

Supplementary

Table S1: *Pinna nobilis* DNA samples PCR amplified with HapF1-HapR2 primers for Sanger sequencing.

Sample/ amplicon size	[DNA] (ng/μL)	A ₂₆₀ /A ₂₈₀	Origin
PN4 (300 pb)	19,4	1,68	Faeces
PN9 (300 pb)	41,9	1,68	Necropsy mantle and digestive gland <i>Pinna nobilis</i>
PN13 (300 pb)	37,6	1,81	
PN14 (300 pb)	56,5	1,8	

Table S2: *Pinna nobilis* DNA samples PCR amplified with COI1-COIL1 primers.

Sample/ amplicon size	[DNA] (ng/μL)	A ₂₆₀ /A ₂₈₀	Origin
PN1 (710 pb)	16,7	1,88	Mantle biopsy <i>Pinna nobilis</i>
PN2 (710 pb)	57	1,78	
PN3 (710 pb)	37,1	1,71	

Table S3: Blast analysis for Sanger sequences obtained from PCR positive *Pinna nobilis* samples

Sample	Most similar species (BLAST)	Query Cover	Identity
PN1	<i>Pinna nobilis</i> (KY321790.1)	87%	95.77%
PN2	<i>Pinna nobilis</i> (KY321794.1)	66%	84.14%
PN3	<i>Pinna nobilis</i> (JX854841.1)	100%	100%

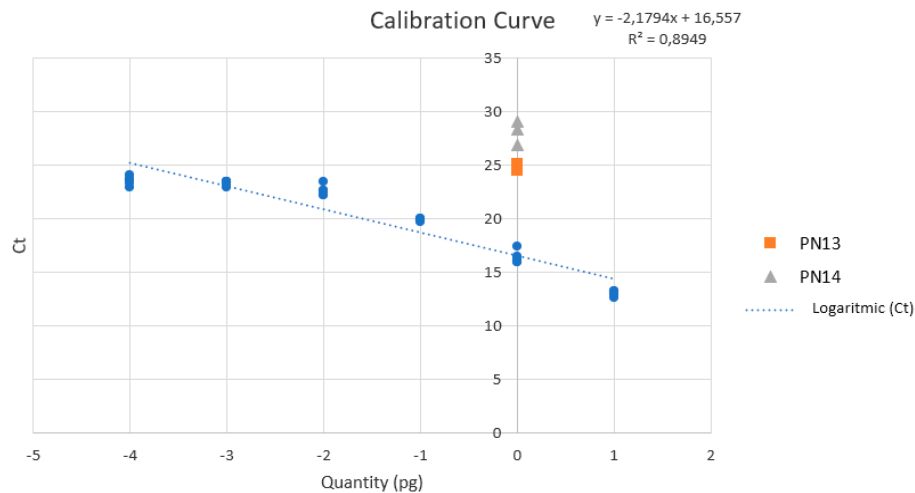


Figure S1: Calibration curve for qPCR for quantification of Haplosporidium pinnae.