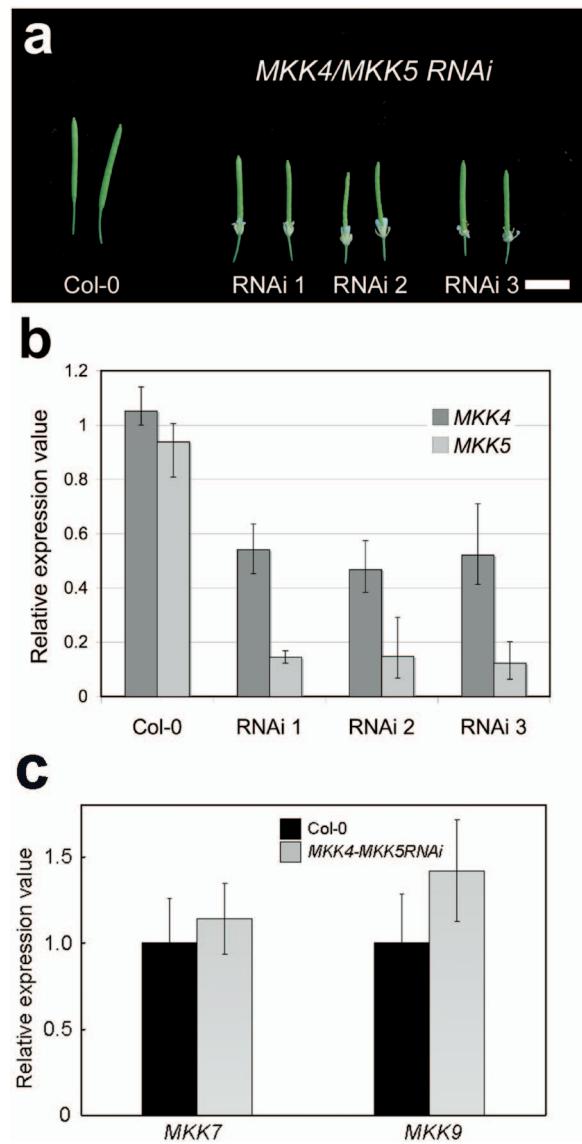
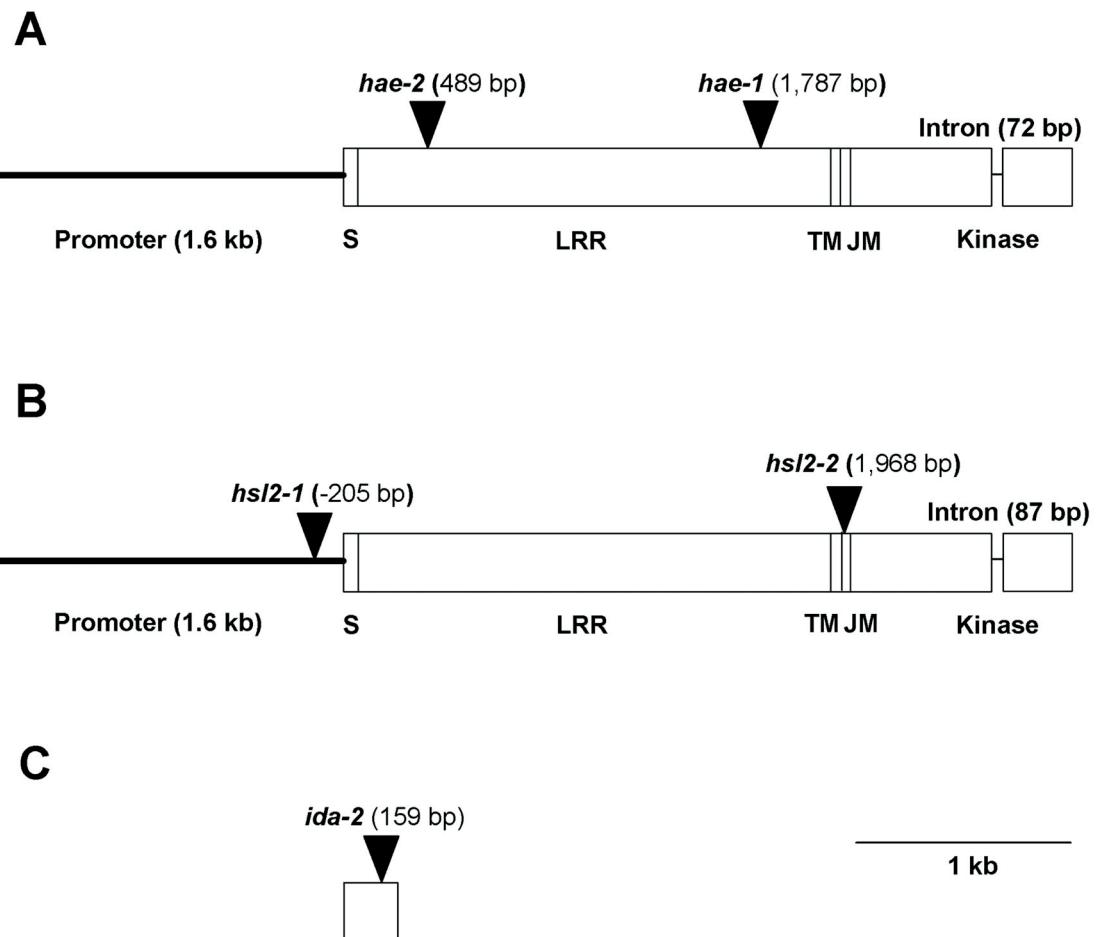


# Supporting Information

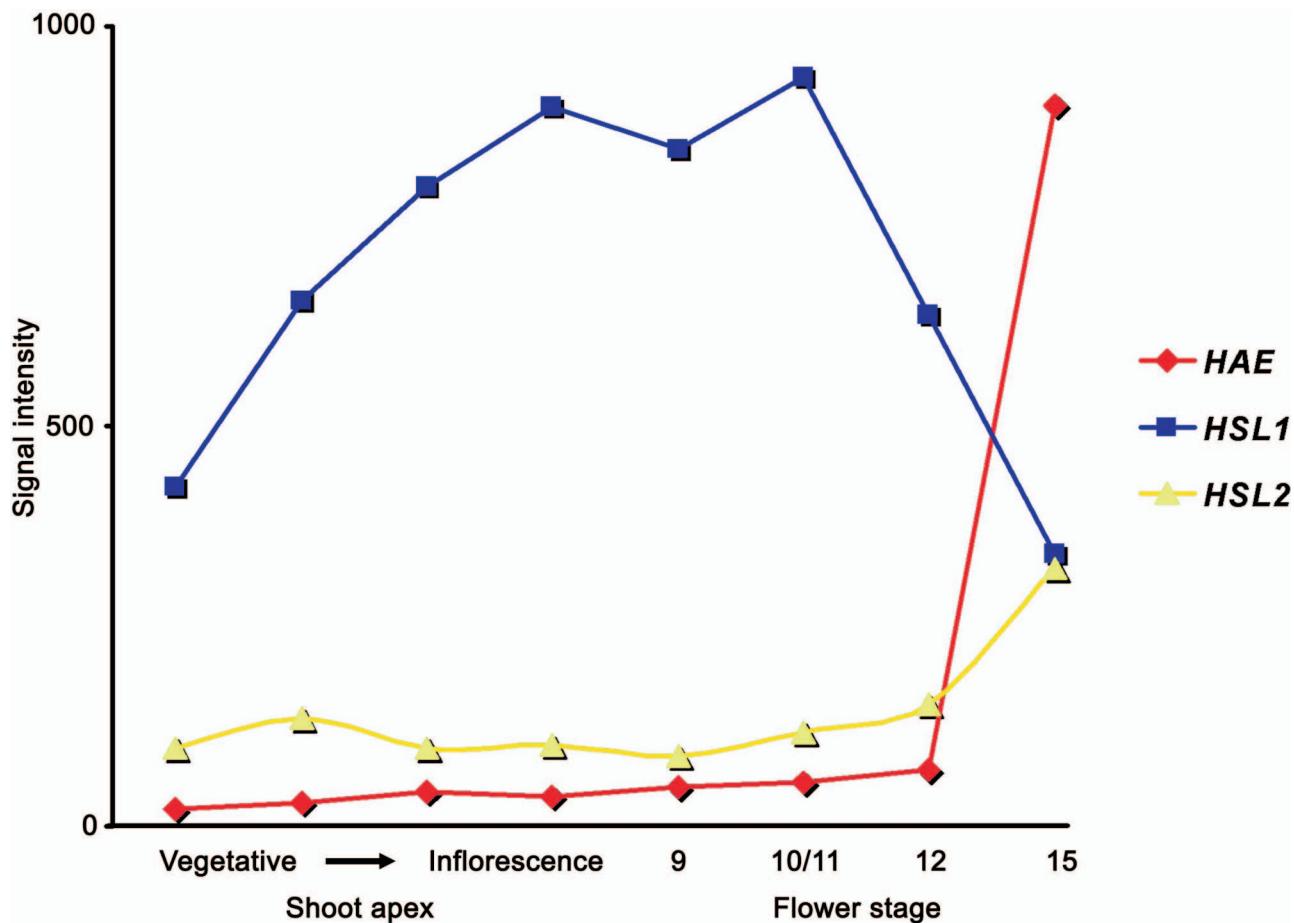
Cho et al. 10.1073/pnas.0805539105



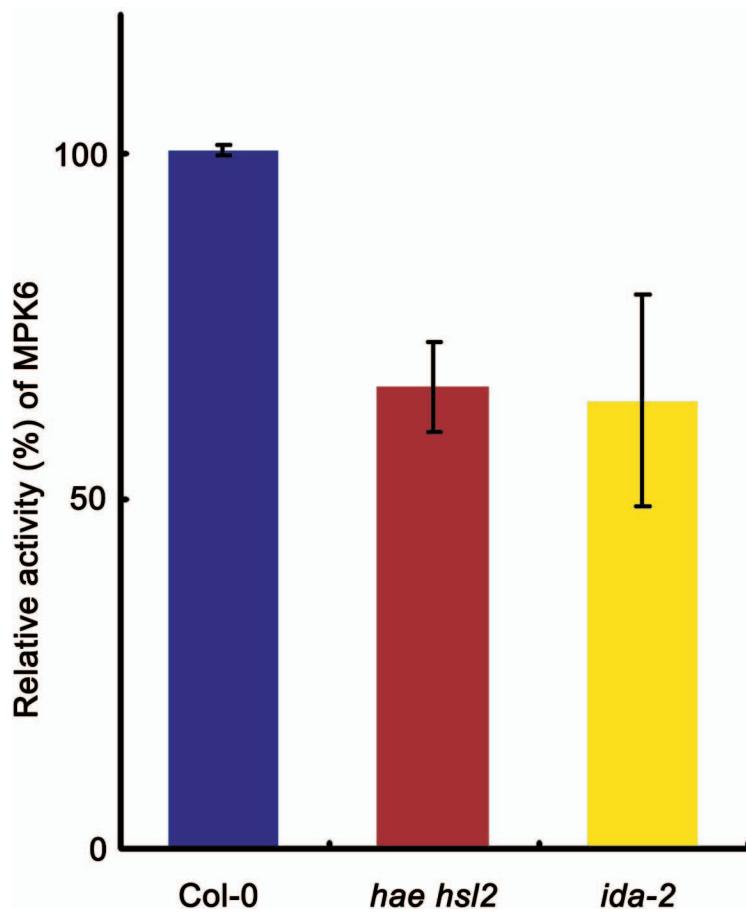
**Fig. S1.** *MKK4* and *MKK5* regulate the floral organ abscission. (a) Three independent lines of *MKK4-MKK5RNAi* transgenic plants show an abscission defective phenotype. (b) Expression of *MKK4* and *MKK5*. RNA was isolated from unopened flower bud clusters of Col-0 and *MKK4-MKK5RNAi* plants. Three biological replicas showed significantly reduced *MKK4* and *MKK5* mRNA accumulation from flowers of the *MKK4-MKK5RNAi* plants. Solid bars indicate the mean and error bars indicate the data ranges ( $n = 3$ ). (c) Expression of *MKK7* and *MKK9* from flowers of Col-0 and *MKK4-MKK5RNAi* plants. Solid bars indicate the mean, and error bars indicate the data ranges ( $n = 4$ ). (Scale bar: 1 cm.)



**Fig. S2.** T-DNA insertion alleles. (A) The *hae-1* T-DNA insertion (SALK\_105975) is located at nucleotide 1787 from start codon. The *hae-2* T-DNA insertion (SALK\_015074), is located at nucleotide 489 from start codon. The *hae-1* allele was used in this study. (B) The T-DNA of the *hsl2-1* allele (SALK\_057117) is located at nucleotide -205, upstream of start codon. The T-DNA of *hsl2-2* (SALK\_030520) is located at nucleotide 1968 from start codon. The *hsl2-1* allele was used in this study. (C) *ida-2* (SALK\_133209) has a T-DNA insertion 159 nucleotides from start codon. The bold line showed promoter region that used for GUS transcriptional fusion, and the thin line in the kinase domain indicates an intron. (S) signal peptide; LRR, leucine rich repeat; TM, transmembrane domain; JM, juxtamembrane region. (Scale bar: 1 kb.)



**Fig. S3.** Expression of *HAE*, *HSL1*, and *HSL2*. Expression pattern of *HAE*, *HSL1*, and *HSL2*. Publicly available microarray data were compiled using AtGenExpress ([www.weigelworld.org/resources/microarray/AtGenExpress](http://www.weigelworld.org/resources/microarray/AtGenExpress)).



**Fig. S4.** MPK6 is activated in receptacle. Relative MPK6 activity from the receptacle of Col-0, *hae hsl2*, and *ida-2* plants. Biological replications ( $n = 3$ ) show the reduced activity of MPK6 from receptacle of the mutants. Error bars indicate SE for the different mutants.

**Table S1. List of primers**

Method used		Primer	Sequence
Genotyping	1	hae-1-F	5'-CAAACTGGGTTATCCGACCCGG-3'
		hae-1-R	5'-GTCCCCAACGTTCAACCTCGCTG-3'
	2	hae-2-F	5'-ATGGGAGAATCTGAATTACTGAG-3'
		hae-2-R	5'-CGAGAAGTGACAAGCGAGGC-3'
	3	hsl2-1-F	5'-CCCACTCGGAAGCTTACCATAGTCTG-3'
		hsl2-1-R	5'-CGGGAATCACACCGGAGAAGTTGTTAGC-3'
	4	hsl2-2-F	5'-GTCTCGACGACAGATTCTCG-3'
		hsl2-2-R	5'-TCCATTGCAAACCTAACATGTCC-3'
	5	ida-2-F	5'-GTTTTGATCAGGAGAGAGTTG-3'
		ida-2-R	5'-CTTCTCACGCCAAAAGATAAGAGTTGG-3'
MPK6	1	MPK6-F	5'-TCTGAATTGTCGGTTACAGAGATCTCACAGA-3'
		MPK6-B	5'-TCTCCGGGCATGACCGGTAAGAGATGAAAGCTT-3'
mutagenesis	2	cMPK6-F	5'-TCTCCGGGATGGACGGTGGTCAGGTCAACCG-3'
		cMPK6-Noterminator-B	5'-CTCTCTAGAAATTCCGATCTAGAACATAGAT-3'
35S::IDA		IDAOE-F	5'-ATGGCTCCGTGTCGTACGATG-3'
		IDAOE-R	5'-TCAATGAGGAAGAGAGTTAACAAAAGAG-3'
HAE::GUS		GUS-HAE-F	5'-CAGAAAGGATTAACAACTAAACTCTGCACC-3'
		GUS-HAE-R	5'-TTTTTGAAAGGAATCGTTATTCTC-3'
HSL2::GUS		GUS-HSL2-F	5'-ACTTATATTGTGAAACAAAAAATTGGTGC-3'
		GUS-HSL2-R	5'-CGTGTGGAAAGAGAGGTATGAAAC-3'
qRT-PCR	1	QRT-MKK4-F	5'-GAGGTTTCCTTCCCTGTGA-3'
		QRT-MKK4-R	5'-CTCTCTGCAAGCAACACGAG-3'
	2	QRT-MKK5-F	5'-CGTCGTATCGTTCATCG-3'
		QRT-MKK5-R	5'-CATTGTTGTGCCAAGATCC-3'
	3	QRT-MKK7-F	5'-GTTCGTAAACGCCGTCAAAT-3'
		QRT-MKK7-R	5'-CTTCTCTCCGAGAACGTG-3'
	4	QRT-MKK9-F	5'-CAGTTGATGCGAGAGATGGA-3'
		QRT-MKK9-R	5'-GCGAGTTTGCTCCGTTAC-3'
	5	QRT-EF-1 $\alpha$ -F	5'-TGAGCACGCTTCTGCTTCA-3'
		QRT-EF-1 $\alpha$ -R	5'-GGTGGTGGCATCCATCTGTTACA-3'

qRT-PCR, quantitative real-time PCR.