

Table S1. Primers used in this study

Primer	Sequence(5' → 3')	Description
Fgwc1-5F	TGCTCCTCATTTGAATAATCA	
Fgwc1-5R	gcacaggtaacttgttagagAAGATGTTGAGCGGGTGTAT	Forward and reverse primers for amplification of the 5'-flanking region of <i>FgWc-1</i> with a tail for geneticin resistance gene cassette fusion
Fgwc1-3F	cctcaatatacatttctgtcgGCGCTTAGGGCTCAGA	
Fgwc1-3R	CCGATAATGCAGCAACTACAGT	
Fgwc1-5N	GTTGTGGCGATTATTTTATCAATGTAC	
Fgwc1-3N	CGCATATCGTCGGGAATCGTTAG	
Fgwc2-5F	TCGAATGGGTAGACTGAGGAC	
Fgwc2-5R	gcacaggtaacttgttagagGTGGCAGCGACAACGATAGGAG	Forward and reverse primers for amplification of the 5'-flanking region of <i>FgWc-2</i> with a tail for geneticin resistance gene cassette fusion
Fgwc2-3F	cctcaatatacatttctgtcgATGGCGCGATACGGCAGGATT	
Fgwc2-3R	GCGTTGATGAGCTTATCCAT	
Fgwc2-5N	TGAGGCACCATCATGACAAGTATT	
Fgwc2-3N	CTGCGCCTATTGTCTCAAGTACATCA	
FgFphA-5F	CGACGCAGACACAGAACTGA	
FgFphA-5R	gcacaggtaacttgttagagCTTGGTCGCGGTCGTAGAGTAAT	Forward and reverse primers for amplification of the 5'-flanking region of <i>FgFph</i> with a tail for geneticin resistance gene cassette fusion
FgFphA-3F	cctcaatatacatttctgtcgCCGACTATTCTTTGATTTCTG	
FgFphA-3R	AGGGGAGCCTAGGAAACAAAT	
FgFphA-5N	CCGGGTACTCCAGTGCTGACATT	
FgFphA-3N	GCGTTGAACGTTGAGGTAGGC	
Gen-F	CGACAGAAGATGATATTGAAGG	
Gen-R	CTCTAAACAAGTGTACCTGTG	
Fgwc1-rev5	GGCGACAGCATGGCTGGA	Reverse primers for amplification of the 5'-flanking region of <i>FgWc-1</i>
AnLreA-F	tccagecagatgtcgccATGCCAATCGAGATATCAACGA	Forward and reverse primers for amplification of whole <i>LreA</i> gene
AnLreA-R	aatgaateacggcaacgcTCAACCCTCACCGCCAGACTTG	
Fgwc1-for3	GCGTGGCGTGATTCAATT	Forward primer for amplification of the 3'-flanking region of <i>FgWc-1</i>
Fgwc2-rev5	GGTTATCAAGGTTACGTTGCG	reverse primers for amplification of the 5'-flanking region of <i>FgWc-2</i>
AnLreB-F	caaacgttaaccttgataaccATGGATCCCACCCACCTCCAAC	Forward and reverse primers for amplification of whole <i>LreB</i> gene
AnLreB-R	gcacatttgcactgtcgagtTCATAAGCCGAATTTGCCGTT	
Fgwc2-for3	ACTCGACAGTCGCAAATGTGCCAA	Forward primer for amplification of the 3'-flanking region of <i>FgWc-2</i>
Gen-RevN	CCGCTTGGGTGGAGAGGCTATT	
Gen-ForN	CCACAGTCGATGAATCCAGAAAAG	Screening primer for deletion mutants
Fgwc1-SF	GTGCAAACCTGCCATACAAGGAACA	
Fgwc1-SR	ATGTAATGGCGACGAGGGAGAAT	Forward and reverse primers for amplification of ~ 0.2 kb region of <i>FgWc-1</i>
Fgwc2-SF	AGACCCACGAACAGGGATAA	
Fgwc2-SR	GTGGCTAGTGGCCTCTTCTC	Forward and reverse primers for amplification of ~ 0.2 kb region of <i>FgWc-2</i>

FgFphA-SF	AGGAGAGTATTGAGACGCTTGAAGTC	
FgFphA-SR	GCGAGTTCGACGTTGTTCTGATG	Forward and reverse primers for amplification of ~ 0.2 kb region of <i>FgFph</i>
AnLreA-SF	CGCAATTGTCCGCTTCAGAG	
AnLreA-SR	GCGCACCGGGATCATCGTCACCA	Forward and reverse primers for amplification of ~ 0.2 kb region of <i>LreA</i>
AnLreB-SF	TCAGTGCAGACGGAGTAGTTCATT	
AnLreB-SR	CCTGGTACATCCGGAAGTGACAAT	Forward and reverse primers for amplification of ~ 0.2 kb region of <i>LreB</i>
EF1-PS1	GGCTTCAACCGACTACCCTCCTCT	
EF1-PS2	ACTTCTCGACGGCCTTGATGACAC	Forward and reverse primers for qRT-PCR of <i>EF1</i> gene
clucWC1for	gtcccgggcggtacc GATGGCTTCTACTCCCCCAT	
clucWC1rev	tggatccccgggtacc TCAAGATTGGCTTGTCTCGCG	Forward and reverse primers for amplification of <i>FgWc-1</i> , which was cloned into pFCLuc
nlucWC2for	acgagatctggtcgac ATGTCTCACGGACCTCCTCGCC	
nlucWC2rev	agctcgagtagtcgac GGCGACCGATTGATCCACTGTCG	Forward and reverse primers for amplification of <i>FgWc-2</i> , which was cloned into pFNLuc
nlucFphAfor	acgagatctggtcgac ATGAGCCA CTCCTATCTGC	Forward and reverse primers for amplification of <i>FgFph</i> , which was cloned into pFNLuc
nlucFphArev	agctcgagtagtcgac GTCATTGTCGGCTTGGCTT	
clucFphA/for	gtcccgggcggtacc AGCCACTCCTATCTGCGAG	
clucFphA/rev	tggatccccgggtacc TCAGTCATTGTCGGGCTTGG	Forward and reverse primers for amplification of <i>FgFph</i> , which was cloned into pFCLuc