

Table S5. Primers used for molecular cloning and qRT-PCR in this study

Primer pair	Sequence(5'-3')	Cutting sites	Description or purpose
Primers used for molecular cloning			
hrpB-F/hrpB-R	TACGAATTCGCCGTGCAGGCCGGCA CCTGCAGCGCGCTCGGTACAGTT	<i>Eco</i> RI- <i>Pst</i> I	A 696-bp <i>hrpB</i> gene cloned in pK19mpbGII
TRV-SIWAKL20-F/ TRV-SIWAKL20-F	GCGAATTCCTTAGAGTTCTTGTCATCTGTGGG GCGGATCCACAATGTGTTCTTACAGTTCCTTCC	<i>Eco</i> RI- <i>Bam</i> HI	A 300-bp fragment of <i>SIWAKL20</i> cloned in TRV2
TRV-NbWAKL20-F/ TRV-NbWAKL20-R	CGGAATTCGTGATTTATCTTTAGGTGGC/ TCGGATCCATGCACTAATGGAGTCTTCAG	<i>Eco</i> RI- <i>Bam</i> HI	A 305-bp fragment of <i>NbWAKL20</i> cloned in TRV2
Primers used for PCR analysis			
Gene	Sequence(5'-3')	Description	
gusA-F/gusA-R	TCCTGTAGAAACCCCAACCC; CAGTTCATAGAGATAACCTTCACCC	748-bp of partial <i>GUS</i> gene	
qRT-ripAA	GACAAGCGGCTGGGAATACA; GTGCTGGTCGGGATAAACAT	91-bp	
qRT-ripP1	CTGGAGATGTCGGCGGTG; GCGCTTGGCGTCTTCATAG	138-bp	
qRT-ripI	CAGTTGCGCCTCCATGAGT; GCTCGGTCTTCAGGTTCCA	91-bp	
qRT-gyrA	CGACTGGAACCGTCCCTAC; TCCGCACGATGGTGTGATA	104-bp	
qRT-SIWAKL20	AACATATCTTCGCGTAATACGGTCA;CTTCATTAGCTTCCTCGAATCGCCT	92-bp	
qRT-SlActin	ATGTTGCTATTCAGGCGGTTTTGTC;CAGGATTTTCATCAGACAGTCAGTA	182-bp	
qRT-NbEF1 $\alpha$	TGGTGTCTCAAGCCTGGTATGGTTG;ACGCTTGAGATCCTTAACCGCAACATTCTT	160-bp	
qRT-NbWAKL20	CTTAGAGTTCTTGTCCTCCGTAGGA;GTATGGAAGGAGGGCTAATAACAAG	92-bp	