

Table S1 SPECIFIC PRIMERS SEQUENCES

Sequence of the primers employed for PCR amplification in this study. The table includes oligonucleotide sequences employed for cloning (lower case letters indicate the *SalI* and *NotI* restriction sites, respectively) and site-directed mutagenesis

Primer nomenclature	5'-3' oligonucleotide sequence
<i>BdPepR2</i> (for cloning)	
BdPepR2 forward	GAATTCCCGGgtcgacAATGCCTGGGCGCCTATCG
BdPepR2 reverse	AGTCACGATgcgggccgcTTACTTGTCCATTCTCACCAAGACA
<i>BdTTM3</i> (for site-directed mutagenesis)	
BdPepR2 M1066A forward	TGCAGAATCCACAGGCGCCTTCCTGCACAGGAGC
BdPepR2 M1066A reverse	GCTCCTGTGCAGGAAGGCGCCTGTGGATTCTGCA
BdPepR2 M1066R forward	ATCCACAGGCCTCTTCCTGCACAGGAGCTC
BdPepR2 M1066R reverse	GAGCTCCTGTGCAGGAAGAGGCCTGTGGAT