

Supplementary 2. PCR primers used in this study. PCR conditions of all markers were in 35 cycles.

PCR primers					
Markers	Primers	Sequences (5'-3')			References
<i>COI</i>	FishF1	TCAACCAACCACAAAGACATTGGCAC			Ward <i>et al.</i> (2005)
	FishF2	TCGACTAATCATAAAGATATCGGCAC			
	FishR1	TAGACTTCTGGGTGGCCAAAGAATCA			
	FishR2	ACTTCAGGGTGACCGAAGAATCAGAA			
<i>cytb</i>	LopCytbF1	ARCCAGGRCNAAYGGNTYGAAA			Huang (2015)
	LopCytbF2	TTAACCAGGACNARYGGCTTGA			
	LopCytbR2	TTCGRSTTACAARNCCGAKGCTCT			
<i>rhodopsin</i>	RH193F	CNTATGAATAYCCTCAGTACTACC			Chen <i>et al.</i> (2003)
	RH1039R	TGCTTGTTTCATGCAGATGTAGA			
<i>RAG1</i>	2553F	CTGAGCTGCAGTCAGTACCATAAGATGT			López <i>et al.</i> (2004)
	4090R	CTGAGTCCTTGTGAGCTTCCATRAAYTT			
PCR conditions (Huang, 2015)					
Markers	Primary denaturation	Denaturation	Annealing	Extension	Additional extension
<i>COI</i>	94°C , 1 min	94°C , 30 sec	51°C , 30 sec	72°C , 40 sec	72°C , 2 min
<i>cytb</i>	95°C , 4 min	95°C , 40 sec	53°C , 40 sec	72°C , 1 min 15 sec	72°C , 7 min
<i>rhodopsin</i>	95°C , 5 min	95°C , 40 sec	55°C , 40 sec	72°C , 1 min	72°C , 7 min
<i>RAG1</i>	95°C , 5 min	95°C , 40 sec	53°C , 40 sec	72°C , 1 min 30 sec	72°C , 7 min

References

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