

SUPPLEMENTARY INFORMATION

T-DNA alleles of the receptor kinase THESEUS1 with opposing effects on cell wall integrity signaling

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Table S1 List of Primers used

Name	Sequence	Specificity
GFP-F	ACATGAAGCAGCACGACTTC	GFP construct
GFP-R	TGTAGTTGACTCCAGCTTGT	GFP construct
THE1-Start	ATGGTGTTCACAAAATCATTAC	ECD-TM ^{THE1} -YFP construct
TM-JTM-Lo	GAACTTATTAGTAGCGTCCAT	ECD-TM ^{THE1} -YFP construct
THE1_3endF	AAGGTGAACCCTGCCTCGT	expression downstream T-DNA insertion
THE1_3endR	ATTGGAATCCCTGGAATGTGG	expression downstream T-DNA insertion
THE1_5endF	GTGGTCTTGTCAACGCTAT	expression upstream T-DNA insertion
THE1_5endR	CATCATTATCCCATTGTCTCC	expression upstream T-DNA insertion
THE1_midF	GCTGATATCACTAATGCGAC	upstream T-DNA insertions, <i>the1-3</i> , <i>the1-4</i> genotyping, expression, antisense cDNA
THE1_midR2	CAGCAATCAGCAGAACATG	upstream T-DNA insertions
LB4	CGTGTGCCAGGTGCCACGGAATAGT	<i>the1-3</i> genotyping; T-DNA specific expression
LB3	CTGAATTCATAACCAATCTCGATACAC	<i>the1-4</i> genotyping; T-DNA specific expression
TUB9-F	GTACCTTGAAGCTTGCTAACCTTA	reference gene qPCR
TUB9-R	GTTCTGGACGTTCATCATCTGTT	reference gene qPCR
UBQ-F	CTCCTTCTTCTGGTGGTAAACGT	reference gene qPCR
UBQ-R	AACCCTGAGGTTGAATCATCC	reference gene qPCR
5g19110/EDGP-F	CTACAATGCTCTGCTCAGTC	<i>THE1</i> target gene, qPCR
5g19110/EDGP-R	ACCGTCGATAAACGCCAAC	<i>THE1</i> target gene, qPCR
2g26530/AR781-F	CATCAGCTTGCCTGTCTTGC	<i>THE1</i> target gene, qPCR
2g26530/AR781-R	CCCTCTCCTCGTGAACTCG	<i>THE1</i> target gene, qPCR

Supplementary Figures

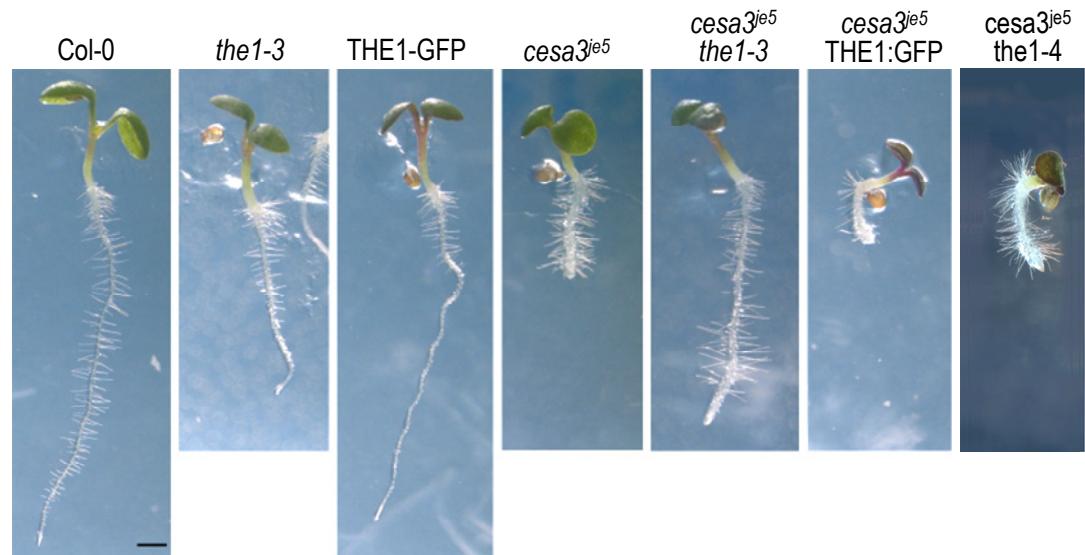


Figure S1 Root growth of *thel-3*, *thel-4* and 35S::THE1:GFP seedlings in cellulose deficient genetic backgrounds. Black bar corresponds to 1 mm.

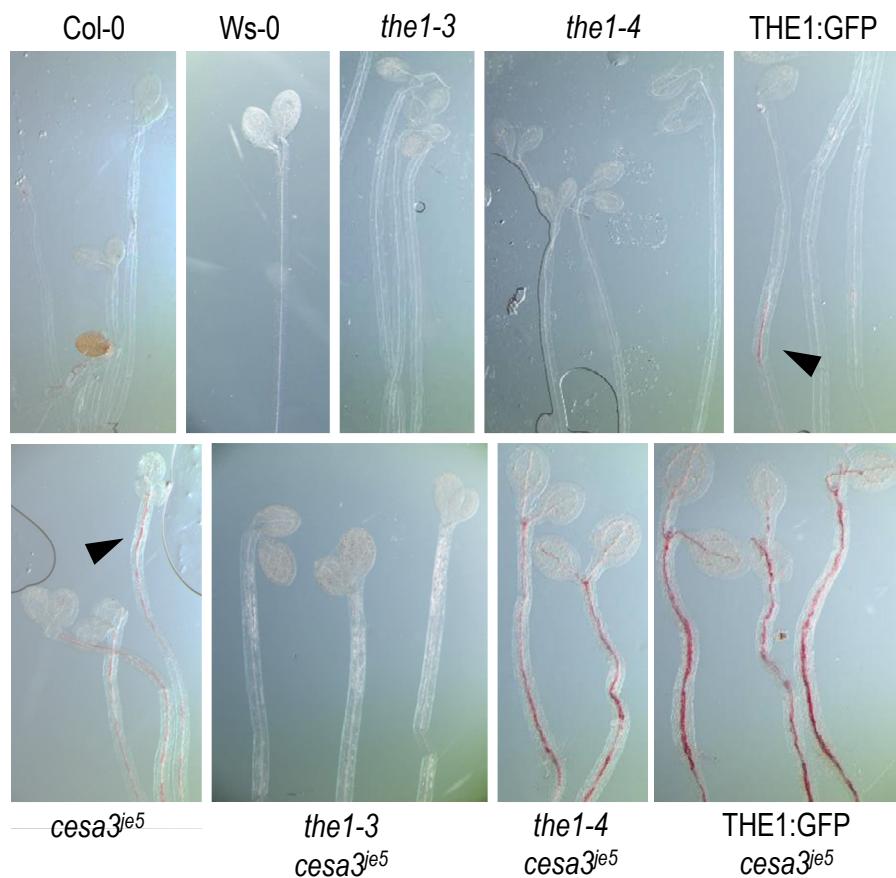


Figure S2 Ectopic lignification of etiolated seedlings of *thel* alleles in cellulose deficient genetic backgrounds.

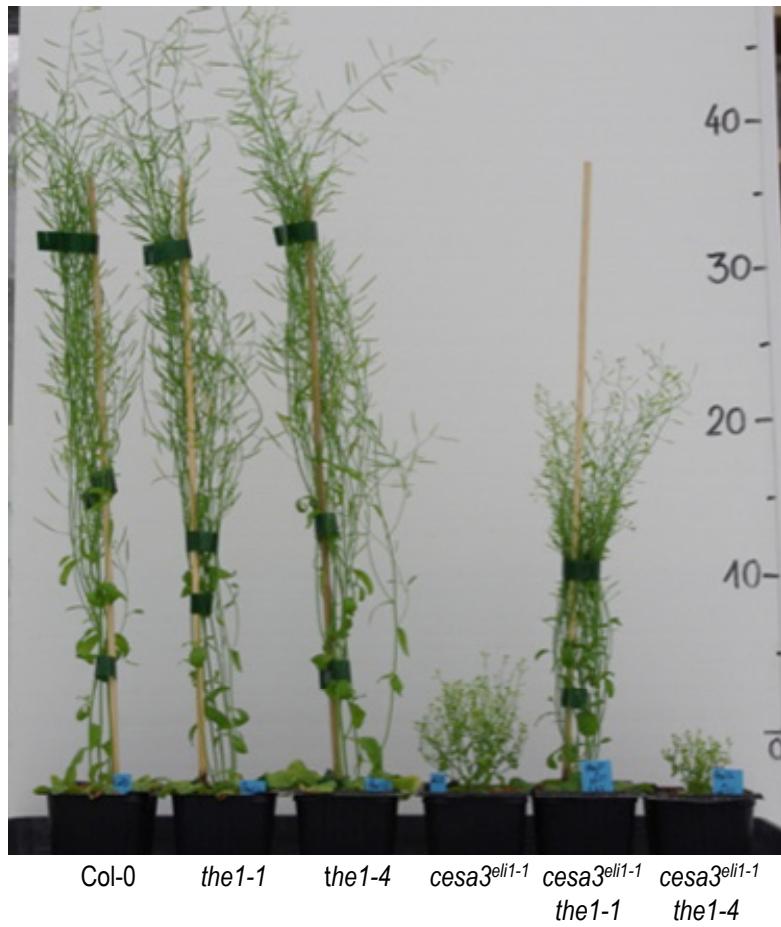


Figure S3 Inflorescence phenotypes of loss- and gain-of-function alleles in combination with cellulose deficient backgrounds.

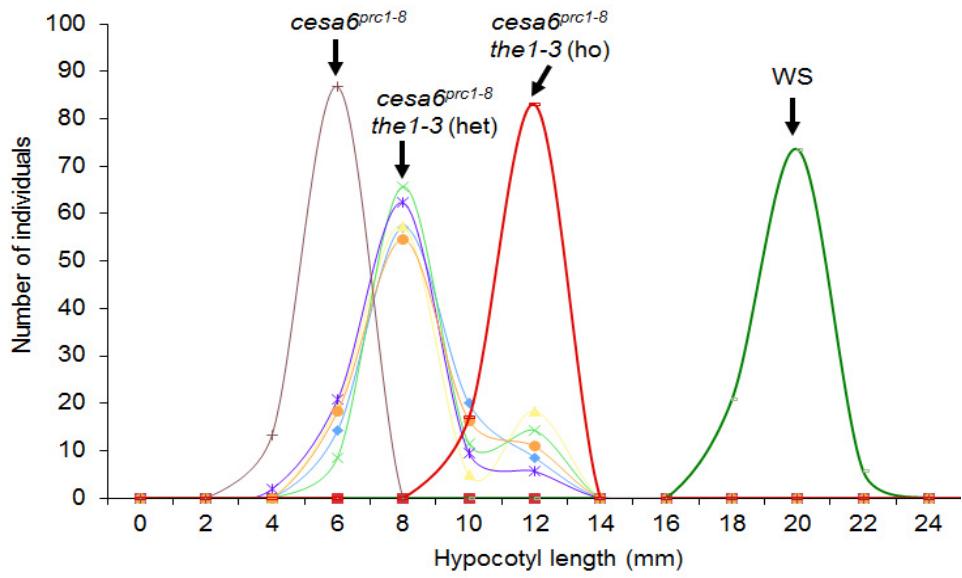


Figure S4 Semidominance of *the1-3* in combination with *cesa6^{prc1-8}*. Seedlings were cultivated on MS medium without sucrose for 7 days in the dark. Data represent frequency distributions of hypocotyl lengths. All seedlings except wild type (WS) were homozygous for *cesa6^{prc1-8}* and either wild type, heterozygous (het) or homozygous (ho) for *the1-3*.