

**Table S2.** Primer pairs used for DNA amplification.

Gene	Size	Primer	Primer sequence	Annealing temp.	Publication
<i>COI</i>	379 bp	L6625 <sup>1</sup> H7005 <sup>2</sup>	5' CCGGATCCTTYTGRTTYTTYGGNCAYCC 3' 5' CCGGATCCACNACRTARTANGTRTCRTG 3'	50 °C	Hafner <i>et al.</i> (1994)
<i>COI<sup>L</sup></i>	670 bp	LCO1490 <sup>1</sup> HCO2198 <sup>2</sup>	5' GGTCACAAATCATAAAGATATTGG 3' 5' TAACTTCAGGGTGACCAAAAAATCA 3'	50 °C	Folmer <i>et al.</i> (1994)
<i>EF-1<math>\alpha</math></i>	347 bp	EF1-For 3 <sup>1</sup> EF1-Cho10 <sup>2</sup>	5' GGNGACAAYGTTGGYTTCAACG 3' 5' ACRGCVACKGTYTGHCKCATGTC 3'	50 °C	Danforth & Ji (1998)

bp - base pair, <sup>L</sup> - longer fragment of mitochondrial gene *COI*, <sup>1</sup> - forward, <sup>2</sup> - reverse

**Table S3.** PCR protocols.

Steps	1.	-	2.	3.	4.	-	5.	6.	7.	8.	9.
Genes ↓											
<i>COI</i>	94 °C 10 min.	<i>steps</i> 2–4 35x	94 °C 30 s.	46 °C 30 s.	65 °C 30 s.	-	65 °C 7 min.	4 °C ∞	-	-	-
<i>COI<sup>L</sup></i>	94 °C 3 min	<i>steps</i> 2–4 5x	94 °C 30 s.	45 °C 30 s.	72 °C 1 min.	<i>steps</i> 5–7 30x	94 °C 30 s.	51 °C 1 min.	72 °C 1 min.	72 °C 10 min.	4 °C ∞
<i>EF-1α</i>	95 °C 10 min.	<i>steps</i> 2–4 35x	94 °C 1m.	45 °C 45 s.	72 °C 45 s.	-	72 °C 10 m.	4 °C ∞	-	-	-

**Table S4.** Partitions with their models used in Bayesian Analysis (BA).

<b>Partition</b>	<b>Nucleotide codon position</b>	<b>Models</b>
1	1st position in COI-L; 3rd in COI (380-1040\3; 1-379\3)	GTR+G
2	1st position in COI (2-379\3)	GTR+I+G
3	3rd position in COI-L; 2nd in COI (382-1040\3; 3-379\3)	GTR+I+G
4	2nd position in COI-L (381-1040\3)	GTR+G
5	1st position in EF1 $\alpha$ (1041-1387\3)	HKY+I
6	2nd position in EF1 $\alpha$ (1042-1387\3)	GTR+G
7	3rd position in EF1 $\alpha$ (1043-1387\3)	GTR+I+G