

Mitochondrial DNA Part B



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MITOGENOME ANNOUNCEMENT

3 OPEN ACCESS

Complete mitochondrial genome of the winghead shark, *Eusphyra blochii* (Elasmobranchii: Sphyrnidae)

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ABSTRACT

The complete mitochondrial genome (16,726 bp) of the winghead shark, *Eusphyra blochii* is presented. This species is exploited throughout parts of its range, and is currently listed as Near Threatened by the IUCN Red List. A phylogenetic analysis placed *E. blochii* within the Carcharhiniformes, as a sister taxon to *Sphyrna lewini* and *S. zygaena*.

ARTICLE HISTORY

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KEYWORDS

Carcharhiniformes; *Eusphyra blochii*; mitogenome; Sphyrnidae

Eusphyra is a monotypic genus of hammerhead shark in the family Sphyrnidae, order Carcharhiniformes. It is distinctive in having the most proportionally extreme narial separation among hammerhead sharks, which is suggested to increase odour resolution (Kajiura et al. 2005). Eusphyra blochii is found in shallow waters over continental and insular shelves of the Indian and Pacific Oceans, from the Persian Gulf to the Philippines, and from China to Australia (Simpfendorfer 2003; Ebert et al. 2013). Members of the family Sphyrnidae have been exploited by multiple fisheries leading to two species being listed as Endangered, two as Vulnerable and two as Near Threatened by the IUCN Red List (Pérez-Jiménez 2014). Life history traits coupled with fishing pressures make these species particularly vulnerable to over-exploitation. Eusphyra blochii is currently listed as Near Threatened due to heavy exploitation by fisheries in parts of its range (i.e. Gulf of Thailand, India and Indonesia) and light exploitation in Australian waters (Simpfendorfer 2003).

In this study, we determined the complete mitochondrial genome of *E. blochii* using a female specimen collected from Fog Bay, in the Timor Sea of northern Australia. A tissue sample from the specimen is deposited in the collection of Janine Caira at the University of Connecticut under field number AU-70 and extracted DNA is kept at the Hollings Marine Laboratory in Charleston, SC, under collection number GN5092. Total gDNA was extracted from liver tissue using the

E.Z.N.A Tissue DNA Kit (Omega Bio-Tek, Inc., Norcross, GA) following the protocol of the manufacturer and the sequence was annotated using MitoAnnotator (Iwasaki et al. 2013). PCR primers and protocols are available upon request.

Twenty-eight shark and four batoid whole mitochondrial genome sequences were downloaded from GenBank and protein-coding regions of the 32 mitochondrial genomes were concatenated and aligned with corresponding regions from *E. blochii*. The complementary strand sequences were used for *ND6*, and incomplete stop codons of genes were removed from alignment. The final length of the protein-coding portion of the alignment was 11,424 bp. The dataset was subjected to maximum likelihood analysis using PAUP*4.0a145 (Swofford 2002) under the general-time reversible (GTR)+invariable sites (I)+gamma model with parameter values estimated from an initial parsimony analysis.

The complete mitochondrial genome (16,726 bp in length) of *E. blochii* has typical vertebrate mitochondrial gene arrangement (Anderson et al. 1981) consisting of 13 proteincoding genes, 22 tRNA genes, two rRNA genes and one control region (GenBank accession no. KJ128290). All genes are encoded on the H-strand, except for *ND6* and eight tRNA genes, which are encoded on the L-strand.

The maximum likelihood tree is given in Figure 1. Based on this analysis and taxon sampling, *E. blochii* is placed within the Carcharhiniformes as sister taxon to *S. lewini* and *S. zygaena*, forming a monophyletic Sphyrnidae.

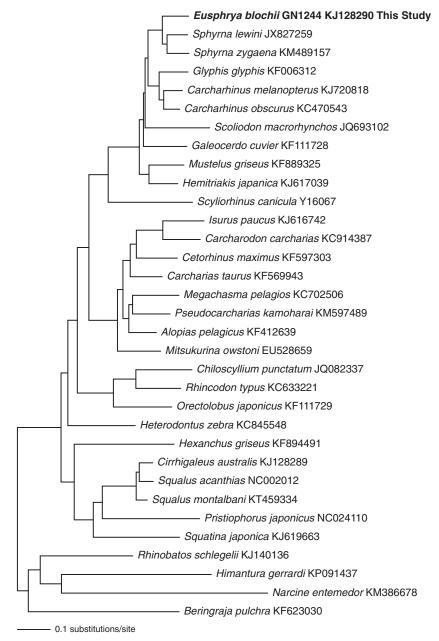


Figure 1. Maximum likelihood tree based on the protein-coding portion of the mitogenome dataset using the GTR+I+gamma model. GenBank accession numbers follow species names. *Eusphyra blochii* position is indicated in bold. Representative batoid species are used as outgroup.

Disclosure statement

The authors report no conflict of interest. The authors alone are responsible for the content and writing of the paper.

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