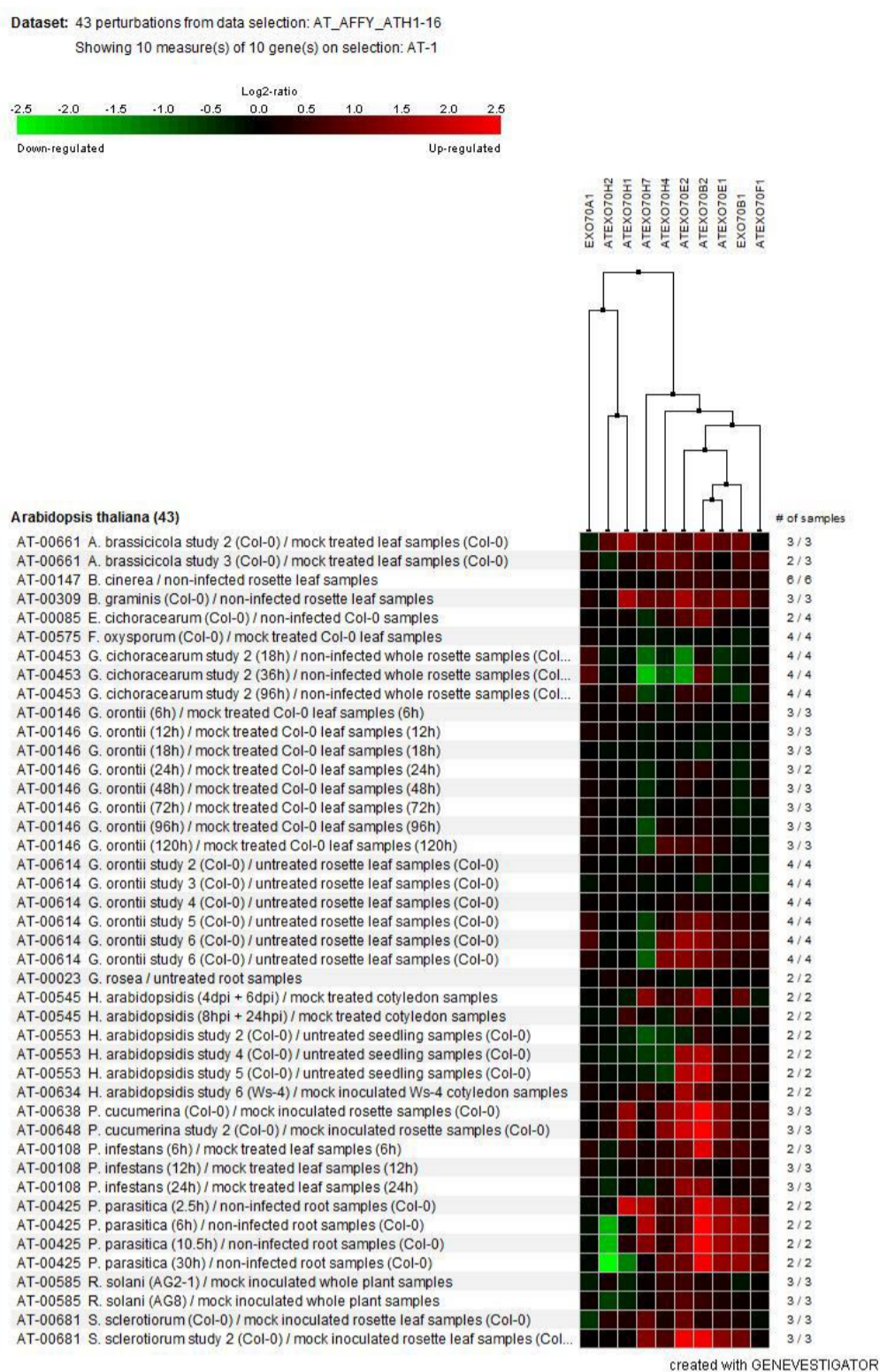
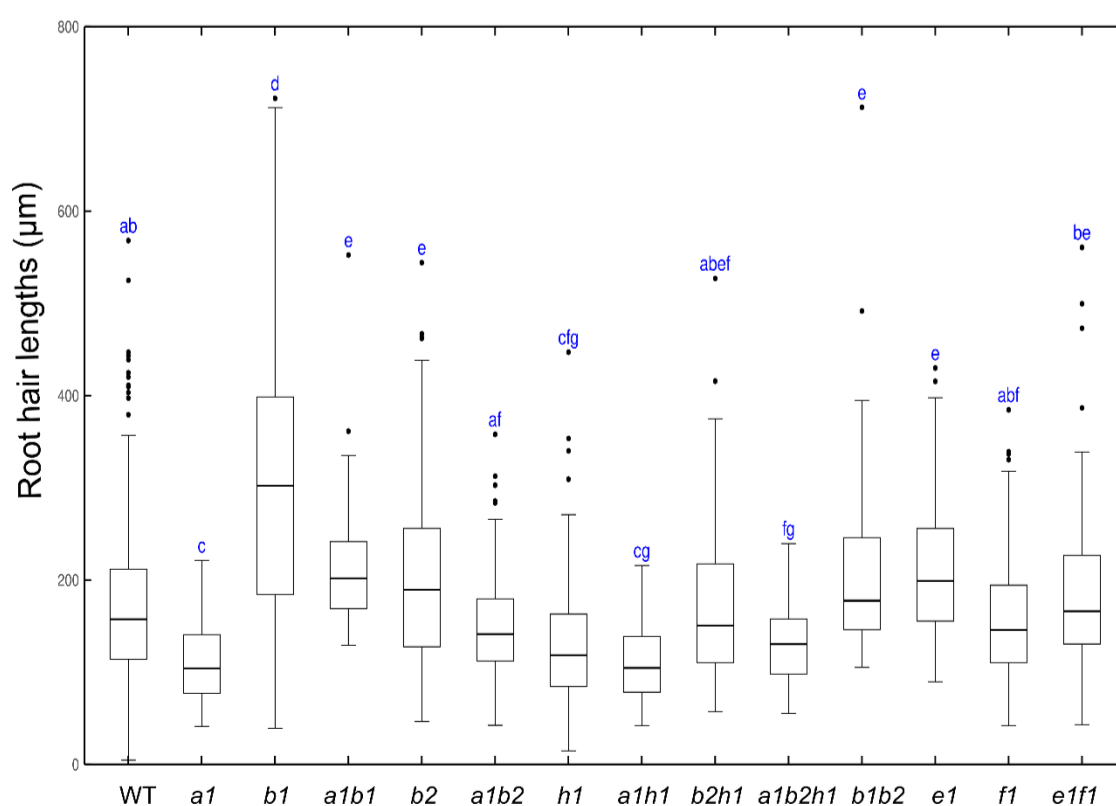




**Supplementary figure 2.** Hierarchical clustering of EXO70 isoforms according to expression levels upon fungal and oomycete treatments.

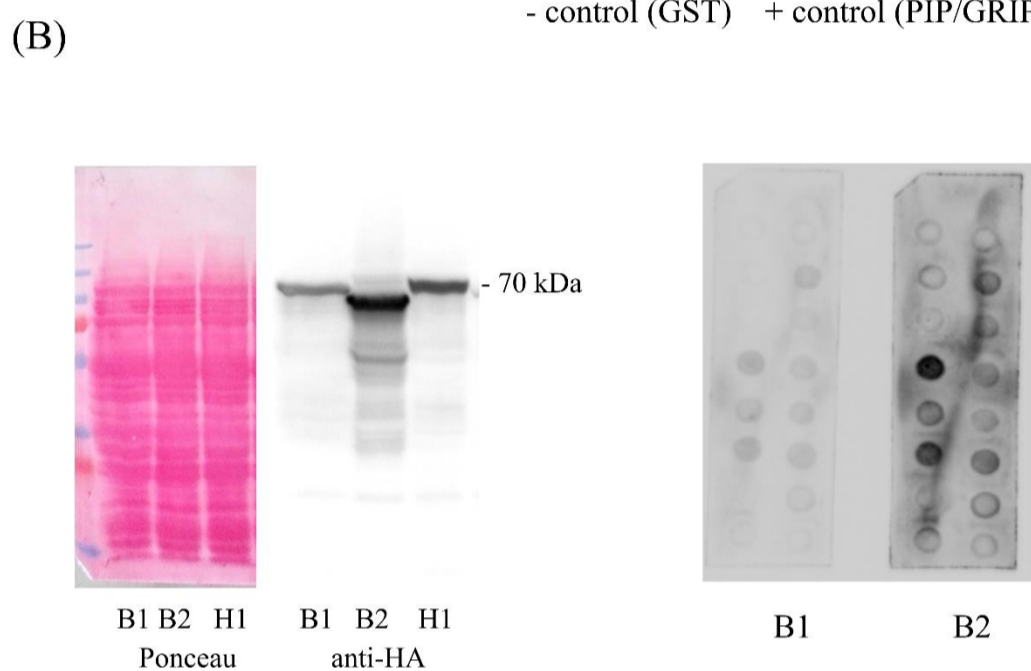
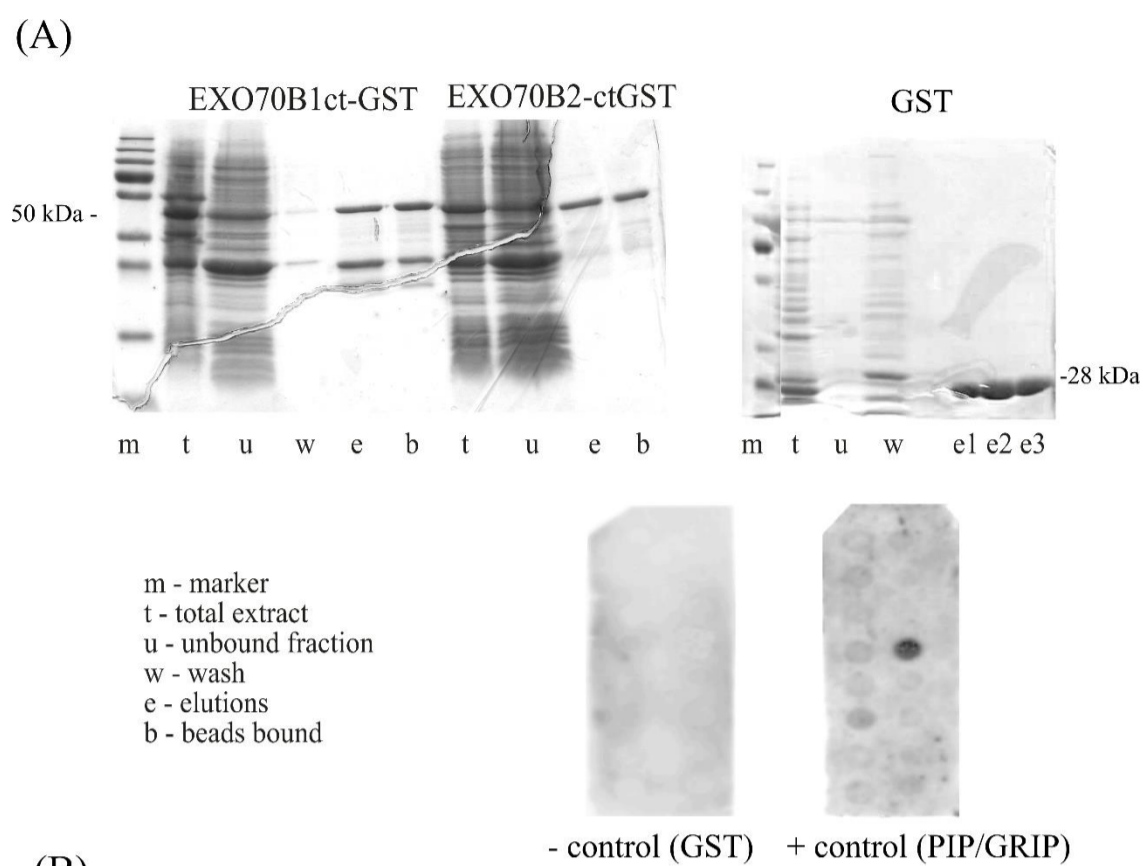


**Supplementary figure 3.** Absolute root hair lengths of mock treated seedlings for analyzed single and multiple mutant lines.





**Supplementary figure 4.** (A) Coomassie stained gels with aliquots of GST-fused EXO70B1 and B2 C termini as well as free GST (upper row); PIP/GST as negative and PIP/GRIP protein (Echelon, Israel) as positive strip controls (lower row). (B) Ponceau stained membranes and anti-HA (Sigma) detection of *in vitro* translated HA-tagged EXO70B1, B2 and H1 (4x more amount used for incubation with strips); comparison of the strength of signal detected on PIP strips for EXO70B1 and B2 when equal amount of proteins and exposure times for signal detection were used.



**Supplementary table 1.** Primers used for cloning and genotyping.

Purpose	Forward primer <sup>1</sup>	Reverse primer <sup>1</sup>
GST-EXO70B1-ct (EcoRI-SalI)	<u>aagaattct</u> cctaaagattcaactctggtgactg	aagg <u>tcgact</u> cattttgttcccgtgg
GST-EXO70B2-ct (EcoRI-XhoI)	aagaatt <u>cggat</u> cagatacagagaccggtg	aaact <u>cgagt</u> caacttgagctttcctt
HA-EXO70B1	gatgttccagattacgctatggcggagaatggtg	atag <u>cgccgct</u> cattttcttcccgtggt
HA-EXO70B2	gatgttccagattacgctatggctgaagccggt	atag <u>cgccgct</u> caacttgagctttccttg
HA-EXO70H1	gatgttccagattacgctatggcgaaaatggcga	atag <u>cgccgct</u> cagcctgaaacacacc
pTNT-HA-overhang_SalI <sup>2</sup>	aag <u>tcgacg</u> ccgccaccatgtaccatacagatggtccagattacgctatg	
pTNT-HA-overhang_XhoI <sup>3</sup>	ata <u>ctcgagg</u> ccgccaccatgtaccatacagatggtccagattacgctatg	
<i>exo70E1</i>	ctgatgatggttcttcaaatagctatgg	acagaggttatggagtagaaatgtcc

<sup>1</sup> All primer sequences are presented in 5' to 3' direction. Introduced restriction sites are underlined.

<sup>2</sup>Used for B1,H1 SalI-NotI cloning. <sup>3</sup>Used for B2 XhoI-NotI cloning.

**Supplementary table 2.** Sample sizes (n) used for analyzes in biotic stress assays. For mock and bacteria treated root hairs (RH) total numbers of root hairs and in parenthesis numbers of seedlings analyzed are presented (ca 20 root hairs per seedling). For flooding experiments numbers of analyzed samples (each sample is pool of 3-4 seedlings) are presented.

	<b>n for mock treated RH</b>	<b>n for bacteria treated RH</b>	<b>n for hrpH-flooding</b>
<b>WT</b>	407 (20)	303 (15)	36
<b><i>a1</i></b>	120 (6)	160 (8)	18
<b><i>b1</i></b>	80 (4)	80 (4)	18
<b><i>alb1</i></b>	80 (4)	80 (4)	12
<b><i>b2</i></b>	160 (8)	160 (8)	18
<b><i>alb2</i></b>	123 (6)	103 (5)	12
<b><i>h1</i></b>	120 (6)	100 (5)	18
<b><i>alh1</i></b>	60 (3)	120 (6)	12
<b><i>b2h1</i></b>	80 (4)	80 (4)	12
<b><i>alb2h1</i></b>	80 (4)	80 (4)	10
<b><i>b1b2</i></b>	100 (5)	100 (5)	12
<b><i>e1</i></b>	135 (7)	116 (6)	6
<b><i>f1</i></b>	152 (5)	119 (6)	6
<b><i>e1f1</i></b>	120 (6)	120 (6)	6