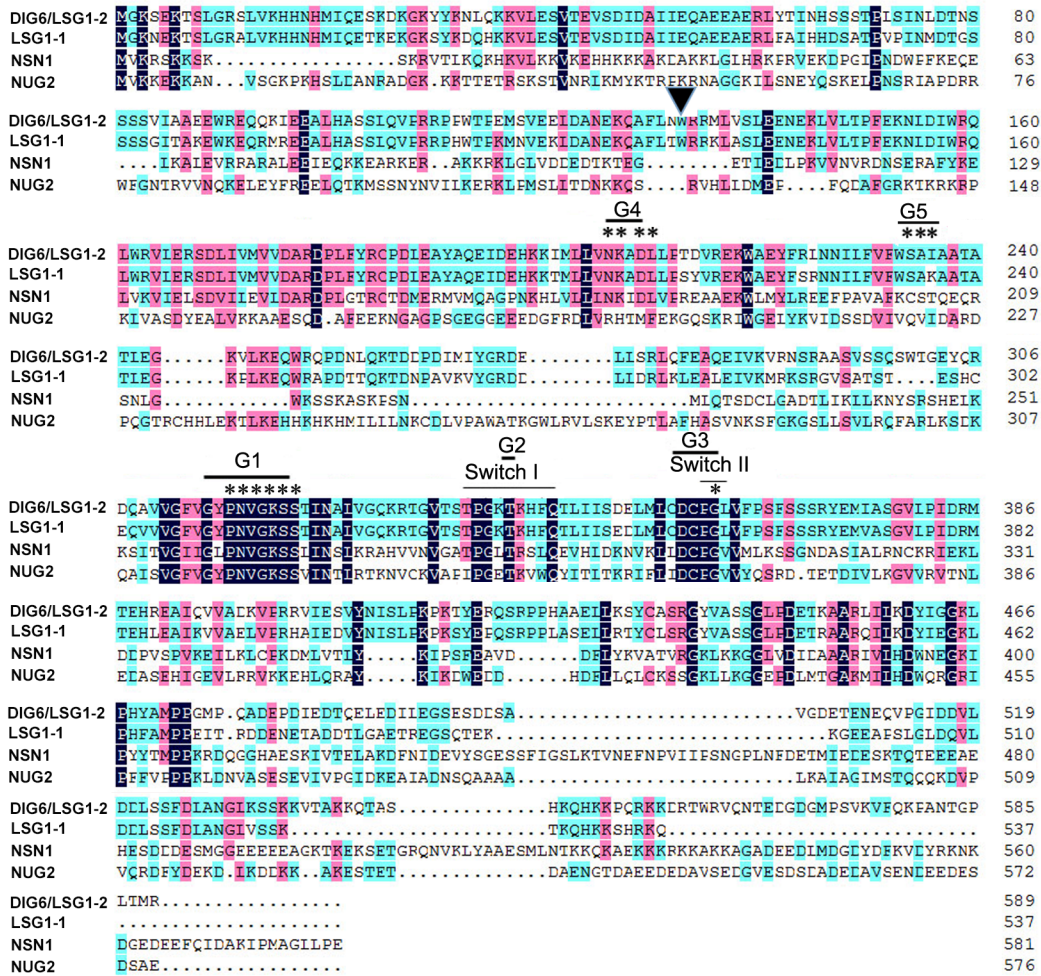
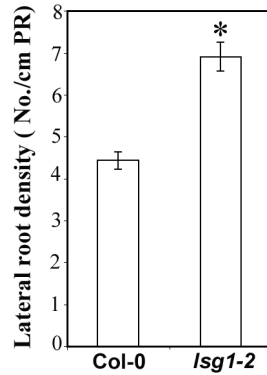


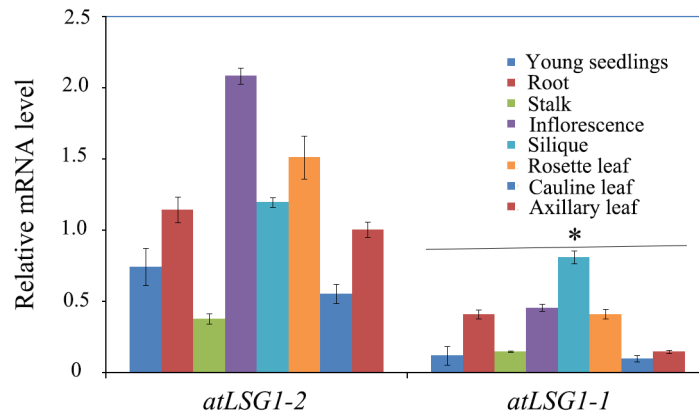
**Figure S1.** The phenotypes of *dig6* mutant. (A-C) Twelve-day-old seedlings grown on horizontally (A-B) or vertically (C) placed agar plates. Seedlings in (A) and on the left of panel (C) are wild type (*Col-gll*) and in (B) and on the right of panel (C) are *dig6*. (D) Twenty-eight-day-old *Col-gll* (left) and *dig6* (right). (E-F) Newly emerged axillary leaves of *Col-gll* (E) and *dig6*.



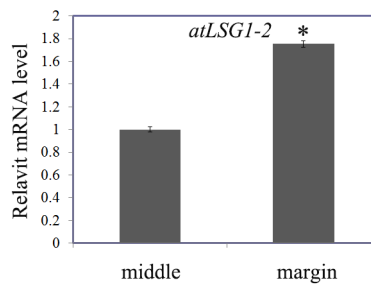
**Figure S2.** Multiple alignment of AtLSG1-related proteins. The accession numbers of DIG6/LSG1-2, LSG1-1, NSN1 and NUG2 are Atlg08410, At2g27200, At3g07050, and At1g52980, respectively. The putative motifs were marked by black lines above the sequence. Asterisks represent GTP/Mg<sup>2+</sup> binding sites.



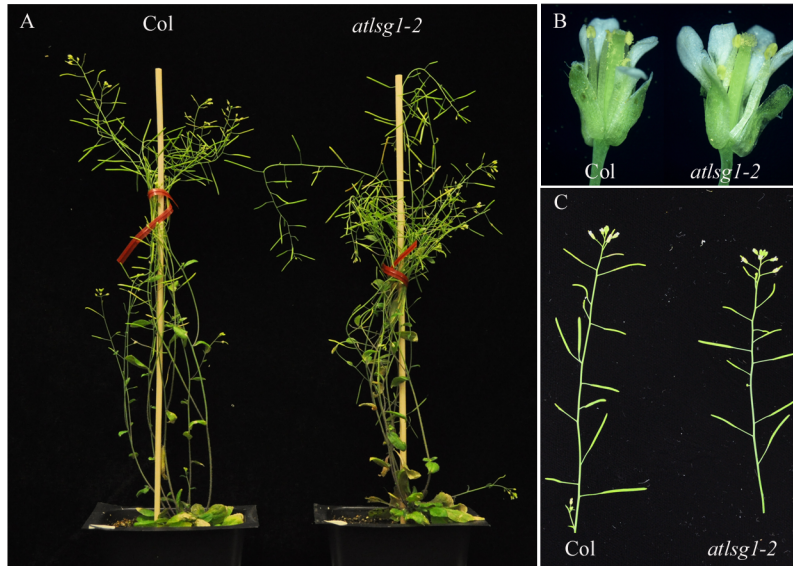
**Figure S3.** Lateral root density. Lateral root number was counted in the primary root of 12-day-old seedlings. Asterisk indicates significant difference from wild type plants (*t*-test, \*  $p < 0.01$ ).



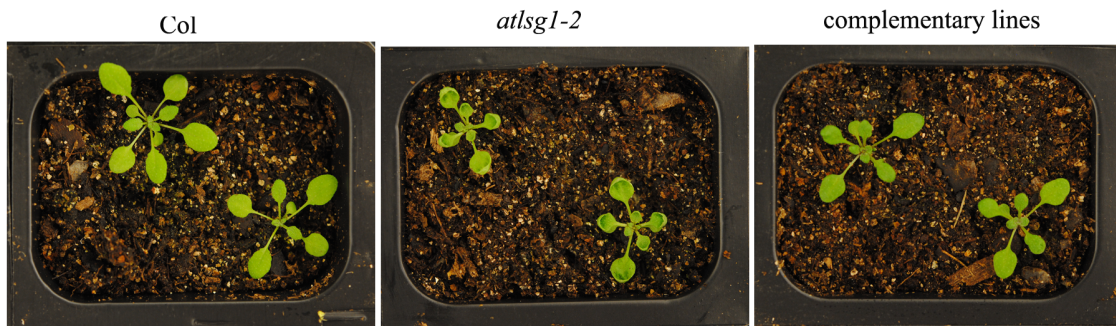
**Figure S4.** The tissue-specific expression pattern of *atLSG1-1* and *atLSG1-2*. RNA was extracted from 12-day-old seedlings and different organs of mature plants. Real time RT-PCR was carried out. Student *t*-test was carried out for comparing the different expression levels of *atLSG1-1* and *atLSG1-2* in various organs. Asterisk indicates significant differences between two genes (*t*-test, \*  $p < 0.01$ ).



**Figure S5.** The expression of *atLSG1-2* in leaf middle and margin regions. Total RNA was extracted from the leaf mid vein or margin regions, and real time RT-PCR was carried out using *atLSG1-2* specific primers. Asterisk indicates significant difference from wild type plants (*t*-test, \*  $p < 0.01$ ).



**Figure S6.** The *atlsg1-2* mutant displayed no visible phenotype at late development stages. Morphology of (A) the mature plants, (B) flowers, and (C) siliques of the wild type Col-0 and *atlsg1-2*.



**Figure S7.** *35S::YFP-AtLSG1* complements the phenotypes of the *lsg1-2* mutant.

**Table S2 Primer list in this study**

<b>purpose</b>	<b>gene name</b>	<b>sequence</b>
<b>for mapping</b>	Chr1-1.070MF	GAGCTGGTGAGAGCCAAAGTTTCC
	Chr1-1.070MR	CACGTAGAAGCGATCACAGAGATAC
	Chr1-3.824MF	CCCTAGTCGAGTATTACAGCGACTC
	Chr1-3.824MR	TACAGCCATTAGGTCTTATAACCGAT
<b>for checking the transcripts of</b>	OXH89-F	CTCGAGAACTACTGCGACAGG
	OXH76-R	TGGAGACAACCCGATAACTTG
<b>for vector construction</b>	AtLSG1-F	CACCATGGGGAAGAGCGAAAAGAC
	AtLSG1-R	CCGCATCGTAAGAGGACCTG TG
	AtLSG1-R	TCACCGCATCGTAAGAGGAC
	AtLSG1 like-F	CACCATGGGGAAGA ACGAGAAAAC
	AtLSG1 like-R	TTGTTTCTATGTGACTTCTT
<b>for making yeast constructs</b>	LSG1-F	CTGACTAGTAAAATGCCACAAAAGAAGCTCC
	LSG1-R	TCTGTCGACCTAATTATTTCAATGCTAAAAACT
	AtLSG1-F	CTGACTAGTAAAATGGGGAAGAGCGAAAAGAC
	AtLSG1-R	ACTGTCGACGGATCCTCACCGCATCGTAAGAGGAC
	AtLSG1 like-F	CTGACTAGTAAAATGGGGAAGAACGAGAAAAC
	AtLSG1 like-R	ACTGTCGACGGATCCTCATTGTTTCTATGTGACTT
<b>for making PAtLSG1::GUS construct</b>		CACCTTGACAGACTGTGAGTTGGAT
		CTCTCTGTTTCAAGTAAACC
<b>For making artificial microRNA</b>	I miR-s	gaTTACTTGAAACTACTCGCGTtctctctttgtattcc
	II miR-a	gaACGCGAGTAGTCTTCAAGTAAtcaagagaatcaatga
	III miR*s	gaACACGAGTAGTCTACAAGTATcacaggtcgtgatg
	IV miR*a	gaATACTTGTAGACTACTCGGTtctacatatattcct
<b>For real-time PCR</b>	PIN3-F	CGCCGAGATGGTCTAAACAAA
	PIN3-R	AAAGCGACGAGAGCCAAATAA
	IAA10-F	GTGAAGGTGACAATGGACGG
	IAA10-R	GGAACATCTCCTACAAGCATCC
	IAA20-F	GAGGAGGAAGAAGAGAATGAGTGT
	IAA20-R	TCAGCCCAGAGAATGGATGC
	IAA30-F	GAACCTAAGCACGGACCTGA
	IAA30-R	TCATCATTTCTCTGCCGA
	IAA13-F	GATGCTTGTGGTGTGTTCCA
	IAA13-R	TGCTTTCGCTGTCTCTCGTT
	GH3.6-F	ATGTCCTAAGCGTGGCTGGT
	GH3.6-R	GGCGTGTACCCTCAAGCA
	LBD33-F	GATTGAGTTTCTCGGGTCACAG
	LBD33-R	CCCATCATAGCAGTTTCTCCG
	ACS6 -F	TCCAGGCGTTCGTTCCATT
	ACS6 -R	AGCCATCGGTTTAGTCTCTCC