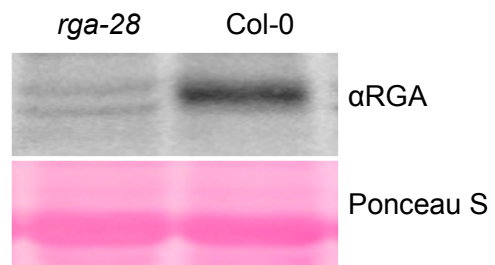
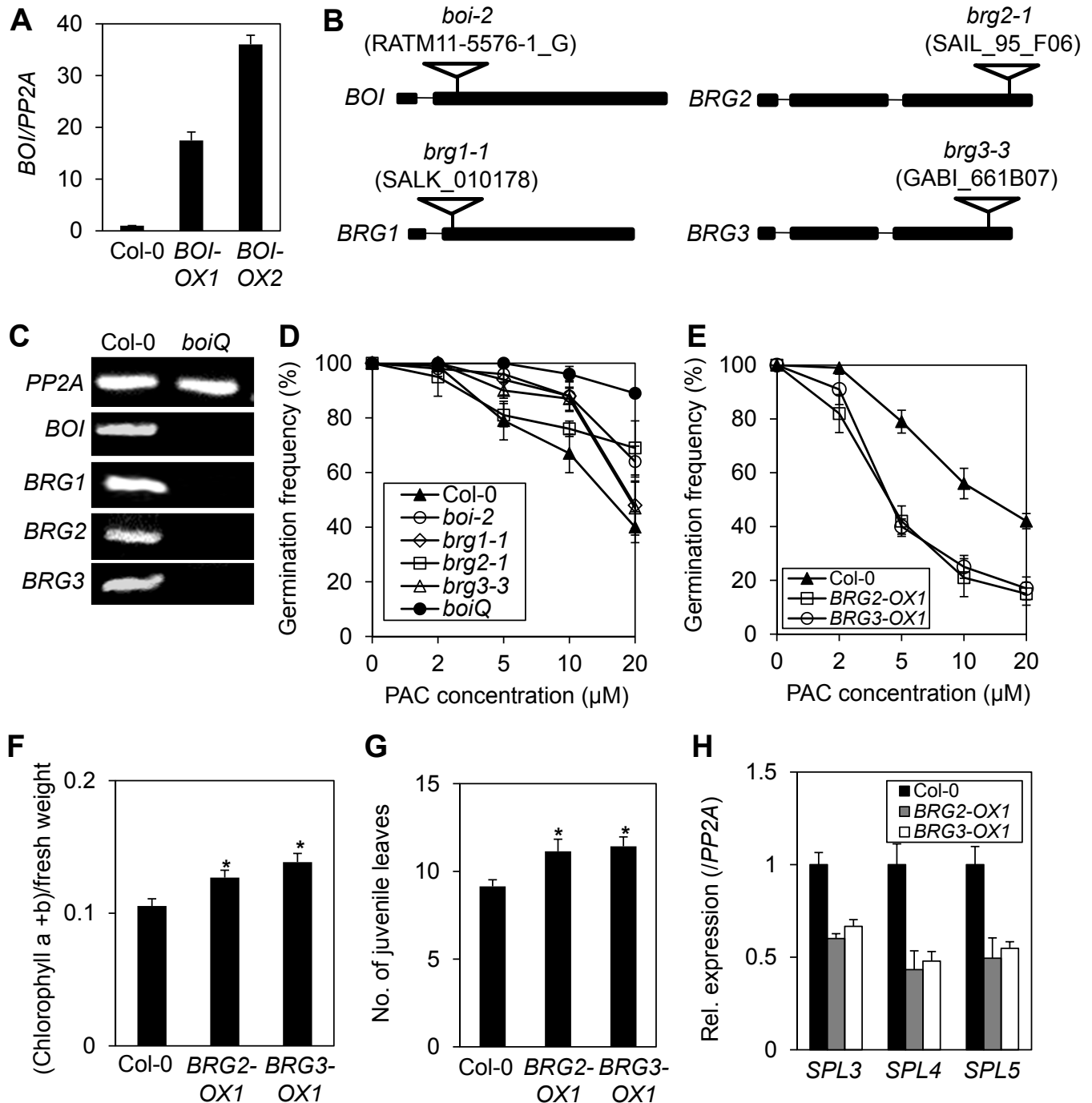


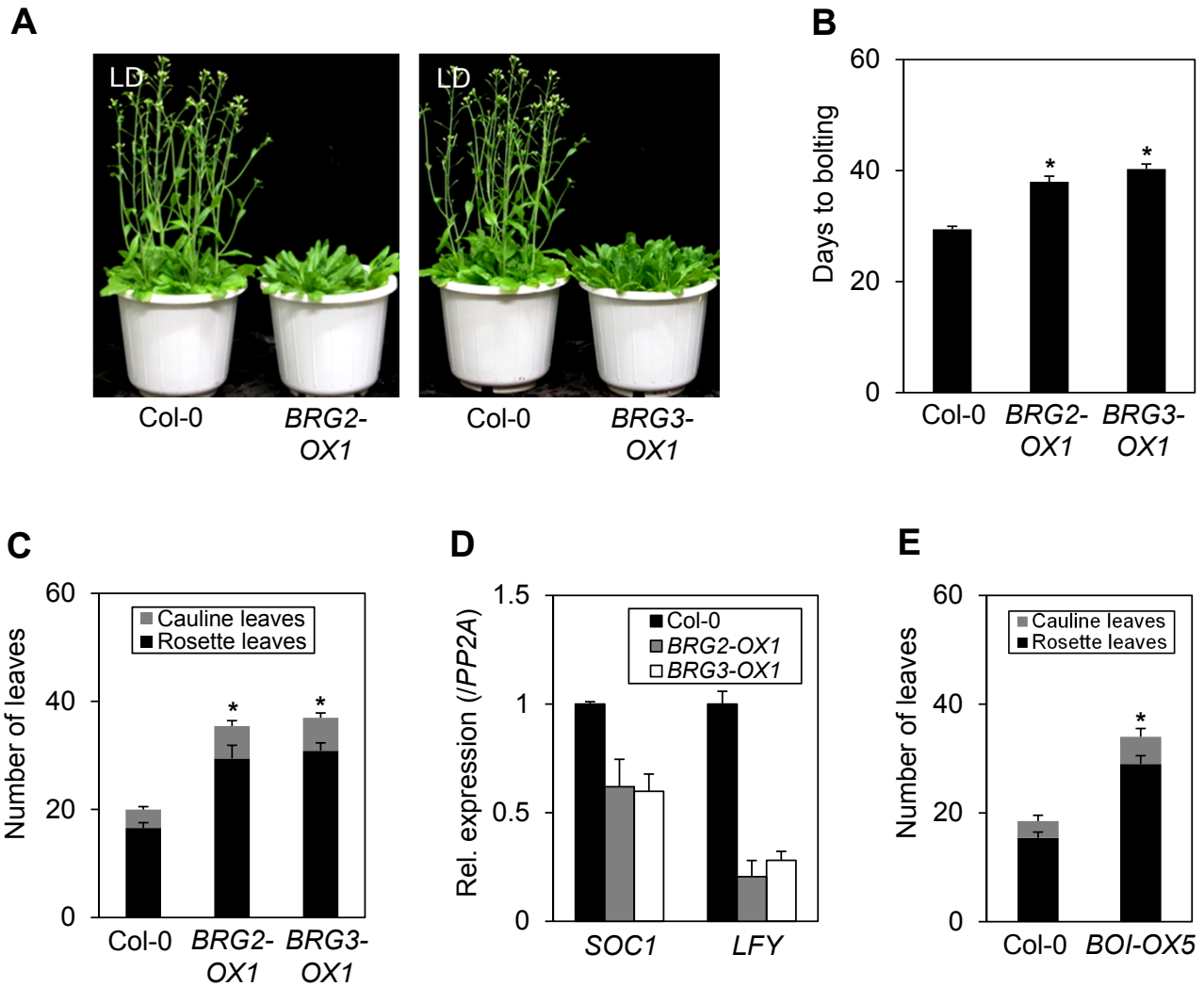
Supplemental Figure 1. Amino acid alignment of the four BOIs (A) and a neighbor-joining tree showing the relationships among the BOIs and their *Arabidopsis* and rice homologs (B). Sequence alignment was generated by Clustal W multiple alignment tool of BioEdit (www.mbio.ncsu.edu/bioedit/bioedit.html) and the phylogenetic tree was generated by MEGA5 program (www.megasoftware.net). The evolutionary distances were computed using the Poisson correction method. The scale bar indicates the evolutionary distance of 0.1 amino acid substitution per position.; * indicates zinc-coordinating Cys or His residues.



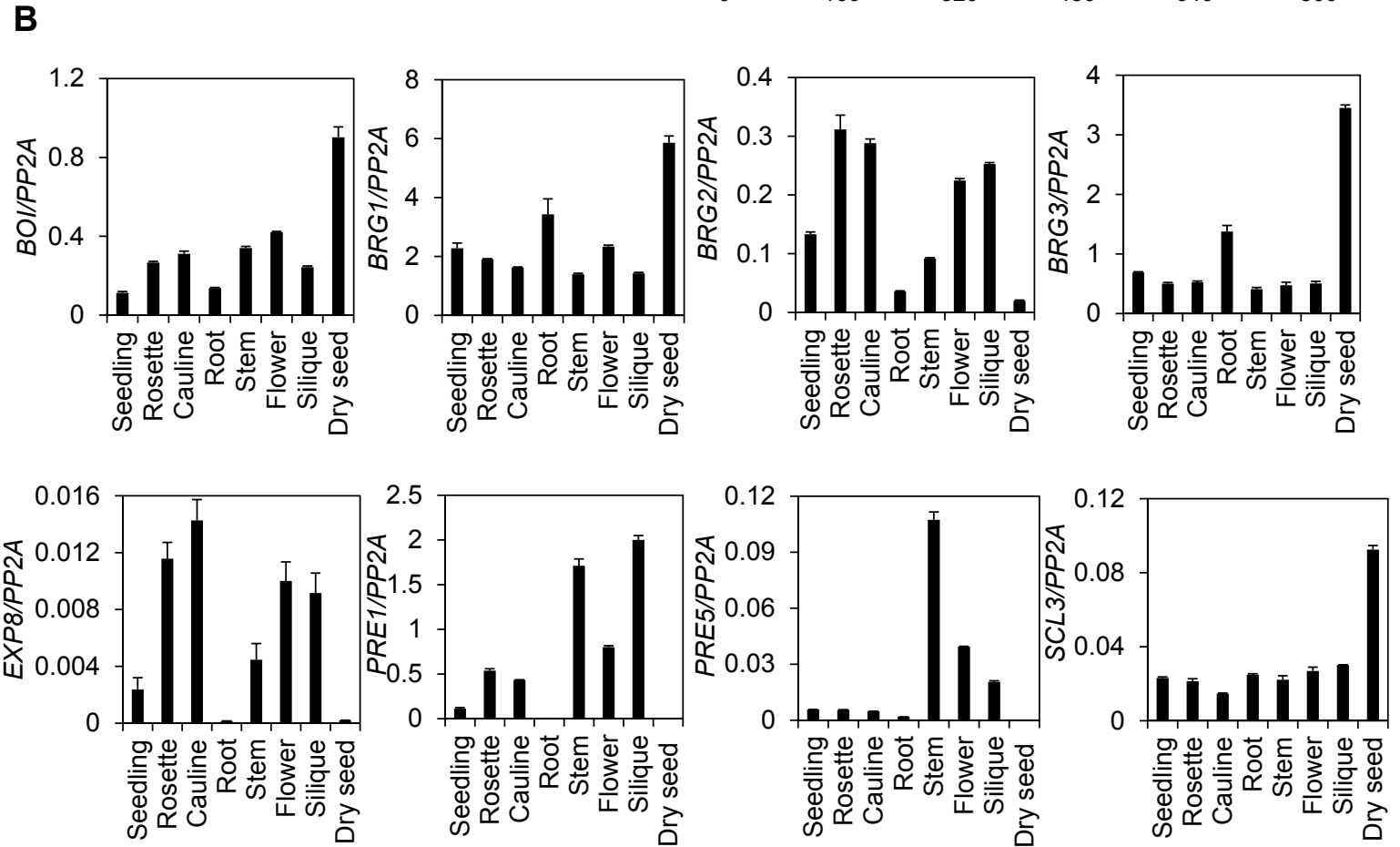
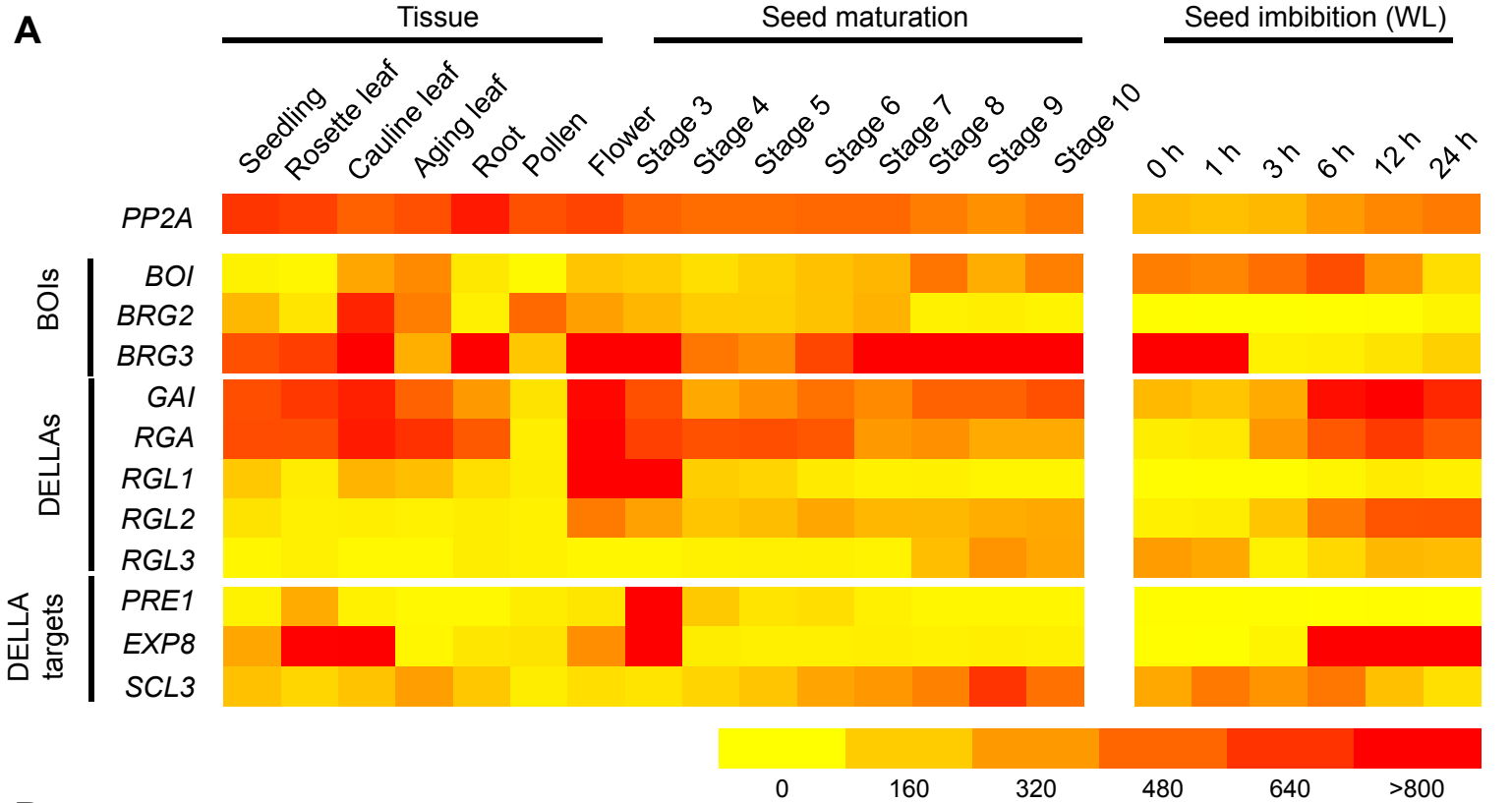
Supplemental Figure 2. The specificity of RGA antibody. 7-day-old white light-grown seedlings of the *rga-28* and Col-0 were used for the analysis of RGA antibody.



Supplemental Figure 3. BOI family members redundantly inhibit GA responses. (A) *BOI* mRNA expression levels in the two *BOI*-overexpressing lines. *BOI* mRNA levels (endogenous + transgenic) were determined in 7-day-old seedlings grown under LD (SD, n=3). (B) T-DNA insertion mutants of *BOI* family members. Exons are indicated by filled rectangles and T-DNAs are by inverted triangles. (C) The full-length *BOI* mRNAs are not detected in the *boi* quadruple mutant (*boiQ*) seedlings, as determined by RT-PCR. (D) Germination frequencies of single and quadruple mutants of *BOI* family members in the presence of PAC (SD, n=3). (E) Decreased germination frequencies of *BRG2*- and *BRG3*-overexpressing lines in the presence of PAC (SD, n=3). (F) Increased chlorophyll accumulation by *BRG2*- and *BRG3*-overexpressing lines (SD, n=6; *, p<0.05, Student's t-test). (G) Delayed juvenile-to-adult phase transition in *BRG2*- and *BRG3*-overexpressing lines (SD, n=10; *, p<0.05, Student's t-test). (H) Decreased expression of *SPL* mRNAs in *BRG2*- and *BRG3*-overexpressing lines. 17-day-old plants were sampled for expression analysis at the 12th hour of light (SD, n=3 biological replicates).

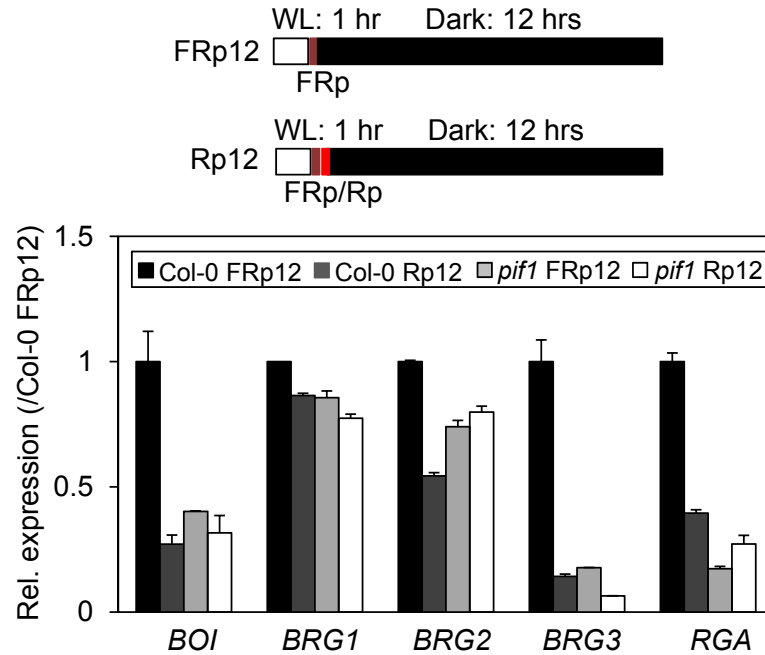


Supplemental Figure 4. Overexpression of *BRGs* inhibit flowering and mRNA expression of *SOC1* and *LFY*. (A) Delayed flowering in *BRG2*- and *BRG3*-overexpressing plants under the LD condition. Pictures were taken at 5 weeks. (B) Quantification of bolting days in *BRG2*- and *BRG3*-overexpressing plants under the LD condition (SD, n=10; *, p<0.05, Student's t-test). (C) Quantification of rosette leaves and cauline leaves in *BRG2*- and *BRG3*-overexpressing plants under the LD condition (SD, n=10; *, p<0.05, Student's t-test). (D) Decreased expression of *SOC1* and *LFY* mRNAs in *BRG2*- and *BRG3*-overexpressing lines under the LD condition. 17-day-old plants were sampled for expression analysis at the 12th hour of light (SD, n=3 biological replicates). (E) Quantification of rosette leaves and cauline leaves in *BOI-OX5* plants under the LD condition (SD, n=10; *, p<0.05, Student's t-test).

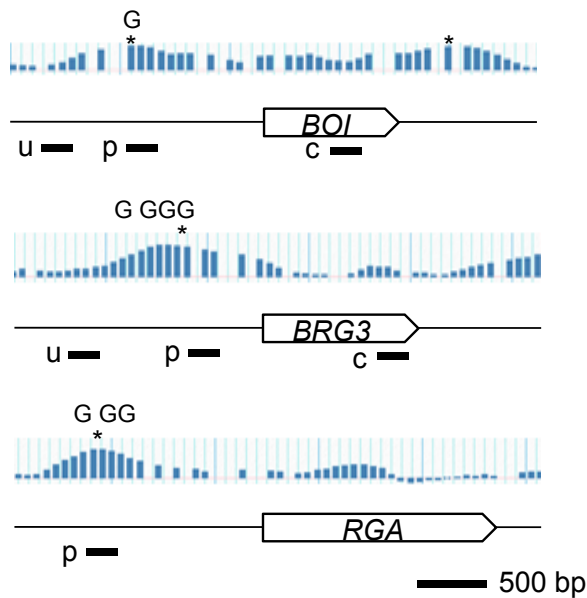


Supplemental Figure 5. Expression patterns of BOIs, DELLAs, and their target genes analyzed by the BAR expression browser and visualized by the BAR heatmapper tool (<http://bar.utoronto.ca>) (A) or by quantitative RT-PCR (SD, n=3) (B).

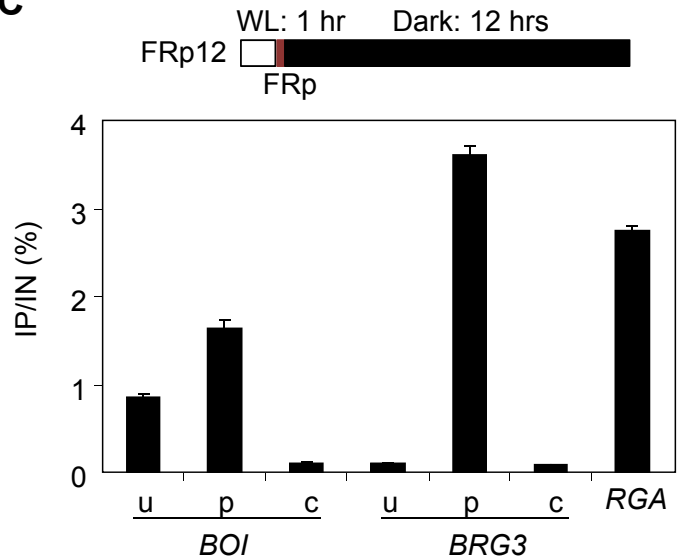
A



B



C



Supplemental Figure 6. Direct regulation of *BOI* and *BRG3* expression by PIF1. (A) Repression of *BOI* and *BRG3* mRNA expression by PIF1 during seed germination. Upper diagrams indicate light irradiation scheme for FRp12 and Rp12. Each mRNA level is presented relative to that of Col-0 FRp12 (SD, n=3). (B) Diagrams showing PIF1 ChIP-Chip signals around *BOI*, *BRG3*, and *RGA*. Vertical blue bars indicate PIF1 signal intensities; * indicates a PIF1 binding site; G indicates G-box elements; and underlines with u, p, and c indicate the amplicons used for ChIP-PCR. (C) Enrichment of *BOI* and *BRG3* promoter fragments by PIF1 ChIP-PCR. The upper diagram indicates light irradiation scheme. The *RGA* promoter was used as a PIF1-binding positive control (SD, n=3).

Supplemental Table 1. Primers used in this study.

Name	Forward primer	Restriction site	Reverse primer	Restriction site	Purposes
<i>pETM_MCS</i>	TATGACTAGTTCTAGAGAAATCCCGGGCCTAGGCTC GAGTCGGATCCCGTCGACA	SpeI-XbaI-EcoRI-XmaI- AvrII-XhoI-BamHI-SalI	TCGATGTCGACGGGATCCGACTCGAGCCTAGGCCCGG GAATCTCTAGAAGACTGCA		Adapter ligation pET29a: NdeI/XhoI
<i>pGADM_MCS</i>	//	//	//		Adapter ligation pGADT7: NdeI/XhoI
<i>pGBKM_MCS</i>	//	//	//		Adapter ligation pGBKT7: NdeI/SalI
<i>pMALM_MCS</i>	AATTGTCTAGAGAATTCGCCGGCCTAGGCTCGAGTC GGATCCCG	XbaI-EcoRI-XmaI-AvrII- XhoI-BamHI-SalI	TCGACGGGATCCGACTCGAGCCTAGGCCCGGGAATTC CTAGAC		Adapter ligation pMAL-c2X: EcoRI/SalI
<i>phNIL_MCS</i>	CTAGAAGTCTAGTCCCGGGCCTAGGCTCGAGTCGGATCC CGTCGACTAATTGATTAAGAGCT	XbaI-SpeI-XmaI-AvrII- XhoI-BamHI-SalI-SacI	CTAATCAATTAGTCGACGGGATCCGACTCGAGCCTAGG CCCGGACTAGTT		Adapter ligation pCAMBIA1300: XbaI/SacI
3XFLAG	GTGT <u>ACTAGT</u> atgGGAGGTGGCAGCGTCGAGATGG	SpeI	GGGTCTAGACCCCCCTCGACTTTATCGTCA	SmaI	phNF construction
<i>BOI</i>	GTGT <u>CCTAGG</u> atgGCTGTTCAGCTCATCACATG	AvrII	GTGTGGATCCGGGAAGACATGTTAACATGCACA	BamHI	Cloning
<i>BRG1</i>	GTGTGT <u>CTAGA</u> atgGCTGTGAAGCAAGACACA	XbaI	GTGTGTGTCGACTGATGACATGTTAACATGTACA	Sall	Cloning
<i>BRG2</i>	GTGTGT <u>CTAGA</u> atgGCCGTCGATGCTCACCATC	XbaI	GTGTGTGTCGACAGATGACATGTTGACATGAACG	Sall	Cloning
<i>BRG3</i>	GTGT <u>GAAATC</u> atgGCCGTTGAAGCTCACCATCTA	EcoRI	GTGTCTCGAGAGAGGAAAGATTAACATGTAGACT	XhoI	Cloning
<i>BOIdC</i>	GTGT <u>CCTAGG</u> atgGCTGTTCAGCTCATCACATG	AvrII	GTGTGGATCCAGCGGTTGGAACGTTTCGAGT	BamHI	Cloning
<i>RGA</i>	GAG <u>CTAGA</u> atgAAGAGAGATCATCACAAT	XbaI	GAGGGATCCCGTACGCCCGCTCGAG AG	BamHI	Cloning
<i>GAI</i>	GAG <u>CTAGA</u> atgAAGAGAGATCATCATCTC	XbaI	GAGGGATCCCGATTGGTGAGAGTTTCCA	BamHI	Cloning
<i>RGL1</i>	GAG <u>CTAGA</u> atgAAGAGAGAGACAACCA	XbaI	GAGGGATCCCGTCCACACGATTGATTG	BamHI	Cloning
<i>RGL2</i>	GAG <u>CCTAGG</u> atgAAGAGAGGATACGGAG	AvrII	GAGAGATCTCCGGCGAGTTTCCACGCCGA	BglII	Cloning
<i>RGL3</i>	GTGT <u>ACTAGT</u> atgAAACGAAGCCATCAAGAAACG	SpeI	GTGTCTCGACCCCGCGCAACTCCGCCGCTAGTTT	Sall	Cloning
<i>boi-2</i>	ATGGCTGTCAAGCTCATCACAT		AGAAGACATGTTAACATGCAC		Genotyping
<i>brg1-1</i>	TAAATAGCATTATGCACGGG		CGCCAGATCTGATTCTCTACG		Genotyping
<i>brg2-1</i>	CGTCTGTCTCCACGCTCTAC		ACGTAAGTTTTGCATCAACGG		Genotyping
<i>brg3-3</i>	AATCCCACCGATCTCTTATG		TCAAGCTTTGAAAAGTGACG		Genotyping
<i>BOI</i>	TGCTCAACTCGAAACGTTTCC		ACCAACACTCGTCTCTCTC		RT-qPCR
<i>BRG1</i>	CGTCGTCCTTTGGTTGAAGAAGC		AGCCGACGATCCACAAACCGTA		RT-qPCR
<i>BRG2</i>	GAACGAAGAGGACGATCGCGA		TCCTCCCCACAGTTTCTACACA		RT-qPCR
<i>BRG3</i>	GAGGCGCAGGATACGAAAAAGATG		CTAAGAGGAAAGATTAACATGTAG		RT-qPCR
<i>SOC1</i>	TCGCCAGCTCCAATATGCAAGATAC		CGATTGAGCATGTTCTATGCGCTTC		RT-qPCR
<i>LFY</i>	TACGCTCTCCACTGCCTAGAC		AGACGGCGTCTATATCCCAGC		RT-qPCR
<i>SPL3</i>	TGAGAAGAAGCAAAGCGGAA		TATCCGCGGTACAACCTCTCG		RT-qPCR
<i>SPL4</i>	GTAGCATCAATCGTGGTGGC		CTTCGCTCATTGTGTCCAGC		RT-qPCR
<i>SPL5</i>	ATGCAGCAGGTTTCATGAGC		GCCTGACCCTTCTCCAAAAAC		RT-qPCR
<i>EXP8</i>	CATGTATGAAGAAAGGAGGAATAAG		AACTGCCAATTAGAAGGAGCCACG		RT-qPCR
<i>PRE1</i>	CAGCCTCGAAAGTATTGCAAG		TTCTAATAACGGCGGCTTCAG		RT-qPCR
<i>PRE5</i>	TCGAATGCTTCGAGGATCTCC		CAGAGTCAAGAAGCTGCGACA		RT-qPCR
<i>SCL3</i>	CAGCTGAGGCACGTGAGAAATGAT		ACCACCATGACCTTTGGAGACAAAAC		RT-qPCR
<i>PP2A</i>	TATCGGATGACGATTCTTCGTGACG		GCTTGGTCGACTATCGGAATGAGAG		RT-qPCR
<i>BOlu</i>	TTGCTACTTGCGAATTACTTGC		TCTTTGACTCGTATCTCGTC		ChIP-qPCR
<i>BOlp</i>	CACGCTTCTCAGTGCAACTTA		GCCATGTCATCTCCACCGT		ChIP-qPCR
<i>BOlc</i>	TGCTCAACTCGAAACGTTTCC		ACCAACACTCGTCTCTCTC		ChIP-qPCR
<i>BRG3u</i>	ATGGAGCTGACTTGATTGGTT G		ATGACCTATGTGAGACTGTGAC		ChIP-qPCR
<i>BRG3p</i>	GTCCTCAACGGATAAGCAAG		GGCTCATGTGACAAGACTTGC		ChIP-qPCR
<i>BRG3c</i>	GAGGCGCAGGATACGAAAAAGATG		CTAAGAGGAAAGATTAACATGTAG		ChIP-qPCR
<i>EXP8</i>	CCCATTCTTCCATTTGTAGATTTC		GGAGGAATAATACGGATATAATTG		ChIP-qPCR
<i>PRE1</i>	CTCTCTTGTGGTCCATTGGCCTA		ATCGGCTATGTCTATGTGGCAAG		ChIP-qPCR
<i>PRE5</i>	TCGGATCAACCACAACTTCTAG		GTGCATGTCACTAAAAGGCAATAC		ChIP-qPCR
<i>SCL3</i>	GCCTCAGCCTCATCTCTTTT		GGAATCATGACTATATATTCTACATCA		ChIP-qPCR
<i>PIL1</i>	ATGAATCACGCGGCATTC		ACGTGAGCGGAAAGAACC		ChIP-qPCR
<i>RGA</i>	GTGACGTAACATGACAATAAACAT		CTAAAACGGTAGGGACCGAGTCTG		ChIP-qPCR

* Underlined are restriction enzyme sites. Start codon (atg) is in lowercase.

Supplemental Table 2. Vectors used in this study.

Name	Transgenic lines and protein name	Plant selection	Usage
phNF-BOI	<i>BOI-OX3, BOI-OX4</i> , FLAG-BOI	Hygromycin	Transgenic, transient
pbFLAG2-BOI	<i>BOI-OX5</i>	Phosphinothricin	Transgenic
pBARH8-BOI	<i>BOI-OX1, BOI-OX2</i>	Phosphinothricin	Transgenic
phNF-BRG2	<i>BRG2-OX1</i>	Hygromycin	Transgenic
phNF-BRG3	<i>BRG3-OX1</i>	Hygromycin	Transgenic
phNF-BOIdC	<i>BOIdC-OX1, BOIdC-OX2</i>	Hygromycin	Transgenic
pbFLAG2-GFP	<i>GFP-OX</i>	Phosphinothricin	Transgenic
phNF-GFP	FLAG-GFP	Hygromycin	Transient
pMALM-RGA	MBP-RGA	n/a	Pull-down
pMALM-GAI	MBP-GAI	n/a	Pull-down
pMALM-RGL2	MBP-RGL2	n/a	Pull-down
pETM-BOI	BOI-His	n/a	Antibody production
pGADM-RGA	AD-RGA	n/a	Yeast two-hybrid
pGADM-GAI	AD-GAI	n/a	Yeast two-hybrid
pGADM-RGL1	AD-RGL1	n/a	Yeast two-hybrid
pGADM-RGL2	AD-RGL2	n/a	Yeast two-hybrid
pGADM-RGL3	AD-RGL3	n/a	Yeast two-hybrid
pGBKM-BOI	BD-BOI	n/a	Yeast two-hybrid
pGBKM-BRG1	BD-BRG1	n/a	Yeast two-hybrid
pGBKM-BRG2	BD-BRG2	n/a	Yeast two-hybrid
pGBKM-BRG3	BD-BRG3	n/a	Yeast two-hybrid

* n/a : not applicable.