Table S1. Primers used in RT-PCR analysis and vector construction

PRIMERS	SEQUENCE
For qRT-PCR	
RhMED15a-RT-Up	ACCCACTGTTGTGCCGTGAT
RhMED15a-RT-Dn	ACCGTAATAAGCAAATCGAAAGC
RhMED26B-RT-Up	TTCCGGGATTTGAATTTGATTG
RhMED26B-RT-Dn	CAGACTTGGCCGTGGTCG
RhDREB1B-RT-Up	CAAGGAGACGAGGCACCCGGTGTACCG
RhDREB1B-RT-Dn	ATGCAATCCGTACTTCTCAAAAGAAGTTAT
RhUBI2-RT-Up	GCCCTGGTGCGTTCCCAACTG
RhUBI2-RT-Dn	CCTGCGTGTCTGTCCGCATTG
For the promoter cloning	
RhMED15A-CDS-Up	AAAACCGACTTGATTCCTACGC
RhMED15A-CDS-Dn	AGACAGACGGACCGAGCAG
RhMED26B-CDS-UP	TTGTTCTCCGGTGAAGTTTG
RhMED26B-CDS-Dn	ACTACAGGATGGTCTTTGCATTT
For vector construction of subcellular localization	
pCambia2300-GFP-*RhMED15a*-Up	tacaagggtacccggggatccATGGACACGAATAATTGGAGGC
pCambia2300-GFP-*RhMED15a*-Dn	cttgcatgcctgcaggtcgacTCAAGCGGCACTCAGGCA
PCambia2300-Up	CCACCCACGAGGAGCA
PCambia2300-Dn	TGTGGAATTGTGAGCGGAT
For vector construction of VIGS	
pTRV2-*RhMED15a*-Up	gtgagtaaggttaccgaattcTTGGTACTCAAACTGGTAACTCAAGC
pTRV2-*RhMED15a*-Dn	cgtgagctcggtaccggatccCGTTGATGACGTCTCAGGAAGG
pTRV2-Up	TGGGAGATGATACGCTGTT
pTRV2-Dn	CCTAAAACTTCAGACACGGAT
pTRV1-Up	TTACAGGTTATTTGGGCTAG
pTRV1-Dn	CCGGGTTCAATTCCTTATC