

Table S1. Primers used for RT-PCR analyses

Gene	Forward	Reverse
<i>Lj1CPrx</i>	GCAGGAACATGGATGAAGTGC	TTTGAGGCCTTCTGCAGTGAC
<i>Lj2CPrxA</i>	TATTTTAAGCGGCAGGGTCT	TCCAAAATCCAGGTTACACCA
<i>Lj2CPrxB</i>	TGCGCATCTGTATAATTGGCG	CCTGCGGCTCAAATTTTAAGC
<i>LjPrxQ1</i>	TACCAAACAGGCTTGCCT	CCTCAGCTCCTGCTTTCTTGA
<i>LjPrxIIB</i>	ACCCCCAAACTGTCTCCATT	TGATGACCTTTTTGCCGGC
<i>LjPrxIIE</i>	TGAGGTGCTGCTATTGTCCGA	CCCAATGGCTCTGGTGAAATC
<i>LjPrxIIF</i>	GTGTGATGAAGCGAGCTGGTT	CCTGATTCTGAAGAGCAGCAGA
<i>LjTrxh1</i>	CATGCTGAAGCTCCTGGAA	TTTCGCAAACACCATAGCTG
<i>LjTrxh3</i>	ACAAAGTGGTGGGAGCAAAG	CACTGCGTCTCTGGAGCTAA
<i>LjTrxh4</i>	GCTTGAGAAGAAGGTTGAGCA	CAGCATCATCAAATATGTCCAA
<i>LjTrxh6</i>	AGACAAAGTTGCTGGCAAGG	GGCCATGTGCAAGATCATACT
<i>LjTrxh8</i>	GCTGGAGAAGAAGGTTGTCTG	AGGAGAAATCCCACAACAGC
<i>LjTrxh9</i>	TTTGTTCACTCCACTCATGC	AAAGACTTTTGACCCGGATG
<i>LjTrxf</i>	CGCTGCCATAGAAACTGTCC	CAAAGGGTGTGTCAAGCAT
<i>LjTrxm1</i>	CCCAAGTCCACTTTGACTGC	TCGTGAGATAATCCTTGGTTCA
<i>LjTrxm2</i>	ATCGGTGCTGTGCCTAAAAC	GGCTTAATGGCAGGAAGCTA
<i>LjTrxm4</i>	GTCAACATTGACCTCGAGCA	GCAATTTTCCTTCGCAACAT
<i>LjTrxx</i>	TTGGATGCTTTATTGGAATCG	AAACTGTAAAAGCCTGTAAACTGC
<i>LjTrxy</i>	GAACGCATAGAAACCAGTCTCA	TCGAACATTCCAATAGACAAGC
<i>LjTrxz</i>	CCGAAGGACTCATCCCAATA	CAGGATCCAGGGACTTCAAC
<i>LjTrxo</i>	AGCTTGTAGGCGCTGATGTT	CAGCCATTCCCAAAGTCAGT
<i>LjNTRA</i>	GCAGCGGTAGACTTGCATTATT	CATGTACGTGTGGTGTCGATTG
<i>LjNTRB</i>	CATGGCAGCCTTGGATGTAGA	ATTCATTACCCCTACGCTCG
<i>LjNTRC</i>	AGGCTACCCCGAGAAGATGAA	GAGAGGTGATGCACCATCACA
<i>LjNTRA (L)^a</i>	CTACTGGAAAAAGAGTCAAAGG	TGTGAACGGCAATCTCTT
<i>LjNTRA (LS)^a</i>	AAGAGATTGCCGTTTCAACA	CAGTGAGTGCTGCCTCTAA
<i>LjNTRB (L)^a</i>	TACTGGAAAACAGAAATAGTTGAA	GGATTCTTCATCACCTTCTC
<i>LjNTRB (LS)^a</i>	GAGAAGGTGATGAAGAATCC	CAGAAGTAGCTTTTCACTATCTTATG
<i>LjFTRB</i>	AATTGCCCATGTGTTCCAAT	GGTGATAGCCTGTTCCCTGC

^a Forward and reverse primers (5'→3') used for conventional RT-PCR analyses (*L*, long mRNA; *LS*, long plus short mRNAs). All other primers (5'→3') were designed for qRT-PCR.