evolution of terrestrial larvae from pond larvae.

The morphological characters that differentiate among the genetically defined lineages are, for the most part, characters involving reduction or loss as the derived state. *Crinia* species have an outer metatarsal tubercle; *Geocrinia* species lack the tubercle. *Crinia* species have an omohyoideus muscle; *Paracrinia* lacks the muscle. *Geocrinia* species lack an outer metatarsal tubercle and toe fringing; *Paracrinia* has a metatarsal tubercle and toe fringing. Of these characters, the only one that does not involve reduction or loss as the derived state is the metatarsal tubercle (Heyer and Liem 1976).

Derived states involving reduction or loss are usually considered to contain little or no phyletic information (Hecht and Edwards 1976). That morphological states of loss or reduction are the primary criteria permitting morphological discrimination of the genera within the *Crinia* complex (which the genetic data show to be greatly differentiated) suggests that the use in systematics of characters of loss and reduction needs re-evaluation. It also further documents the extreme morphological conservatism so often observed in anuran evolution (Maxson and Wilson 1975; Wilson *et al.*, 1977) and reinforces the need to examine genetic and other categories of characters when conducting phylogenetic studies (Blake 1973).

Generic Redefinition

Crinia is the only genus requiring redefinition. The format used is comparable to that of Heyer and Liem (1976), which may be referred to for definitions of the genera *Geocrinia* and *Paracrinia*.

Crinia Tschudi, 1838

Synonyms.—Ranidella Girard, 1853 Camariolus Peters, 1863 Pterophrynus Lutken, 1863 Pterophryne Gunther, 1867 Australocrinia Heyer and Liem, 1976. Type species.—Crinia georgiana Tschudi, 1838.

Diagnosis.—A myobatrachine genus; cervical cotyles widely separated; vomer and vomerine teeth present or absent; narrow sacral diapophyses; depressor mandibulae muscle with or lacking a slip from the dorsal fascia; omohyoideus muscle present; tympanum present; belly smooth or granular; toes with or without fringe; outer metatarsal tubercle present; eggs placed in water or on land; pond or stream larvae.

Content.—Crinia bilingua, deserticola, georgiana, glauerti, insignifera, parinsignifera, pseudinsignifera, remota, riparia, signifera, perhaps sloanei (see Thompson 1981), subinsignifera, tasmaniensis, tinnula, and undescribed species. Present knowledge of variation within this genus (as here defined) does not support the recognition of species groups (also see Daugherty and Maxson, in press).

Acknowledgments

We thank P. Baverstock and his family for assistance in collecting sufficient *C. signifera* to produce an antibody. We thank the many other people who provided us with specimens. Portions of this work were carried out at the Institute of Medical and Veterinary Science in Adelaide, Australia. We thank M. Krieg, P. Baverstock, M. Adams, and C. Watts for stimulation and support while in Adelaide. Financial support came in part from NSF grant INT 79-24146 to LRM and NIH grant 1-F32FM06788-01 to CHD.

Literature Cited

Blake, A. J. D. 1973. Taxonomy and relationships of Myobatrachine frogs (Leptodactylidae): A numerical approach.—Australian Journal of Zoology 21:119–149.

- Hecht, M. K., and J. L. Edwards. 1976. The determination of parallel or monophyletic relationships: the proteid salamanders—a test case.—American Naturalist 110:653-677.
- Heyer, W. R., and D. S. Liem. 1976. Analysis of the intergeneric relationships of the Australian frog family Myobatrachidae.—Smithsonian Contributions to Zoology 233:1-29.
 - , and L. R. Maxson. 1982. Distributions, relationships, and zoogeography of lowland frogs: The Leptodactylus complex in South America, with special reference to Amazonia. Pages 375– 388 in G. T. Prance (ed.), Biological Diversification in the Tropics. Columbia University Press, New York. 714 pp.
- Maxson, L. R. 1981. Albumin evolution and its phylogenetic implications in African toads of the genus *Bufo*.—Herpetologica 37:96–104.
 - ——, and A. C. Wilson. 1974. Convergent morphological evolution detected by studying proteins of tree frogs in the *Hyla eximia* species group.—Science 185:66–68.
- —, and —, 1975. Albumin evolution and organismal evolution in tree frogs (Hylidae).— Systematic Zoology 24:1–15.
- Thompson, M. B. 1981. The systematic status of the genus *Australocrinia* Heyer and Liem (Anura: Leptodactylidae).—Australian Journal of Zoology 29:93–102.
- Wilson, A. C., S. S. Carlson, and T. J. White. 1977. Biochemical evolution.—Annual Review of Biochemistry 46:573–639.

(WRH) Department of Vertebrate Zoology, Smithsonian Institution, Washington, D.C. 20560. (CHD & LRM) Department of Genetics and Development, University of Illinois at Urbana-Champaign, Urbana, Illinois 61801.

Daugherty, C. H., and L. R. Maxson. [In press.] A biochemical assessment of the evolution of myobatrachine frogs.—Herpetologica.



Heyer, W R, Daugherty, Charles H., and Maxson, L R. 1982. "Systematic Resolution Of The Genera Of The Crinia Complex Amphibia Anura Myobatrachidae." *Proceedings of the Biological Society of Washington* 95, 423–427.

View This Item Online: <u>https://www.biodiversitylibrary.org/item/108716</u> Permalink: <u>https://www.biodiversitylibrary.org/partpdf/48280</u>

Holding Institution Smithsonian Libraries and Archives

Sponsored by Biodiversity Heritage Library

Copyright & Reuse

Copyright Status: In copyright. Digitized with the permission of the rights holder. Rights Holder: Biological Society of Washington License: <u>http://creativecommons.org/licenses/by-nc-sa/3.0/</u> Rights: <u>https://biodiversitylibrary.org/permissions</u>

This document was created from content at the **Biodiversity Heritage Library**, the world's largest open access digital library for biodiversity literature and archives. Visit BHL at https://www.biodiversitylibrary.org.