

Table S4. Primers used for qRT-PCR and RT-PCR amplification reactions:

Primer name	5'-3' sequence	Fragment	Length of amplification product/s (bp)
RTPnIAA30F	TGGTGTGTCAACTGTCCTATCTG	<i>PnIAA30</i> fragment used in qRT-PCR assays	109 ¹
RTPnIAA30R	TTCATGGCTTCTCTCTTTCGTAG		
RTPnIAA30IF	ATGAAAGGATCAGAAGCCATTG	<i>PnIAA30</i> fragment spanning between adjacent exons	200 ¹ , 418 ²
RTPnIAA30IR	GTAGAGGCTTGGAGAACACAGG		
IAA30_probeF	CGACGCATGCATCTGAATTA	<i>PnIAA30</i> fragment used as a probe in in situ hybridization experiments	503 ^{1,2}
IAA30_probeR	TGCATGGAACAGTTAACACCA		
β -tubulinF	GTGGAGTGGATCCCCAACAA	β -tubulin fragment used in qRT-PCR assays as normalizer	157 ^{1,2}
β -tubulinR	AAAGCCTTCCTCCTGAACATGG		
GPDHF	CATCAGAGATGAGAAAGTCAAGGTT	GDPH fragment spanning between adjacent exons used in qRT-PCR assays as normalizer	194 ¹ , 520 ²
GPDHR	AGCTTTAAGAATGAAAGGAACACCT		

¹using cDNA as template, ²using genomic DNA as template