

Supporting Information

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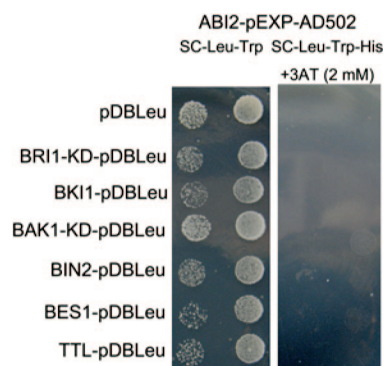


Fig. S1. ABI2 does not interact with several known components of BR signaling, including BRI1, BAK1, TTL, BK11, BIN2, and BES1, respectively, in yeast. *ABI2* was cloned into the pEXP-AD502. The kinase domains of BRI1 (BRI1-KD) and BAK1 (BAK1-KD) and the full-length of TTL, BK11, BIN2, and BES1 were cloned into the pDBLeu vector. Plasmids containing fusion proteins were cointroduced into *Saccharomyces cerevisiae* AH109 and were tested on SD medium lacking Leu, Trp, and His in the presence of 2 mM 3-amino-1,2,4-triazole (3AT). A ProQuest Two-Hybrid system with Gateway technology (Invitrogen) was used.

Table S1. BR and ABA-related mutants used in this study

Mutants	Genetic background	Notes
<i>det2-1</i>	Col-0	Steroid 5 α -reductase
<i>bri1-301</i>	Col-0	A point mutation of <i>BRI1</i>
<i>bri1-116</i>	Col-0	A null allele of <i>BRI1</i>
<i>BK11-YFP-OX</i>	Col-0	Overexpression line of <i>BK11-YFP</i>
<i>BK11-YFP-OX/bri1-116</i>	Col-0	Overexpression line of <i>BK11-YFP</i> in BR mutant <i>bri1-116</i>
<i>BRI1-GFP-OX</i>	Col-0	Overexpression line of <i>BRI1-GFP</i>
<i>aba1</i>	Ler	Zeaxanthin epoxidase
<i>aba2-1</i>	Col-0	Dehydrogenase/reductase
<i>abi1</i>	Ler	A serine/threonine phosphatase
<i>abi2</i>	Ler	A serine/threonine phosphatase