

MITOGENOME ANNOUNCEMENT



The mitochondrial genome of the assassin bug *Scloмина erinacea* (Hemiptera: Reduviidae)

Qian Zhao^a, Ping Zhao^b, Hu Li^a, Wanzhi Cai^a and Fan Song^a

^aDepartment of Entomology and MOA Key Lab of Pest Monitoring and Green Management College of Plant Protection, China Agricultural University, Beijing, China; ^bGuangdong Key Laboratory of Animal Conservation and Resource Utilization, Guangdong Public Laboratory of Wild Animal Conservation and Utilization, Guangdong Institute of Applied Biological Resources, Guangzhou, China

ABSTRACT

The complete mitochondrial genome (mitogenome) of the assassin bug, *Scloмина erinacea* Stål, was determined in the present study. The sequenced mitogenome is a typical circular DNA molecule which is 15,828 bp in length, containing 13 protein-coding genes, two rRNA genes, 22 tRNA genes and a putative control region. All protein-coding genes are initiated by ATN codons and terminated by TAA or TAG codons except *COII*, *COIII* and *ND4* which use a single T residue as the stop codon. All tRNAs have the cloverleaf structure with the exception of *tRNA^{Ser(AGN)}* and the length of them range from 62 to 70 bp. The control region is 818 bp long with an A + T content of 67.1%. The phylogenetic analysis supports the monophyly of Harpactorinae and *Scloмина erinacea* is the closest relative to *Macracanthopsis nodipes*.

ARTICLE HISTORY

Received 28 June 2019
Accepted 1 August 2019

KEYWORDS

Mitochondrial genome;
Reduviidae; Harpactorinae;
Scloмина erinacea

The assassin bug Harpactorinae (Hemiptera: Reduviidae) is the largest subfamily of Reduviidae distributed around the world mainly in tropical and subtropical regions. The Harpactorinae currently contains about 180 species in China, occurring in the southern region of China. *Scloмина erinacea*, which belongs to the subfamily Harpactorinae, is a widely distributed predatory natural enemy insect in China and can prey on a variety of agricultural pests (Zhao and Liu 2015). In this study, the complete mitogenome of *Scloмина erinacea* is sequenced and annotated for the first time. The samples were collected in Meiling, Jiangxi, China (28°48'39.0"N 115°41'52.1"E). Voucher specimen was deposited at the Entomological Museum of China Agricultural University (No. VCim-00168) and the sequence has been submitted to GenBank (Accession number: MK696614).

The mitogenome is a single circular DNA molecule of 15,828 bp in size that encode 37 genes (13 protein-coding genes, 22 tRNA genes, and two rRNA genes) and a control region. Gene order is identical to the putative ancestral gene arrangement of insect (Cameron 2014; Song et al. 2016a; Li et al. 2017; Liu et al. 2019). Except control region, this mitogenome has 3 inter-genic regions > 90 bp. A 126 bp noncoding region could be found between *tRNA^{Cys}* and *tRNA^{Tyr}* and a 94 bp noncoding region exist between *CytB* and *tRNA^{Ser}* (*UCN*). Another 218 bp noncoding region is located between *tRNA^{Ser(UCN)}* and *ND1*, which has also been found in other assassin bugs (Sun et al. 2019). There are totally 51 bp

overlapped nucleotides between neighboring genes in 12 locations, ranging from 1 to 14 bp in size.

The nucleotide composition of the whole mitogenome shows highly A + T biased. The A + T content is 71.6% with positive AT-skew (0.14) and negative GC-skew (−0.17). All protein-coding genes initiate with ATN as the start codon. The stop codon TAA terminates 7 protein-coding genes and TAG terminates 3 protein-coding genes. Whereas *COII*, *COIII* and *ND4* use a single T residue as incomplete stop codon which has repeatedly found in other insect mitogenomes (Song et al. 2016b; Liu et al. 2018; Sun et al. 2019).

The length of the 22 sequenced tRNA genes range from 62 to 70 bp and most tRNA genes could be folded into the typical cloverleaf structure except for *tRNA^{Ser(AGN)}*, due to the deficiency of the dihydrouridine (DHU) arm which is typical feature of insect mitogenomes (Jiang et al. 2016). The *LrRNA* is 1,267 bp in length with an A + T content of 74.1% and the *srRNA* is 778 bp in length with an A + T content of 72.4%. The control region, which is located between *srRNA* and *tRNA^{Ile}*, is 818 bp long and is biased toward A + T (67.1%).

Phylogenetic tree based on the maximum likelihood method shows Harpactorinae is monophyletic (Figure 1), which is also recovered in previous molecular and morphological analyses (Weirauch et al. 2014; Liu et al. 2019). The sister relationship between *Scloмина erinacea* and *Macracanthopsis nodipes* is also highly supported (bootstrap = 100).

CONTACT Fan Song ✉ fansong@cau.edu.cn; Ping Zhao ✉ zpyajl@126.com Guangdong Key Laboratory of Animal Conservation and Resource Utilization, Guangdong Public Laboratory of Wild Animal Conservation and Utilization, Guangdong Institute of Applied Biological Resources, Guangzhou, 510260 China

© 2019 The Author(s). Published by Informa UK Limited, trading as Taylor & Francis Group.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

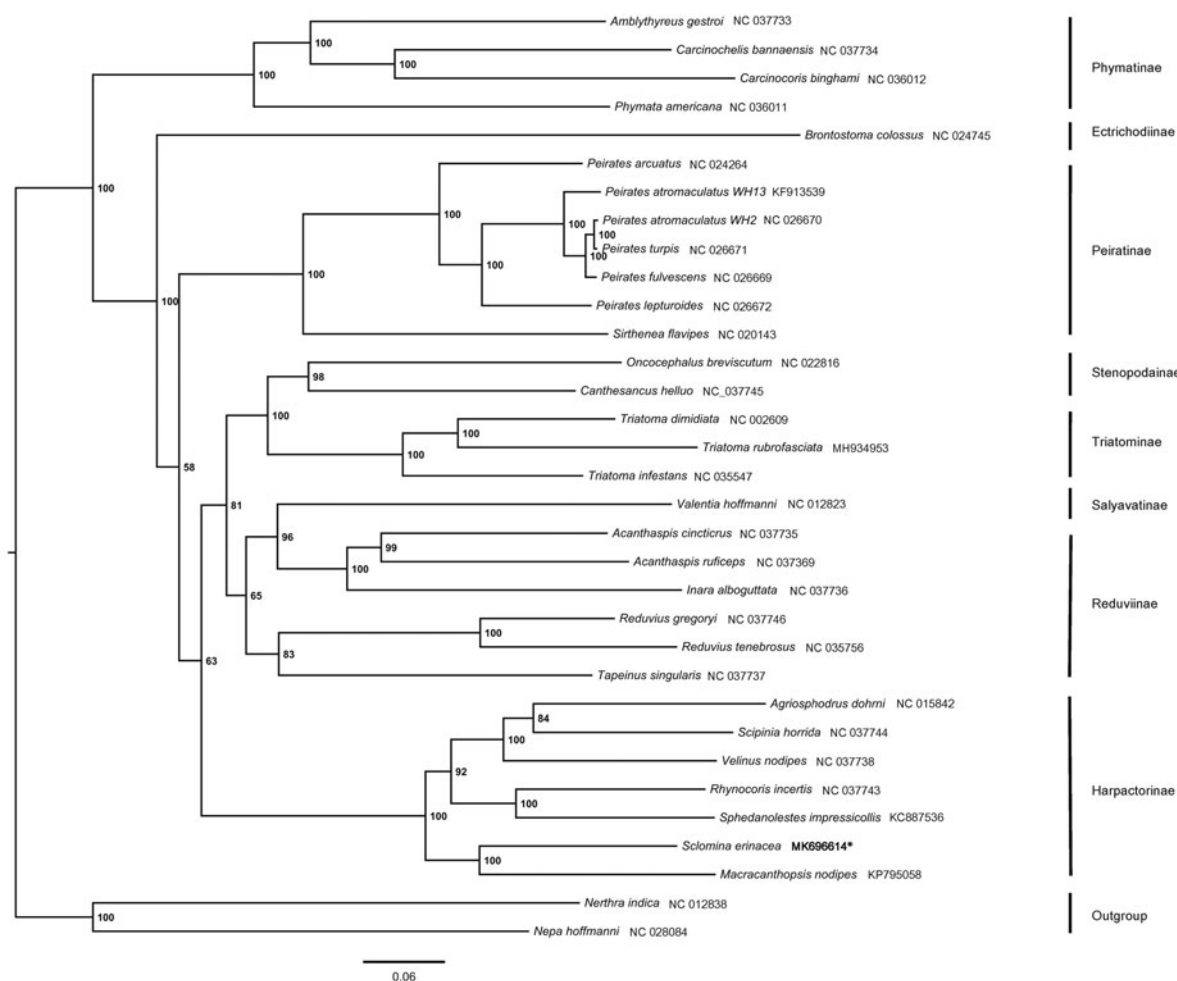


Figure 1. Phylogenetic relationship of *Scloimina erinacea* and other 32 species among Reduviidae. Phylogenetic tree was inferred from ML analysis of the 13 protein-coding genes and two rRNAs genes (12,812 bp) and generated by IQ-TREE 1.6.5 (Trifinopoulos et al. 2016) under the GTR+I+G model. The number beside the nodes are percentages of 1000 bootstrap values. The GenBank accession number of the species is shown behind the species name. The newly sequenced mitochondrial genome is highlighted by the asterisk.

Disclosure statement

No potential conflict of interest was reported by the authors.

Funding

Funding for this study was supported by grant from the National Natural Science Foundation of China [Nos. 31760634 and 31772498].

References

- Cameron SL. 2014. Insect mitochondrial genomics: implications for evolution and phylogeny. *Annu Rev Entomol.* 59:95–117.
- Jiang P, Li H, Song F, Cai Y, Wang JY, Liu J, Cai WZ. 2016. Duplication and remodelling of tRNA genes in the mitochondrial genome of *Reduvius tenebrosus* (Hemiptera: Reduviidae). *IJMS.* 17:951.
- Li H, Leavengood JM, Jr, Chapman EG, Burkhardt D, Song F, Jiang P, Liu J, Zhou X, Cai WZ. 2017. Mitochondrial phylogenomics of Hemiptera reveals adaptive innovations driving the diversification of true bugs. *Proc R Soc B.* 284:20171223.
- Liu YQ, Li H, Song F, Zhao YS, Wilson JJ, Cai WZ. 2019. Higher-level phylogeny and evolutionary history of Pentatomomorpha (Hemiptera: Heteroptera) inferred from mitochondrial genome sequences. *Syst Entomol.* 44:810.
- Liu YQ, Song F, Jiang P, Wilson JJ, Cai WZ, Li H. 2018. Compositional heterogeneity in true bug mitochondrial phylogenomics. *Mol Phylogenet Evol.* 118:135–144.
- Song F, Li H, Jiang P, Zhou X, Liu JP, Sun CH, Vogler AP, Cai WZ. 2016a. Capturing the phylogeny of Holometabola with mitochondrial genome data and Bayesian site-heterogeneous mixture models. *Genome Biol Evol.* 8:1411–1426.
- Song F, Li H, Shao R, Shi A, Bai X, Zheng X, Heiss E, Cai WZ. 2016b. Rearrangement of mitochondrial tRNA genes in flat bugs (Hemiptera: Aradidae). *Sci Rep.* 6:25725.
- Sun ZQ, Liu YQ, Wilson JJ, Chen Z, Song F, Cai WZ, Li H. 2019. Mitochondrial genome of *Phalantus geniculatus* (Hemiptera: Reduviidae): *trnT* duplication and phylogenetic implications. *Int J Biol Macromol.* 129:110–115.
- Trifinopoulos J, Nguyen LT, Haeseler AV, Minh BQ. 2016. W-IQ-TREE: a fast online phylogenetic tool for maximum likelihood analysis. *Nucleic Acids Res.* 44:W232–W235.
- Weirauch C, Berenger JM, Berniker L, Forero D, Forthman M, Frangenberg S, Michael A, Paiero SM, Udah O, Waston C, et al. 2014. An illustrated identification key to assassin bug subfamilies and tribes (Hemiptera: Reduviidae). *Can J Arthrod Ident.* 26:1–115.
- Zhao P, Liu HX. 2015. Investigation of the Subfamily Harpactorinae (Hemiptera: Heteroptera: Reduviidae) in Nonggang National Nature Reserve of Guangxi. *J Kaili Univ.* 33:70–74.