

Supporting Information

Fig. S1 Verification of mutants and transgenic plants.

Fig. S2 Phenotypes of wild type (Col-0) and *agb1-2* under 10 µM GA₃.

Fig. S3 Phenotypes of wild type, *agb1-2*, and *N692967* under 1 µM GA₃ treatment at 1d, 7d, and 10d.

Fig. S4 Phenotypes of wild type, *agb1-2*, and *N692967* under 10 µM GA₃ treatment at 1d, 7d, and 10d.

Fig. S5 Phenotypes of wild type, *agb1-2*, and *N692967* under 100 µM GA₃ treatment at 1d, 7d, and 10d.

Fig. S6 Phenotypes of wild type, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1*, and *MYB62:GFP/agb1-2-4* under 1 µM GA₃ treatment at 1d, 7d, and 10d.

Fig. S7 Phenotypes of wild type, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1*, and *MYB62:GFP/agb1-2-4* under 10 µM GA₃ treatment at 1d, 7d, and 10d.

Fig. S8 Phenotypes of wild type, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1*, and *MYB62:GFP/agb1-2-4* under 100 µM GA₃ treatment at 1d, 7d, and 10d.

Fig. S9 Phenotypic identification of wild type, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1* and *MYB62:GFP/agb1-2-4* under 1 µM GA₃ treatment at 10d.

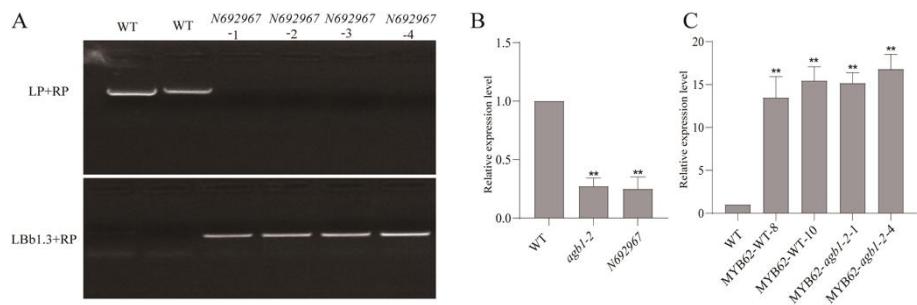
Fig. S10 Phenotypic identification of wild type, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1*, and *MYB62:GFP/agb1-2-4* under 100 µM GA₃ treatment at 10d.

Fig. S11 Phenotypes of wild type, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1*, and *MYB62:GFP/agb1-2-4* under normal conditions and phosphate-free conditions.

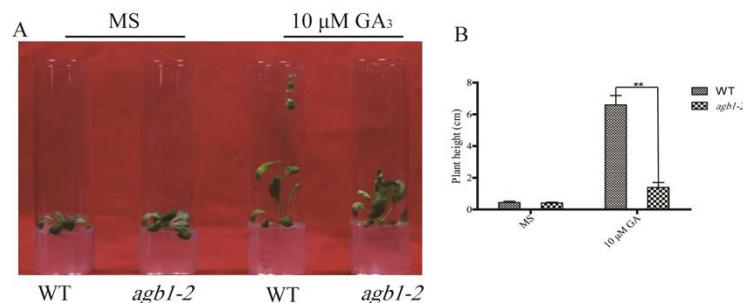
Table S1 Primers used in this study

Table S2 Composition of MS powder

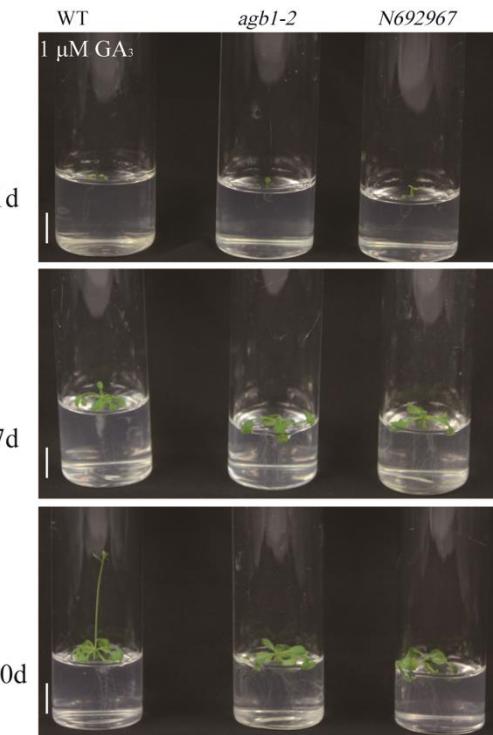
Table S3 Composition of MS powder phosphate-free



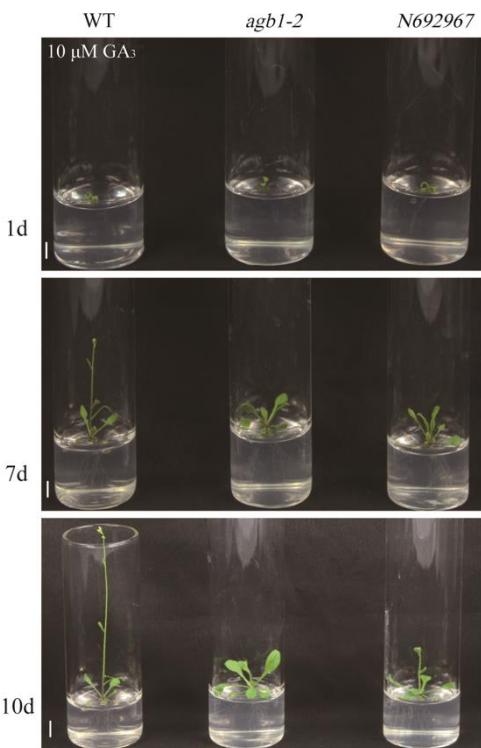
Supplementary Figure S1. Verification of mutants and transgenic plants. (A) Genomic PCR verification of *AGB1* mutant *N692967*. (B) Analysis of *AGB1* expression in *agb1-2* and *N692967* mutants, grown for four weeks, under normal growth conditions. Data is the average of three independent experiments, and the error bar represents SE ($n = 3$). Significant differences were analyzed by Duncan's multiple range test ($P < 0.05$). (C) Expression analysis of *MYB62* in transgenic plants *MYB62:GFP/WT* and transgenic plants *MYB62:GFP/agb1-2*, grown for four weeks, under normal growth conditions. Data is the average of three independent experiments, and the error bar represents SE ($n = 3$). Asterisks indicate significant differences between genotypes. Significant differences were analyzed by Duncan's multiple range test ($P < 0.05$).



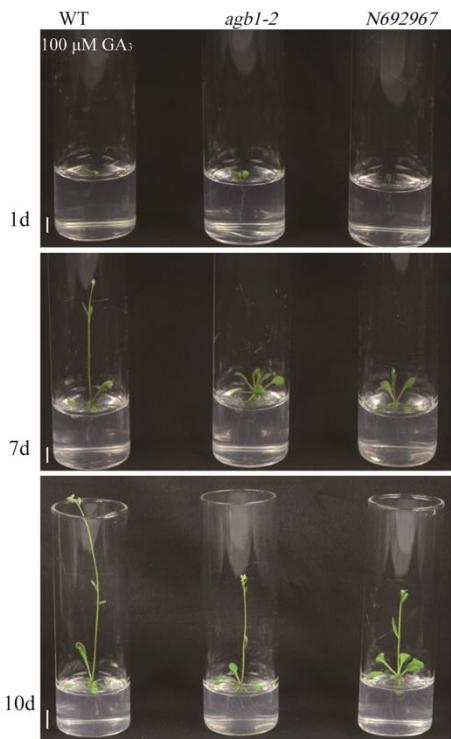
Supplementary Figure S2. Phenotype of mutant *agb1-2* and wild type (*Col-0*) under treatment with 10 μ M GA₃. (A) Plant height phenotype of wild type (*Col-0*), *agb1-2* under normal conditions and 10 μ M GA₃ treatment. Scale bars, 1 cm. (B) Plant heights of wild type (*Col-0*), *agb1-2* under normal conditions and 10 μ M GA₃ treatment. Data is the average of three independent experiments, and the error bar represents SE ($n = 10$). Asterisks indicate significant differences between genotypes. Significant differences were analyzed by Duncan's multiple range test ($P < 0.05$).



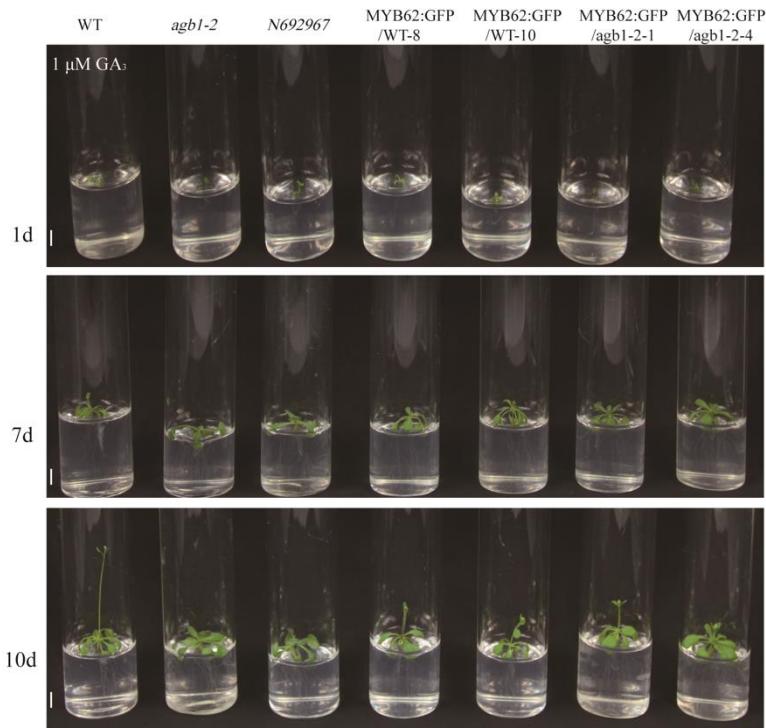
Supplementary Figure S3 Phenotype of wild type, *agb1-2*, and *N692967* under 1 μ M GA₃ treatment at 1d, 7d, and 10d. Scale bars, 1 cm.



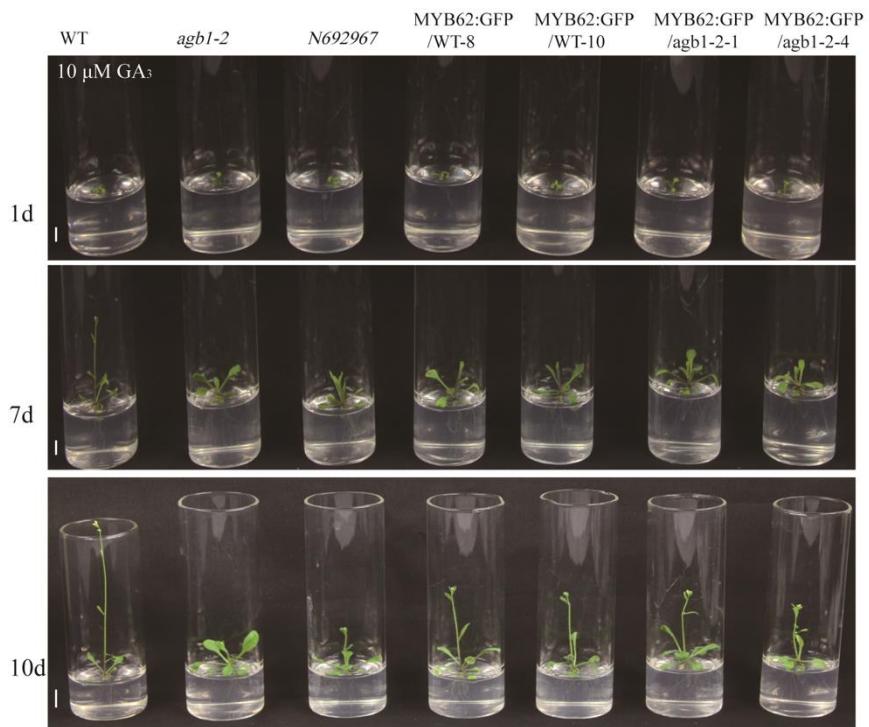
Supplementary Figure S4. Phenotypes of wild type, *agb1-2*, and *N692967* under 10 μ M GA₃ treatment at 1 d, 7 d, and 10 d. Scale bars, 1 cm.



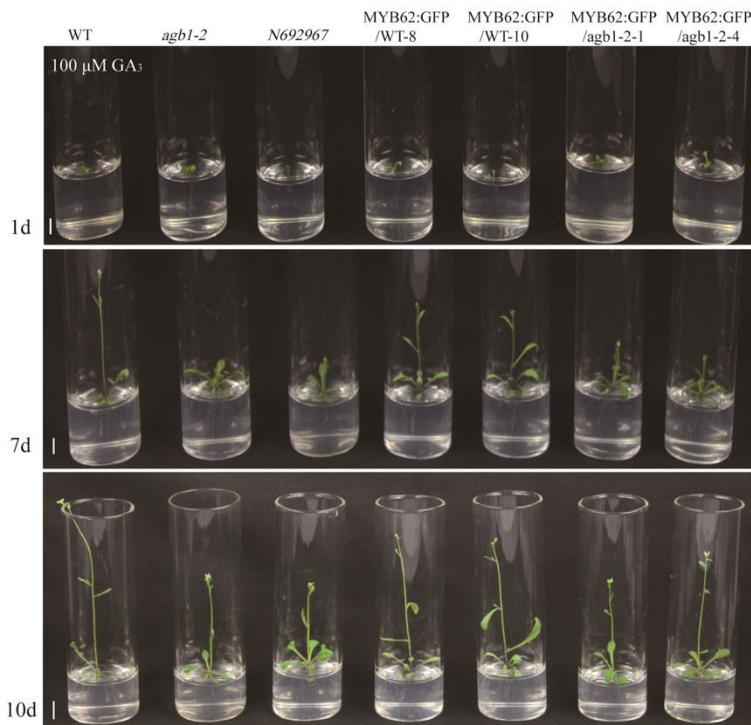
Supplementary Figure S5. Phenotypes of wild type, *agb1-2*, and *N692967* under 100 μM GA₃ treatment at 1 d, 7d, and 10 d. Scale bars, 1 cm.



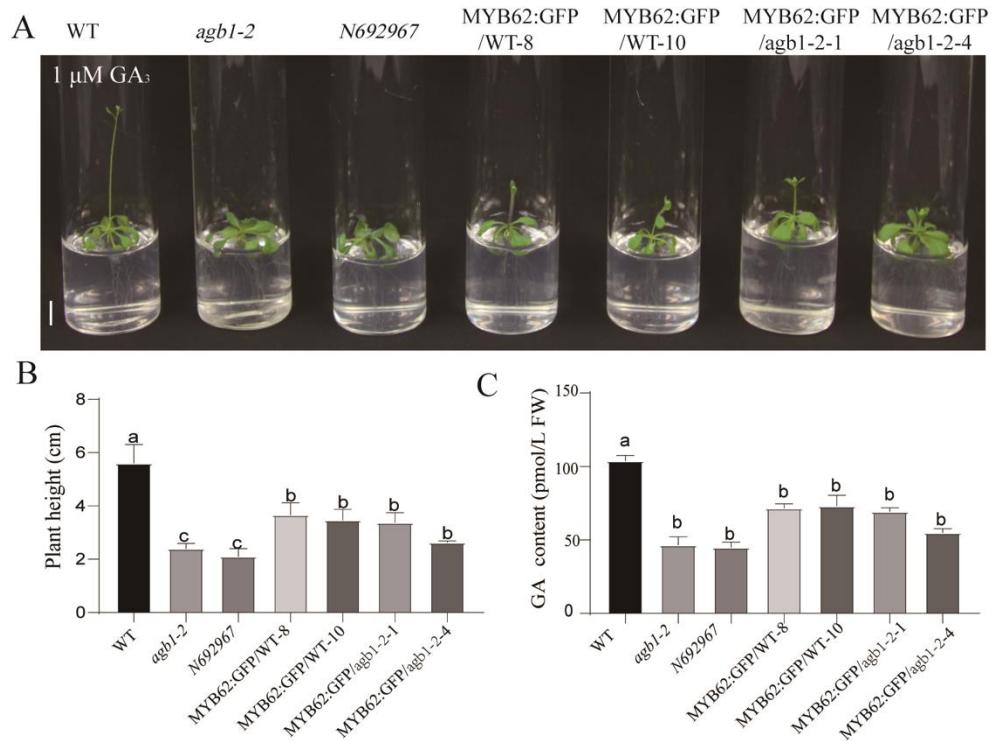
Supplementary Figure S6. Phenotypes of wild type, *agb1-2*, *N692967*, MYB62:GFP/WT-8, MYB62:GFP/WT-10, MYB62:GFP/*agb1-2-1*, and MYB62:GFP/*agb1-2-4* under 1 μM GA₃ treatment at 1d, 7d, and 10d. Scale bars, 1 cm.



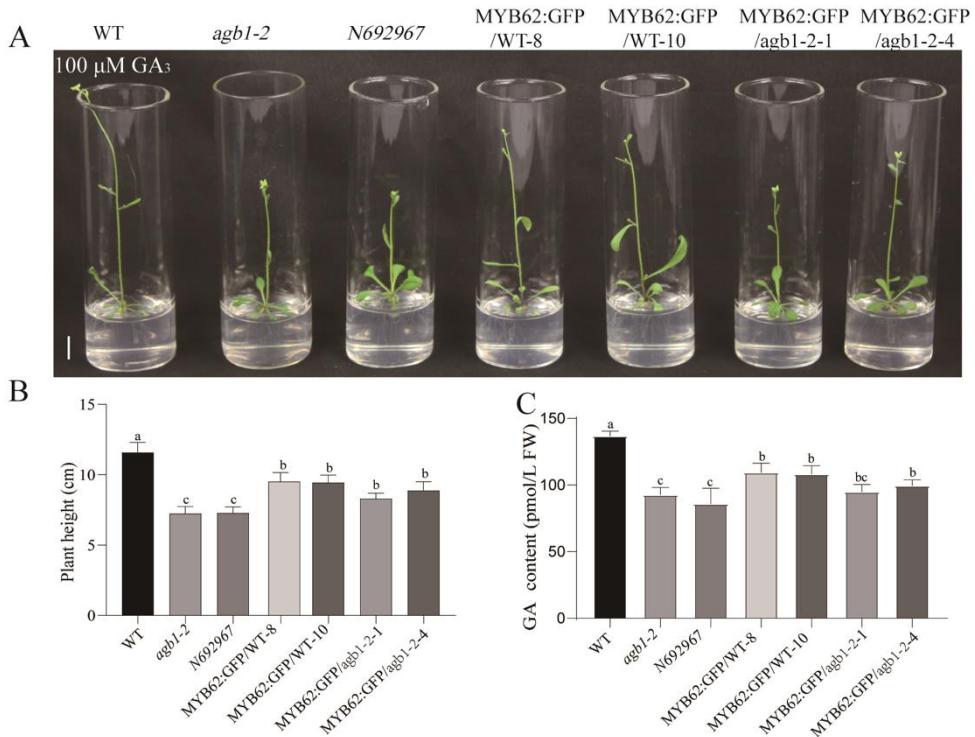
Supplementary Figure S7. Phenotypes of wild type, *agb1-2*, N692967, MYB62:GFP/WT-8, MYB62:GFP/WT-10, MYB62:GFP/*agb1-2-1*, and MYB62:GFP/*agb1-2-4* under 10 μ M GA₃ treatment at 1 d, 7 d, and 10 d. Scale bars, 1 cm.



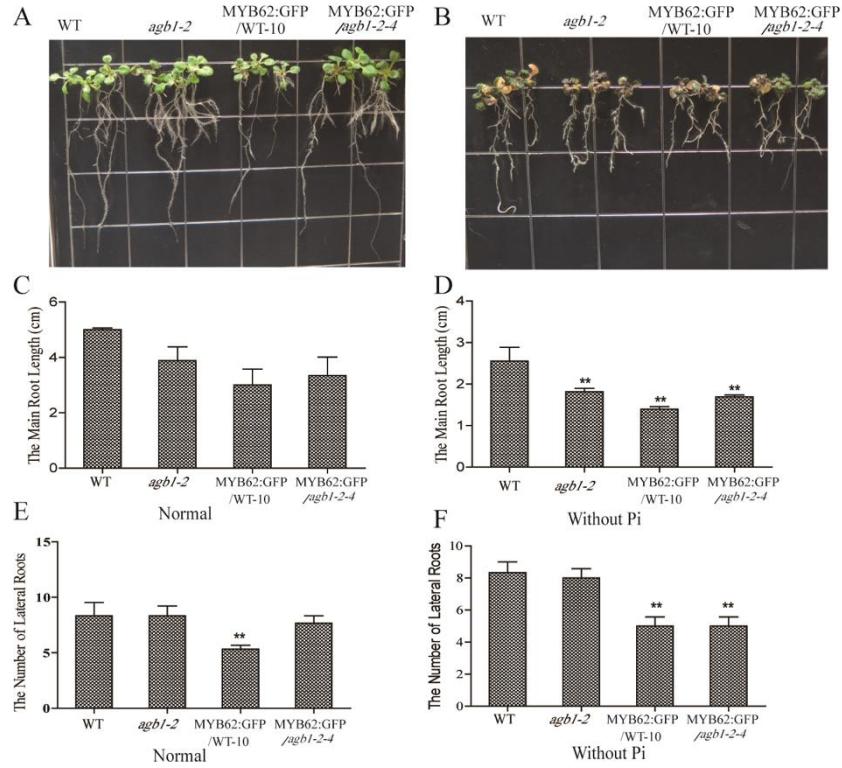
Supplementary Figure S8. Phenotypes of wild type, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1*, and *MYB62:GFP/agb1-2-4* under 100 μ M GA₃ treatment at 1 d, 7 d, and 10 d. Scale bars, 1 cm.



Supplementary Figure S9. Phenotypic identification of WT, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1*, and *MYB62:GFP/agb1-2-4* under 1 μ M GA₃ treatment at 10 d. (A) Plant height phenotypes of wild type, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1* and *MYB62:GFP/agb1-2-4* following 1 μ M GA₃ treatment at 10 d. Scale bars, 1 cm. (B-C) Plant heights and GA contents of wild type, *agb1-2*, *N692967*, *MYB62:GFP/WT-8*, *MYB62:GFP/WT-10*, *MYB62:GFP/agb1-2-1*, and *MYB62:GFP/agb1-2-4* under normal conditions. Data is the average of three independent experiments, and the error bars represent SE ($n = 10$). Asterisks indicate significant differences between genotypes. Significant differences were analyzed by Duncan's multiple range test ($P < 0.05$).



Supplementary Figure S10. Phenotypic identification of WT, *agb1-2*, N692967, MYB62:GFP/WT-8, MYB62:GFP/WT-10, MYB62:GFP/*agb1-2-1*, and MYB62:GFP/*agb1-2-4* under 100 μM GA_3 treatment at 10 d. (A) Plant height phenotypes of wild type, *agb1-2*, N692967, MYB62:GFP/WT-8, MYB62:GFP/WT-10, MYB62:GFP/*agb1-2-1*, and MYB62:GFP/*agb1-2-4* following 100 μM GA_3 treatment at 10d. Scale bars, 1 cm. (B-C) Plant heights and GA contents of wild type, *agb1-2*, N692967, MYB62:GFP/WT-8, MYB62:GFP/WT-10, MYB62:GFP/*agb1-2-1*, and MYB62:GFP/*agb1-2-4* under normal conditions. Data is the average of three independent experiments, and the error bars represent SE ($n = 10$). Asterisks indicate significant differences between genotypes. Significant differences were analyzed by Duncan's multiple range test ($P < 0.05$).



Supplementary Figure S11. Phenotypes of wild type, *agb1-2*, *MYB62:GFP/WT-10*, and *MYB62:GFP/agb1-2-4* under normal conditions and phosphate-free conditions. (A) Phenotype identification of wild type, *agb1-2* mutant, *MYB62:GFP/WT-10*, and *MYB62:GFP/agb1-2-4* under normal conditions. Bar = 1cm. (B) Phenotype identification of wild type, *agb1-2* mutant, *MYB62:GFP/WT-10*, and *MYB62:GFP/agb1-2-4* under phosphate-free conditions. Bar = 1cm. (C) The main root lengths of wild type, *agb1-2* mutant, *MYB62:GFP/WT-10*, and *MYB62:GFP/agb1-2-4* under normal conditions. Error bars represent the means ± SE ($n = 3$). Asterisks indicate significant differences between genotypes. Significant differences were analyzed by Duncan's multiple range test ($P < 0.05$). (D) The numbers of lateral roots of the wild type, *agb1-2* mutant, *MYB62:GFP/WT-10*, and *MYB62:GFP/agb1-2-4* under normal conditions. Error bars represent the means ± SE ($n = 3$). Significant differences were analyzed by Duncan's multiple range test ($P < 0.05$). (E) The main root lengths of wild type, *agb1-2* mutant, *MYB62:GFP/WT-10*, and *MYB62:GFP/agb1-2-4* under phosphorus-free conditions. Error bars represent the means ± SE ($n = 3$). Asterisks indicate significant differences between genotypes. Significant differences were analyzed by Duncan's multiple range test ($P < 0.05$). (F) The numbers of lateral roots of wild type, *agb1-2* mutant, *MYB62:GFP/WT-10*, and *MYB62:GFP/agb1-2-4* under phosphorus-free conditions. Error bars represent the means ± SE ($n = 3$). Asterisks indicate significant differences between genotypes. Significant differences were analyzed by Duncan's multiple range test ($P < 0.05$).

Table S1 Primers used in this study

Name	Primer sequence (5'-3')
Primers used for qRT-PCR	
<i>qRT-AtAGB1-F</i>	ACGTTGGGATACCCTCTTGG
<i>qRT-AtAGB1-R</i>	GTGTCCTCCAAACGCCATA
<i>qRT-AMYB62-F</i>	TGGCTCGATGTGTTCCATGA
<i>qRT-AtMYB62-R</i>	ACGCTGAATAACCATCCGCC
<i>qRT-AtGA2ox7-F</i>	AGTGGTGAGGAGGTCAAACG
<i>qRT-AtGA2ox7-R</i>	GAGAAGTGGCGCTAGGGTT
<i>qRT-AtGA2ox4-F</i>	TGTCCGACCAGTTAACCGAC
<i>qRT-AtGA2ox4-R</i>	GGCAGGGTCGTTAGTGTGAA
<i>qRT-AtGAMT1-F</i>	ATGGAGTCGTACGGAGCC
<i>qRT-AtGAMT1-R</i>	TGGAGTTGATGGCTGTCGTC
Primers used to generate DNA constructs for the yeast two-hybrid assay	
<i>AGB1-BD-F</i>	AGGAGGACCTGCATATGATGTCGTCTCCG
<i>AGB1-BD-R</i>	GCCTCCATGGCCATATG AATCACTCTCCTG
<i>MYB62-AD-F</i>	CATCGATACTGGATCCATGGAAAATTGA
<i>MYB62-AD-R</i>	CGAGCTCGATGGATCCATCTCCCTAAACTG
Primers used to generate DNA constructs for the subcellular localization assay	
<i>AGB1-16318-F</i>	TATCTCTAGAGGATCCATGGGCTTACTCTGC
<i>AGB1-16318-R</i>	TGCTCACCATGGATCCAATCACTCTCCTGT
<i>MYB62-16318-F</i>	TATCTCTAGAGGATCCATGGAAAATTGAT
<i>MYB62-16318-R</i>	TGCTCACCATGGATCCCTCCCTAAACTGCC
Primers used to generate DNA constructs for protein expression	
<i>AGB1-GST-FULL-F</i>	CGGGATCCATGTCTGTCCTCGAGCTCAAAG
<i>AGB1-GST-FULL-R</i>	GCGTCGACAATCACTCTCCTGTGTCCTC
<i>MYB62-MBP-F</i>	CGGGATCCATGTCTACCGATGTGGCGAGGT
<i>MYB62-MBP-R</i>	GCGTCGACAATCACTCTCCTGTGTCCTC
Primers used to generate DNA constructs for LUC assay	
<i>AGB1-nLUC-F</i>	TCGGTACCCGGGATCCATGTCTGTCCTCGAG
<i>AGB1-nLUC-R</i>	CCATTGTTGGATCCAATCACTCTCCTGTG
<i>MYB62-cLUC-F</i>	CGGGCGGTACCATGGAAAATTGA
<i>MYB62-cLUC-R</i>	GTTGCTGCAGGTGACCTCCCTAAACTG
Primers used to generate DNA constructs for Co-IP assays	
<i>AGB1-Flag-F</i>	GGGGCCCGGGTCGACATGTCTGTCCTCGAGCTCAAAG
<i>AGB1-Flag-R</i>	TACCGGATCCACTAGTAATCACTCTCCTGTGTCCTC
<i>MYB62-GFP-F</i>	GATTACGAATTCATGGAAAATTGATG

<i>MYB62-GFP-R</i>	GTGCTCGAATTCCCTCCCTAAACTGCCAAATG
Primers used for genotyping	
<i>AGB1-F (LP)</i>	TCATTAGATTGGACACCGGAG
<i>AGB1-R (RP)</i>	TGTGAATCCTGCTGTAATCCC
<i>MYB62-F (P1)</i>	TGAGATCAATCGGCTAAAAGC
<i>MYB62-R (P2)</i>	CAAAGATTGCGATTCATCGAT
<i>LBB1.3</i>	ATTTGCCGATTTCGGAAC
Primers used to produce transgenic plants	
<i>OE-MYB62-F</i>	GGGGCCCGGGGTCGACATGGAAAATTGATGAAGAA GAAGA
<i>OE-MYB62-R</i>	TACCGGATCCACTAGTCC CTCCCTAAACTGCCAAATGTCAT
Primers used to identify the homozygote N692967 mutants	
<i>LP</i>	TCATTAGATTGGACACCGGAG
<i>RP</i>	TGTGAATCCTGCTGTAATCCC
<i>LB</i>	ATTTGCCGATTTCGGAAC
Primers used for EMSA analysis	
<i>ProGA2ox7-F</i>	BIO-AAGCCCTTGGTGGGATGG
<i>ProGA2ox7-R</i>	BIO-CCATCCCCAA CCAAGGGCTT
Primers used to generate DNA constructs for LUC assays	
<i>GA2ox7-F</i>	GCAGCCCGGGGGATCCTCCATTACTAGAGAA
<i>GA2ox7-R</i>	TAGAACTAGTGGATCCTAAACTGAAT TAAGT

Table S2 Composition of MS medium

Component		
Macro elements	mg/L	mM
CaCl ₂	332.02	2.99
KH ₂ PO ₄	170.00	1.25
KNO ₃	1900.00	18.79
MgSO ₄	180.54	1.50
NH ₄ NO ₃	1650.00	20.61
Micro elements	mg/L	μM
CoCl ₂ • 6H ₂ O	0.025	0.11
CuSO ₄ • 5H ₂ O	0.025	0.10
FeNaEDTA	36.70	100.00
H ₃ BO ₄	6.2	100.27
KI	0.83	5.00
MnSO ₄ •H ₂ O	16.90	100.00
Na ₂ MoO ₄ •2H ₂ O	0.25	1.03

ZnSO ₄ •7H ₂ O	8.60	29.91
Vitamins	mg/L	µM
Glycine (free base)	2.00	26.64
myo-Inositol	100.0	554.94
Nicotinic acid (free acid)	0.50	4.06
Pyridoxine • HCl	0.50	2.43
Thiamine • HCl	0.10	0.30

Table S3 Composition of MS phosphate-free medium

Component		
Macro elements	mg/L	mM
CaCl ₂	332.02	2.99
KNO ₃	1900.00	18.79
MgSO ₄	180.54	1.50
NH ₄ NO ₃	1650.00	20.61
Micro elements	mg/L	µM
CoCl ₂ • 6H ₂ O	0.025	0.11
CuSO ₄ • 5H ₂ O	0.025	0.10
FeNaEDTA	36.70	100.00
H ₃ BO ₄	6.2	100.27
KI	0.83	5.00
MnSO ₄ •H ₂ O	16.90	100.00
Na ₂ MoO ₄ •2H ₂ O	0.25	1.03
ZnSO ₄ •7H ₂ O	8.60	29.91
Vitamins	mg/L	µM
Glycine (free base)	2.00	26.64
myo-Inositol	100.0	554.94
Nicotinic acid (free acid)	0.50	4.06
Pyridoxine • HCl	0.50	2.43
Thiamine • HCl	0.10	0.30