TAXONOMY AND GENETIC RELATIONSHIPS OF PANGASIIDAE, ASIAN CATFISHES, BASED ON MORPHOLOGICAL AND MOLECULAR ANALYSES

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ABSTRACT

Pangasiids are economically important riverine catfishes generally residing in freshwater from the Indian subcontinent to the Indonesian Archipelago. The systematics of this family are still poorly known. Consequently, lack of such basic information impedes the understanding of the biology of the Pangasiids and the study of their aquaculture potential as well as improvement of seed production and growth performance. The objectives of the present study are to clarify phylogeny of this family based on a biometric analysis and molecular evidence using 12S ribosomal mtDNA on the total of 1070 specimens. The study revealed that 28 species are recognised as valid in Pangasiidae. Four genera are also recognized as Helicophagus Bleeker 1858, Pangasianodon Chevey 1930, Pteropangasius Fowler 1937, and Pangasius Valenciennes 1840 instead of two as reported by previous workers. The phylogenetic analysis demonstrated the recognised genera, and genetic relationships among taxa. Overall, trees from the different analyses show similar topologies and confirm the hypothesis derived from geological history, palaeontology, and similar models in other taxa of fishes from the same area. The oldest genus may already have existed when the Asian mainland was still connected to the islands in the southern part about 20 million years ago.

KEYWORDS: taxonomy, genetic, morphology, molecular, catfish, pangasiidae

INTRODUCTION

Pangasiidae are economically important riverine catfishes generally occurring in freshwater from the Indian subcontinent to the Indonesian Archipelago (Gustiano 2003; Gustiano and Pouyaud 2005; 2006). Morphologically, they are recognized by a laterally compressed body, two pairs of barbels, a short dorsal fin with two spines, a well developed adipose fin, a long anal fin, and a strong pectoral spine (Teugels 1996). The systematics of this family are still poorly known (Gustiano 2003; Pouyaud et al. 2005). Consequently, the lack of this basic information is a significant barrier to understanding the biology and hence the study of the aquaculture potential of Pangasiids, the improvement of their seed production and growth performance (Legendre

The objectives of the present study are to clarify phylogeny of this family based on a biometric analysis and molecular evidence using 12S ribosomal mtDNA on the total of 1070 specimens.

MATERIALS AND METHODS

Biometrics

Nine hundred and ninety nine specimens were examined including the type of 49 previous described species housed in various museums. For each specimen, 35 point-to-point measurements covering the possible variations of the body were undertaken (Fig. 1).

Data were subjected to principal component analysis (PCA) (Bookstein *et al.* 1985). Measurements were log-transformed in order to minimise the effect of non-normality. The

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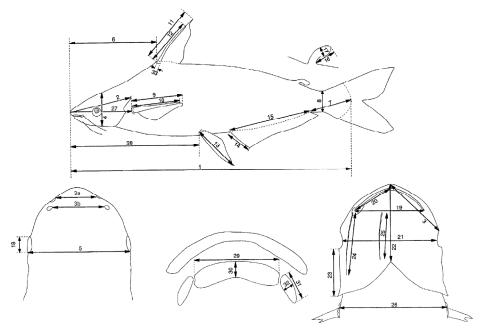


Figure 1. Measurements taken on *Pangasius* specimens: 1. Standard length; 2. Head length; 3. Snout length; 3a. Anterior snout width; 3b. Posterior snout width; 4. Head depth; 5. Head width; 6. Predorsal length; 7. Caudal peduncle length; 8. Caudal peduncle depth; 9. Pectoral fin length; 10. Pectoral spine length; 11. Dorsal fin length; 12. Dorsal spine width; 13. Pelvic fin length; 14. Anal fin height; 15. Anal fin length; 16. Adipose fin height; 17. Adipose fin width; 18. Eye diameter; 19. Mouth width; 20. Lower jaw length; 21. Interorbital length; 22. Distance snout to isthmush; 23. Postocular length; 24. Maxillary barbel length; 25. Mandibulary barbel length; 26. Body width; 27. Prepectoral length; 28. Prepelvic length; 29. Vomerine width; 30. Vomerine width; 31. Palatine length; 32. Palatine width; 33. dorsal spine width.

first factor, considered as the size factor, was not taken into account in order to minimise the size differences between samples. An independent PCA was run on the correlation matrix from untransformed data.

Finally, data analysis consisted of characterising groups from scatter plots between pairs of structuring characters for subsequent use in generic identification keys.

Molecular analysis

Fresh tissue samples were stored in liquid nitrogen for transfer to the laboratory. They were then stored at -20°C. Mitochondrial analysis consisted of sequencing part of the 12S rDNA gene. One to five individuals were analysed for 28 species of Pangasiidae. One hundred mg of muscle were used to isolate total genomic DNA using a hexadecylmethyl-

ammoniumbromide extraction process (Doyle & Doyle, 1987). Primers used for the amplification of the mitochondrial gene included 12S light strand 5'-TTACACATGCAAGTCTCCGC-3' and 12S heavy strand GTTACGACTTGCC TCCCCTT-3' defined on the complete mitochondrial sequence of *Cyprinus carpio* (EMBL X61010, Chang *et al.* 1994).

DNA amplification and sequencing were performed following the procedures developed in Pouyaud *et al.* (2000). In order to check sequence accuracy and to correct any ambiguity bases, both strands were sequenced using each one of the two initial PCR primers.

Sequences of both strands were compared with each other and aligned using the sequence editor ESEE (version 3.1 s; Cabot & Beckenbach, 1989). Comparison of absolute numbers of transitions and tranversions were

calculated and plotted versus corrected genetic distances (using Kimura's distance two-parameter method, Kimura 1980). Phylogenetic inference was based on the neighbour joining method (Saitou & Nei 1987) (NEIGHBOR program in PHYLIP; Felsenstein 1993) from Kimura's distances among species. The reliability of the topologies was assessed with bootstrapping on 1000 replicates (SEQBOOT and CONSENSE programs in PHYLIP; Felsenstein 1993).

RESULTS AND DISCUSSION

During the study twenty eight species were recognised as valid in Pangasiidae. Four genera were also recognized (*Helicophagus* Bleeker, 1858; *Pangasianodon* Chevey, 1930; *Pteropangasius* Fowler, 1937; and *Pangasius* Valenciennes, 1840) instead of two as reported by Vidthayanon (1993). Key below different genera is presented. For each genus, a key to the species is given.

Key to genera

- 1b. 6 pelvic fin rays2
- 2a. Slender anterior part of snout (<16.5% HL), posterior nostrils are in between anterior nostrils and orbit................ *Helicophagus*

Key to species of Helicophagus

- Anal ray counts more than 35; premaxillary teeth divided into two quadratic

Key to species Pangasianodon

- 2b. Head length less than 29% SL; prepectoral length less than 25% SL; anterior part of snout width less than 30% HL; anal fin length more than 27.5% SL; distance between snout and isthmus more than 37% HL Pangasianodon hypophthalmus

Key to species of Pteropangasius

Key to species of Pangasius

- 1a. Vomerine toothplate without additional toothplate2
- 1b. Vomerine toothplate with additional toothplate5

- 3a. Anal fin length more than 31% SL and

	prepelvic length less than 44% SL	14a.Predorsal length 31.8-35.3% SL, snout length 45.5-52.4% HL, anterior part of
3b.	Anal fin length less than 31% SL and prepelvic length 42-52.9% SL4	snout width 32.6-37.8% HL, posterior part of snout width 42.2-49.1% HL
4a.	Dorsal spine width more than 7.7-9.3% HL and head width more than 14.1-15.6% SL	14b. Predorsal length 34.6-42.8% SL, snout length 31.9-49.9% HL, anterior part of snout
4b.	Dorsal spine width 5.5-7.6% HL and head width 13.8-16.4% SL <i>P. nieuwenhuisii</i>	width 22.8-45.8% HL, and posterior part of snout width 26.8-39.5% HL15
5a.	Filamentous fin rays6	15a. Dorsal spine width 4.7-6.2% HL, head length 19.6-23.2% SL, head width 11-14.2%
5b.	Non filamentous fin rays7	SL, and body width 14.9-17% SL
6a.	Body width 21.1-23.2% SL, head length	P. rheophilus
/ h	23.2-30.2% SL, gill rakers on the first branchial arch 17-26	15b.Dorsal spine width 5.4-10.4% HL, head length 21.3-28.8% SL, head width 11.9-20.6% SL, body width 16.5-21.4% SL16
6b.	Body width 16.9-21.9% SL, head length 20.3-28.3% SL, gill rakers on the first branchial arch 13-17	16a. Higher gill rakers number on the first bran- chial arch (more than 27)17
7a.	Maxillary barbel length 100.5-203.9% HL, mandibular barbel 76.8-176.5% HL, and eye	16b. Lower gill raker number on the first bran- chial arch (less than 27)18
	diameter 21.9-45% HL	17a. 35-47 gill rakers on the first branchial arch,
7b.	Maxillary barbel less than 100.5% HL and mandibular less than 76.8% HL8	anterior part of snout width 26.4-31.1% HL
8a.	Predorsal length 25.1-31.2% SL and eye diameter 16.0-30.3% HL	17b.27-39 gill rakers on the first branchial arch, anterior part of snout width 29.3-36.6.5% HL
8b.	Predorsal length more than 30.1% SL9	18a. Predorsal length 33.5-37% SL and head
9a.	Long caudal peduncle (17.6-21.4% SL) and	length 21-24% SL19
	eye diameter 19.2-25.5% HL	18b. Predorsal length 36.1-42.8% SL and head length 23.8-28.8% SL20
9b.	Short caudal peduncle (less than 17% SL)10	19a. Dorsal spine width 8.1-8.9% HL and post ocular length 28.6-33.3% HL
10a.	Eye diameter 22.8-29.4% HL and predorsal	P. myanmar
105	length 30.1-32.7% SL	19b. Dorsal spine width 6.3-7.7% HL and post ocular length 32.3-39.7% HL
100	Eye diameter less than 22.8% HL and predorsal length more than 31.8% SL11	20a.Width of mouth 31-41% HL, vomerine
11a.	Short distance snout isthmus (less than 110% SNL)12	toothplate width 16-21.5% HL, and lower jaw length 17-24% HL
11b	Long distance snout isthmus (more than 110% SNL)14	20b.Width of mouth 41.9-52.5% HL, vomerine
12a.	Maxillary barbel length 79-97%; mandibular length 56-66% HL and dorsal spine width 4.4-5.7% HL	toothplate width 21.9-30.7% HL, and lower jaw length 23.9-31.5% HL
12b	Maxillary barbel length less than 80.7%	The phylogenetic analysis demonstrated
	HL, mandibular barbel length less than 52% HL; dorsal spine width more than 6.5% HL	the recognised genera, and the genetic relationships among the species (Fig. 2). Overall, trees from the osteological (Gustiano 2003),
13a.	Gill rakers on the first branchial arch 16-	molecular and biometric analyses (Gustiano and Pouyaud 2005) show similar topologies and
12h	23	confirm the hypothesis derived from geological history, palaeontology, and similar models
130	32	in other taxa of fishes from the same area

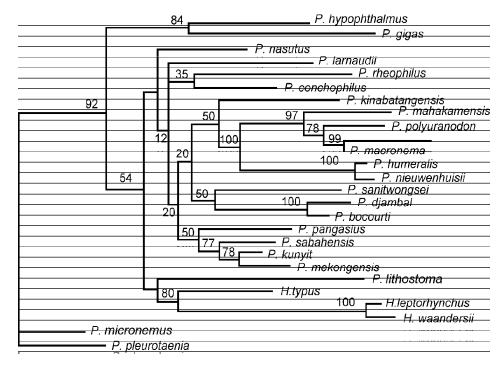


Figure 2. The nearest neighbor dendrogram for 12S rDNA gene within Pangasiidae. Numbers next to node show percentage group occurrence found with the 1000 bootstrapped trees.

(Gustiano 2003). The oldest genus may already have existed when the Asian mainland was still connected to the islands in the southern part about 20 million years ago (Gustiano *et al.* 2003; Pouyaud *et al.* 2005).

CONCLUSION

The analysis demonstrated the genetic relationships among species of Pangasiids. The present study recognizes four genera i.e. *Helicophagus* Bleeker, 1858; *Pangasianodon* Chevey, 1930; Pteropangasius Fowler, 1937; and *Pangasius* Valenciennes, 1840. At this level, species belonging to the same genus are aggregated using the degree of similarity of morphometric and molecular data.

AKNOWLEDGEMENTS

The authors thank J. Slembrouck, M. Legendre, A. Pariselle, Sudarto, A.H. Kristanto, W. Hadie, D. Sadili, O. Komarudin, and E. Siagian for their support and help in collecting fish samples. Sincere thanks to G. Teugels for his encouragement in the present study. This paper forms a part of INCO.DC Project

"Catfish Asia" financed by the European Union (Contract IC 18-CT 96-0043). The Senior Author is also grateful to AARD for the fellowship provided to complete this study through "PAATP" Project at Leuven University, Belgium.

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