Huangshan population of Chinese *Zacco platypus* (Teleostei, Cyprinidae) harbors diverse matrilines and high genetic diversity

Xin ZHENG^{1,2,#}, Tian-Qi ZHOU^{1,2,#}, Tao WAN^{3,#}, Anabel PERDICES⁴, Jin-Quan YANG¹, Xin-Sheng TANG², Zheng-Ping WANG⁵, Li-Qun HUANG⁶, Song HUANG^{2,3,*}, Shun-Ping HE^{7,*}

¹ Key Laboratory of Exploration and Utilization of Aquatic Genetic Resources, Shanghai Ocean University, Ministry of Education, Shanghai 201306, China

² College of Life and Environment Sciences, Huangshan University, Huangshan Anhui 245041, China

³ State Key Laboratory of Genetic Resources and Evolution, Kunming Institute of Zoology, Chinese Academy of Sciences, Kunming Yunnan 650223, China

⁴ Museo Nacional de Ciencias Naturales, CSIC, C/ José Gutiérrez Abascal, 2, 28006 Madrid, Spain

⁵ College of Foreign Languages, Huangshan University, Huangshan Anhui 245041, China

⁶ Landscaping Bureau of Huangshan Scenic Area, Huangshan Anhui 245000, China

⁷ Institute of Hydrobiology, Chinese Academy of Sciences, Wuhan Hubei 430072, China

ABSTRACT

Six main mitochondrial DNA (mtDNA) lineages have been described in minnow (Zacco platypus) samples obtained from northern, western and southern China. Perdices et al. (2004) predicted that further sampling of other tributaries might discover more lineages of this species. In this study, we collected 26 Zacco platypus individuals in the Huangshan area of eastern China and determined the cytochrome b (cytb) sequence variations. Combined with reported data in GenBank, we identified ten matrilines (Zacco A-J) in a total of 169 samples, with relatively high molecular divergence found among them. The Huangshan population had the greatest genetic variation among all sampled regions and hosted six of the ten matrilines. Our results highlight the significance of the Huangshan area for the conservation of Zacco platypus.

Keywords: *Zacco platypus*; Matriline; Huangshan; Phylogenetics; Diversity

INTRODUCTION

Zacco platypus is a common minnow that occurs in sympatry with most Chinese cyprinids (Deng et al., 2013). Topographical barriers may restrict its life history and drive cryptic diversity. The species' distribution encompasses all major river systems in mainland China, as well as the Korean Peninsula and Japan (Chen, 1998). Perdices et al. (2004) analyzed the genetic diversity of *Z. platypus* sampled in the upper and middle Changjiang (Yangtze River) and found four major matrilines that may harbor multiple species. Long-term interruption of dispersal is thought to have driven this diversity. Perdices and Coelho (2006) further studied samples from the Pearl River and northern drainages, and obtained six matrilines in China. Using nuclear DNA data, Berrebi et al. (2005) identified four genetic groups within *Z. platypus* from Sichuan, Hunan and Guangxi provinces in China.

Although Perdices et al. (2004) predicted that exhaustive sampling of other tributaries might discover other lineages of *Zacco*, few specimens have been sampled in eastern China. The Huangshan area in eastern China is a mosaic of mountains with elevations lower than 2 000 m, and exhibits a complex geological history that includes tectonic movements, orogenesis, and periodic climatic change (e.g., Ju et al., 2007; Rüber et al., 2004; Zhang et al., 1990). Based on patterns of intraspecific genetic variation and buffer-zone models, Huangshan hosts refugia of eastern Asian conifers, frogs, non-migratory birds and Asian salamanders (Gao et al., 2007; Li et al., 2009; Murphy et al., 2000; Wu et al., 2013; Zhang et al., 2008).

Received: 18 June 2015; Accepted: 25 February 2016

Foundation items: This research was funded by the National Natural Science Foundation of China (NSFC 30870290, 31071891 and 31471968) ^{*}Corresponding authors, E-mail: snakeman@hsu.edu.cn;clad@ihb.ac.cn [#]Authors contributed equally to this work

DOI:10.13918/j.issn.2095-8137.2016.2.103

In view of modern genetics, genetic diversity in a given species is closely related to its adaptability, variability, and evolutionary potentiality, with genetic variation considered a prerequisite for organisms to cope with environmental uncertainty (Conrad, 1983). Herein, we report on the genetic diversity of *Z. platypus* from eastern China based on extensive sampling of the Huangshan area together with prior *cytb* sequence data from mainland China, Taiwan (Perdices et al., 2004; Perdices & Coelho, 2006; Wang et al., 2007), and Japan (He et al., 2004; Kawamura et al., 2014; Kitamura et al., 2012; Sasaki et al., 2007; Wang et al., 2007). We further evaluated the matrilineal diversity of *Z. platypus* and revealed the possible ecological significance of the Huangshan area.

MATERIALS AND METHODS

Sampling

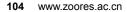
We evaluated 169 sequences in total, including 26 from five Huangshan counties (Table 1), 137 from Perdices et al. (2004) and Perdices & Coelho (2006), one from Wang et al. (2007), and five from Japan (AF309085, He et al. (2004); AB198972, Sasaki et al. (2007); AY958194, Wang et al. (2007); AB620130, Kitamura et al. (2012); and AB366543, Kawamura et al. (2014)). Our new samples were preserved and deposited in the Museum of Huangshan University (Voucher numbers: HUM201201–26). Sampling sites in this study are shown in Figure 1.

PCR amplification and sequencing

Fresh dorsal muscle tissues were removed from the 26 Huangshan individuals and immediately preserved in 95% ethanol for sequencing complete mitochondrial cytb. Total DNA was extracted from tissues using standard phenol/chloroform techniques (Sambrook et al., 1989). Cyt b was amplified using polymerase chain reaction (PCR) with the following sets of primers: LCB1 (5'-AATGACTTGAAGAACCACCGT-3') and HA (5'-CAACGATCTCCGGTTTACAAGAC-3') (Brito et al., 1997; Schmidt & Gold, 1993). Reagents included 100 ng of template DNA, 1 μL of each primer, 5 μL of 10× reaction buffer, 2 μL dNTPs (each 2.5 mmol/L), and 2.0 U of Tag DNA polymerase. The reactions were cycled as follows: an initial preheating at 94 °C for 3 min, 30 cycles of denaturation at 94 °C for 1 min, annealing at 55 °C for 40 s, extension at 72 °C for 1 min, and a final extension at 72 °C for 5 min. We next obtained nucleotide sequences through fractionation, purification, and sequencing according to Tiangen's protocols. Newly obtained haplotype sequences were deposited in GenBank under Accession Nos. KM491716-35 (Table 1).

Matrilineal genealogy and population structure

We used 916 bp out of 1 140 bp of the *cytb* sequences in the following analyses. The newly obtained sequences and those downloaded from GenBank were aligned using Clustal X (Thompson et al., 1997). For phylogenetic reconstruction, two closely related species, *Zacco temmincki* and *Candidia barbatus* (Mayden et al., 2007; Wang et al., 2011), were chosen as outgroups.



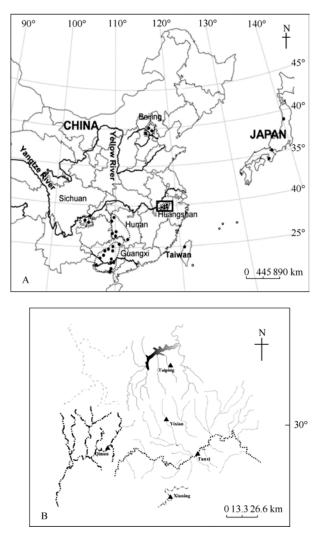


Figure 1 Sampling localities (A) and main drainages of the Huangshan area (B)

Black dots refer to samples from GenBank, and triangles are from this study. This map shows the seven geographic units grouped according to geographic distances and similarities. Xiuning, Qimen, and Taiping drainages belong to the Yangtze River, while Tunxi and Yixian drainages belong to the Qiantang River.

Bayesian inference (BI) and maximum likelihood (ML) were used to reconstruct a bifurcating tree using MrBayes v3.0 (Huelsenbeck & Ronquist, 2001) and RAxML at the CIPRES Science gateway (http: //www.phylo.org/portal2/login!input.action), respectively. JModelTest v. 0.1.1 (Posada, 2008) was used to find the best model of nucleotide evolution for ML based on the Akaike Information Criterion (AIC) and for BI based on the Bayesian Information Criterion (BIC). Analyses selected the TN93+G model. Bayesian posterior probabilities (BPP) and the frequencies of nodal resolution were obtained by Markov Chain Monte Carlo (MCMC) analysis with one cold chain and three heated chains. The BI analysis used 10 000 000 generations, with sampling every 1 000 generations and discarding the first $3x10^6$ generations as burn-in. We ran four analyses starting

with random trees and a consensus of the resulting 36 000 trees was computed from all four runs.

Table 1 Information for samples newly obtained in this study, including localities, rivers, sample sizes, haplotypes, coordinates, voucher specimens and GenBank accession numbers

Localities	Sample Size (<i>n</i>)	Rivers	Coordinates	Haplotypes	Voucher specimens	GenBank Accession No.
Qimen	4	Chang Jiang	N29°84'51", E117°71'77"	Qimen1	HUM201212	KM491716
				Qimen2	HUM201213	KM491717
				Qimen3	HUM201214	KM491718
				Qimen4	HUM201215	KM491719
Xiuning	8	Jiangwan He	N29°43'20", E118°16'75"	Xiuning1	HUM201201	KM491733
				Xiuning2	HUM201202-03	KM491731
				Xiuning3	HUM201204-06	KM491732
				Xiuning4	HUM201207	KM491735
				Xiuning5	HUM201208	KM491734
Tunxi	3	Xinan Jiang	N29°70'43", E118°31'16"	Tunxi1	HUM201209	KM491728
				Tunxi2	HUM201210	KM491729
				Tunxi3	HUM201211	KM491724
Yixian	6	Xinan Jiang	N29°92'06", E118°10'13"	Yixian1	HUM201216	KM491730
				Yixian2	HUM201217	KM491727
				Yixian3	HUM201218	KM491726
				Yixian4	HUM201219-20	KM491725
				Yixian5	HUM201221	KM491723
Taiping	5	Taiping Hu	N30°36'16", E118°04'65"	Taiping1	HUM201222	KM491722
				Taiping2	HUM201223-25	KM491720
				Taiping3	HUM201226	KM491721

Sampling information of extant sequences is not listed in this table.

Estimation of divergence time

Divergence times among the main lineages of *Z. platypus* were estimated using a Bayesian MCMC approach implemented in BEAST V.1.7.5 based on a strict molecular clock (Drummond & Rambaut, 2007). The parameters were: substitution model, TN93+G; tree prior, Coalescent: constant size; normal distribution; 10 million generations; parameters logged every 1 000; burn-in value=1 000. The molecular clock of cyprinids was assumed to be 1.52% site⁻¹ Ma (million years)⁻¹ (Doadrio et al., 2002) for *cyt b*.

RESULTS

A total of 75 haplotypes were defined from all 169 in-group individuals. The topologies of the BI and ML trees were nearly identical (Figure 2). The haplotypes were grouped into main clade 1 and 2. Clade 1 hosted individuals from Huangshan, Sichuan, Hunan and Guangxi and clade 2 contained specimens from Huangshan, Beijing and Japan. We identified ten matrilines of *Z. platypus* according to the topology of the phylogenetic tree and the genetic variation between the ten matrilines.

Six of the ten lineages involved Huangshan individuals, and

four consisted entirely of Huangshan individuals. Moreover, samples from the same Huangshan location were grouped into different clades. For example, Qimen had samples from matrilines A, F and J, and Xiuning had samples from matrilines A and I. The genetic divergences of these samples were significant, and the maximum pairwise differences from the same counties reached 6.0% (Xiuning) and 21.9% (Qimen).

We grouped sampling localities into seven geographic units according to geographic distances and then calculated the nucleotide diversity within them. Huangshan showed remarkably high nucleotide diversity relative to other groups (2.5-111.7 times that of others) although the geographical area of Huangshan was less than that of the other units (Table 3).

Divergence times estimated for the in-group nodes are shown in Figure 3. The initial divergence occurred at about 10.67 Ma.

DISCUSSION

Perdices et al. (2004) and Perdices & Coelho (2006) divided Z. *platypus* sampled in southern, western and northern China into matrilines A–F. They suggested that the long-term interruption of gene flow might have caused the diversification and an underestimation of the number of species. Our analyses identified ten matrilines of *Z. platypus* in Chinese and some Japanese populations. This confirms the prediction of Perdices

et al. (2004, page: 9) that "exhaustive sampling of other tributaries might evidence other *Zacco* lineages". This is also in accordance with that found for *Opsariichthys bidens*, a sympatric species of *Z. platypus* (Perdices et al., 2005).

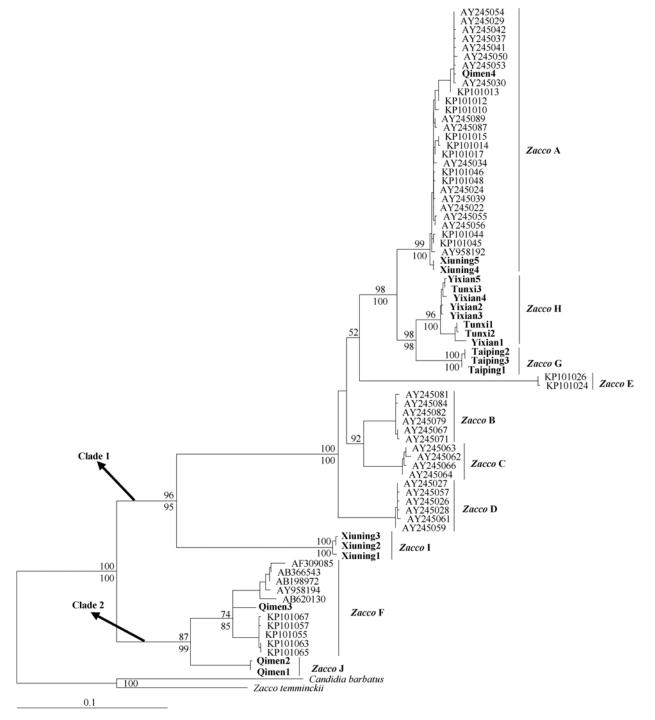


Figure 2 Phylogenetic tree derived from the maximum likelihood of the cyt b sequences

Values above branches represent the support level of ML (BSP) and values below branches represent the popularity rating of BI (BPP). Vertical bars indicate the mtDNA lineage assignment (A-J). Zacco A-F follow the nomenclature of Perdices & Coelho (2006) and the others follow the alphabet. Bold types are Huangshan populations.

	-	-								
	А	В	С	D	E	F	G	Н	Ι	J
А										
В	0.067									
С	0.078	0.044								
D	0.076	0.059	0.065							
Е	0.118	0.112	0.109	0.108						
F	0.146	0.148	0.163	0.158	0.174					
G	0.052	0.073	0.074	0.074	0.122	0.152				
Н	0.059	0.076	0.091	0.08	0.128	0.158	0.044			
I	0.157	0.157	0.158	0.146	0.184	0.155	0.152	0.155		
J	0.157	0.151	0.156	0.151	0.181	0.067	0.153	0.164	0.135	

Table 2 Matrix of pairwise genetic variation between matrilines (A-J) of Z. platypus

Table 3 Matrilines, haplotype (h), and nucleotide diversity (π) with standard errors (SE) for each geographic unit

Geographic unit	mtDNA lineage	No. of samples (n)	h±SE	π±SE
Huangshan	A, F, G, H, I, J	26	0.9631±0.0219	0.085751±0.042501
Beijing	F	23	0.6087±0.0761	0.000768±0.000670
Hunan	A, C, D	49	0.9092±0.0262	0.034648±0.017036
Sichuan	В	19	0.6374±0.1045	0.001098±0.000863
Guangxi	A, E	46	0.6963±0.0740	0.029668±0.014733
Taiwan	А	1	_	_
Japan	F	5	_	0.011681±0.007511

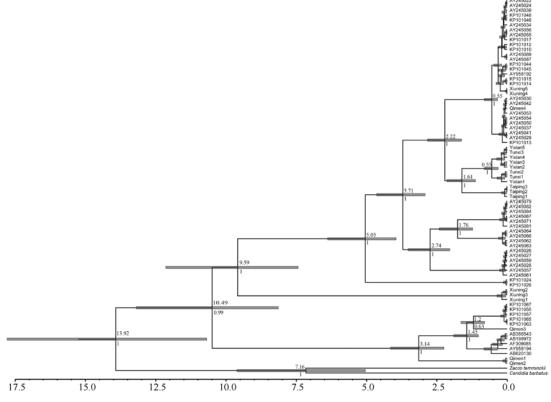


Figure 3 Time tree of Z. platypus

Tree topology derived from BEAST analyses of all 77 haplotypes. Numbers above branches represent node age and values below are support rates. Gray bar represents 95% posterior credible intervals.

Some drainages still await sampling, such as the Yellow River, one of the most important drainages in China. Future research should detect additional matrilines of *Zacco*, while morphological analyses may help differentiate morphological differences of taxonomic significance.

Grant & Bowen (1998) interpreted four basic population history scenarios based on haplotype and nucleotide diversities, which can also be used to clarify the history of *Z. platypus* populations. Our results revealed a pattern of high haplotype and nucleotide diversity in the Huangshan population (Table 3), which likely indicates large stable populations with long evolutionary histories or secondary contact between differentiated lineages (Grant & Bowen, 1998). The highest levels of genetic variation may occur in the region of origin. For example, Savolainen et al. (2002) claimed an East Asian origin for the domestic dog in part due to the area having the highest level of genetic diversity.

Genetic variability in mtDNA has been reported in fish species. Several scenarios have been proposed to explain the maintenance of high haplotype diversity within populations, including large population size, environmental heterogeneity, and life history traits that favor rapid population increase (Han et al., 2008; Ju et al., 2013; Yang et al., 2012). Huangshan has a heterogeneous topography, with mountains of elevation lower than 2 000 m maintaining stable climatic conditions during the Pleistocene. This condition likely provided glacial refugia for many species (Gao et al., 2007; Li et al., 2009; Qian & Ricklefs, 2000; Wu et al., 2013; Zhang et al., 2008). At least three other species or species groups have high levels of nucleotide diversity in the Huangshan area, including the Chinese giant salamander, sharpsnouted pit viper and Asian salamander (Huang et al., 2007; Murphy et al., 2000; Wu et al., 2013). These co-occurrences indicate that Huangshan hosts old lineages.

We only used mtDNA for genetic analyses. Therefore, it will be necessary to gather and analyze nuclear DNA data in the future to assess population structure and gene flow and thus better inform the demographic history of this fish species.

ACKNOWLEDGEMENTS

We extend our thanks to Dr. Robert W. MURPHY for his valuable comments on this manuscript. We thank Jin-Min CHEN (Yunnan University), Bao-Lin ZHANG (Kunming Institute of Zoology, CAS) and Li-Fang PENG (Nanjing Forestry University) for their help in data processing. We further thank Dian-Cheng YANG (Nanjing Forestry University) and Jun-Sheng CUI (Anhui Agricultural University) for sampling.

REFERENCES

Berrebi P, Boissin E, Fang F, Cattaneo-Berrebi G. 2005. Intron polymorphism (EPIC-PCR) reveals phylogeographic structure of *Zacco platypus* in China: a possible target for aquaculture development. *Heredity*, **94**(6): 589-598.

Brito RM, Briolay J, Galtier N, Bouvet Y, Coelho MM. 1997. Phylogenetic relationships within genus *Leuciscus* (Pisces, Cyprinidae) in Portuguese fresh waters, based on mitochondrial DNA cytochrome *b* sequences. *Molecular Phylogenetics and Evolution*, **8**(3): 435-442.

Chen Y. 1998. Fauna Sinica. Beijing, China: Science Press.

Conrad M. 1983. Adaptability: The Significance of Variability from Molecule to Ecosystem. New York: Plenum Press.

Deng FY, Zhang CG, Zhao YH, Zhou QH, Zhang J. 2013. Diversity and community structure of the fishes in the headstream region of the Dongjiang River. *Chinese Journal of Zoology*, **48**(2): 161-173.

Doadrio I, Carmona JA, Machordom A. 2002. Haplotype diversity and phylogenetic relationships among the Iberian Barbels (*Barbus*, Cyprinidae) reveal two evolutionary lineages. *The Journal of Heredity*, **93**(2): 140-147.

Drummond AJ, Rambaut AJ. 2007. BEAST: Bayesian evolutionary analysis by sampling trees. *BMC Evolutionary Biology*, **7**: 214.

Gao LM, Möller M, Zhang XM, Hollingsworth ML, Liu J, Mill RR, Gibby M, Li DZ. 2007. High variation and strong phylogeographic pattern among cpDNA haplotypes in *Taxus wallichiana* (Taxaceae) in China and North Vietnam. *Molecular Ecology*, **16**(22): 4684-4698.

Grant WAS, Bowen BW. 1998. Shallow population histories in deep evolutionary lineages of marine fishes: insights from sardines and anchovies and lessons for conservation. *Journal of Heredity*, **89**(5): 415-426.

Han ZQ, Li YZ, Chen GB, Gao TX. 2008. Population genetic structure of coral reef species *Plectorhinchus flavomaculatus* in South China Sea. *African Journal of Biotechnology*, **7**(11): 1774-1781.

He SP, Liu HZ, Chen YY, Kuwahara M, Nakajima T, Zhong Y. 2004. Molecular phylogenetic relationships of Eastern Asian Cyprinidae (Pisces: Cypriniformes) inferred from cytochrome *b* sequences. *Science in China Series C-Life Sciences*, **47**(2): 130-138.

Huang S, He SP, Peng ZG, Zhao K, Zhao EM. 2007. Molecular phylogeography of endangered sharp-snouted pitviper (*Deinagkistrodon acutus*; Reptilia, Viperidae) in Mainland China. *Molecular Phylogenetics and Evolution*, **44**(3): 942-952.

Huelsenbeck JP, Ronquist F. 2001. MRBAYES: Bayesian inference of phylogenetic trees. *Bioinformatics*, **17**(8): 754-755.

Ju LX, Wang HJ, Jiang DB. 2007. Simulation of the Last Glacial Maximum climate over East Asia with a regional climate model nested in a general circulation model. *Palaeogeography, Palaeoclimatology, Palaeoecology,* **248**(3-4): 376-390.

Ju YM, Hsu CH, Fang LS, Lin HD, Wu JH, Han CC, Chen IS, Chiang TY. 2013 Population structure and demographic history of *Sicyopterus japonicus* (Perciformes; Gobiidae) in Taiwan inferred from mitochondrial control region sequences. *Genetics and Molecular Research*, **12**(3): 4046-4059.

Kawamura K, Ueda T, Arai R, Smith C. 2014. Phylogenetic relationships of bitterling fishes (Teleostei: Cypriniformes: Acheilognathinae), inferred from mitochondrial cytochrome *b* sequences. *Zoological Science*, **31**(5): 321-329. Kitamura J, Nagata N, Nakajima J, Sota T. 2012. Divergence of ovipositor length and egg shape in a brood parasitic bitterling fish through the use of different mussel hosts. *Journal of Evolutionary Biology*, **25**(3): 566-573.

Li SH, Yeung CKL, Feinstein J, Han LX, Le MH, Wang CX, Ding P. 2009. Sailing through the Late Pleistocene: unusual historical demography of an East Asian endemic, the Chinese Hwamei (*Leucodioptron canorum canorum*), during the last glacial period. *Molecular Ecology*, **18**(4): 622-633. Mayden RL, Tang KL, Conway KW, Freyhof J, Chamberlain S, Haskins M, Schneider L, Sudkamp M, Wood RM, Agnew M, Bufalino A, Sulaiman Z, Miya M, Saitoh K, He SP. 2007. Phylogenetic relationships of *Danio* within the order Cypriniformes: a framework for comparative and evolutionary studies of a model species. *Journal of Experimental Zoology Part B: Molecular and*

Developmental Evolution, 308B(5): 642-654.

Murphy RW, Fu JZ, Upton DE, De Lema T, Zhao EM. 2000. Genetic variability among endangered Chinese giant salamanders, *Andrias davidianus*. *Molecular Ecology*, **9**(10): 1539-1547.

Perdices A, Cunha C, Coelho MM. 2004. Phylogenetic structure of *Zacco platypus* (Teleostei, Cyprinidae) populations on the upper and middle Chang Jiang (=Yangtze) drainage inferred from cytochrome b sequences. *Molecular Phylogenetics and Evolution*, **31**(1): 192-203.

Perdices A, Sayanda D, Coelho MM. 2005. Mitochondrial diversity of *Opsariichthys bidens* (Teleostei, Cyprinidae) in three Chinese drainages. *Molecular Phylogenetics and Evolution*, **37**(3): 920-927.

Perdices A, Coelho MM. 2006. Comparative phylogeography of *Zacco platypus* and *Opsariichthys bidens* (Teleostei, Cyprinidae) in China based on cytochrome b sequences. *Journal of Zoological Systematics and Evolutionary Research*, **44**(4): 330-338.

Posada D. 2008. jModelTest: phylogenetic model averaging. *Molecular Biology and Evolution*, **25**(7): 1253-1256.

Qian H, Ricklefs RE. 2000. Large-scale processes and the Asian bias in species diversity of temperate plants. *Nature*, **407**(6801): 180-182.

Rüber L, Britz R, Kullandr SO, Zardoya R. 2004. Evolutionary and biogeographic patterns of the *Badidae* (Teleostei: Perciformes) inferred from mitochondrial and nuclear DNA sequence data. *Molecular Phylogenetics and Evolution*, **32**(3): 1010-1022.

Sambrook J, Fritsch EF, Maniatis T. 1989. Molecular Cloning: A Laboratory Manual. 2nd ed. New York: Cold Spring Harbor Laboratory Press.

Sasaki T, Kartavtsev YP, Chiba SN, Uematsu T, Sviridov VV, Hanzawa N. 2007. Genetic divergence and phylogenetic independence of Far Eastern species in subfamily Leuciscinae (Pisces: Cyprinidae) inferred from mitochondrial DNA analyses. *Genes & Genetic Systems*, **82**(4): 329-340.

Savolainen P, Zhang YP, Luo J, Lundeberg J, Leitner T. 2002. Genetic evidence for an East Asian origin of domestic dogs. *Science*, **298**(5598): 1610-1613.

Schmidt TR, Gold JR. 1993. Complete sequence of the mitochondrial cytochrome *b* gene in the Cherryfin shiner, *Lythrurus roseipinnis* (Teleostei: Cyprinidae). *Copeia*, **1993**(3): 880-883.

Thompson JD, Gibson TJ, Plewniak F, Jeanmougin F, Higgins DG. 1997. The CLUSTAL_X windows interface: flexible strategies for multiple sequence alignment aided by quality analysis tools. *Nucleic Acids Research*, **25**(24): 4876-4882.

Wang CF, Hsieh CH, Lee SC, Wang HY. 2011. Systematics and phylogeography of the Taiwanese endemic minnow *Candidia barbatus* (Pisces: Cyprinidae) based on DNA sequence, allozymic, and morphological analyses. *Zoological Journal of the Linnean Society*, **161**(3): 613-632.

Wang HY, Wang CF, Du SY, Lee SC. 2007. New insights on molecular systematics of opsariichthines based on cytochrome *b* sequencing. *Journal of Fish Biology*, **71**: 18-32.

Wu YK, Wang YZ, Jiang K, Hanken J. 2013. Significance of pre-Quaternary climate change for montane species diversity: Insights from Asian salamanders (Salamandridae: Pachytriton). *Molecular Phylogenetics and Evolution*, **66**(1): 380-390.

Yang JQ, Tang WQ, Liao TY, Sun Y, Zhou ZC, Han CC, Liu D, Lin HD. 2012. Phylogeographical Analysis on *Squalidus argentatus* Recapitulates Historical Landscapes and Drainage Evolution on the Island of Taiwan and Mainland China. *International Journal of Molecular Sciences*, **13**(12): 1405-1425.

Zhang H, Yan J, Zhang GQ, Zhou KY. 2008. Phylogeography and demographic history of Chinese black-spotted frog populations (*Pelophylax nigromaculata*): evidence for independent refugia expansion and secondary contact. *BMC Evolutionary Biology*, **8**: 21.

Zhang HN, Chen CG, Huang KR, Li ZQ, Zhang FL, Chen G. 1990. The New Geological Structures, Tectonic Movements and Geological Environment in Coastal Line of South China. Beijing: Earthquake Press. (in Chinese)