

Additional file 1: Table of PCR primer sequences, annealing temperatures (T_a), cycle numbers and product sizes used for cloning the candidate 'housekeeping' genes.

Target gene	Primer name	Type	Primer sequence (5'-3')	T _a used for PCR (°C)	No. cycles	Product size (bp)
18S rRNA	18S rRNA Forward	'Core PCR'	GCGGCGACGACTCTTTTCGAATGTC	54.0	35	768
18S rRNA	18S rRNA Reverse	'Core PCR'	AGTTCCGACCGTAAACGATG			
<i>rpl8</i>	<i>rpl8</i> Forward	'Core PCR'	CTCCGTCTTCAAAGCCCATGT	54.0	35	708
<i>rpl8</i>	<i>rpl8</i> Reverse	'Core PCR'	TGTTCTCTCGCAGTCTGCCAG			
<i>efla</i>	<i>efla</i> Forward 1	'Core PCR 1'	GTGACAAACGTTGGCTTCAACG	54.8	25	377
<i>efla</i>	<i>efla</i> Reverse 1	'Core PCR 1'	CCTCATGTCACGCACAGCAAA			
<i>efla</i>	<i>efla</i> Forward 2	'Core PCR 2'	AGGCTGGTATCTCCAAGAACG	54.0	25	564
<i>efla</i>	<i>efla</i> Reverse 2	'Core PCR 2'	CGTTGAAGCCCAACGTTGTCAC			
<i>efla</i>	<i>efla</i> 5'-RACE	'5'-RACE'	GGTCTGCCGTTCTTGGAGATACCAGC	68.0	30	470
<i>efla</i>	<i>efla</i> 3'-RACE	'3'-RACE'	GCAAGAAGCTTGAGGACAACC	55.0	40	494