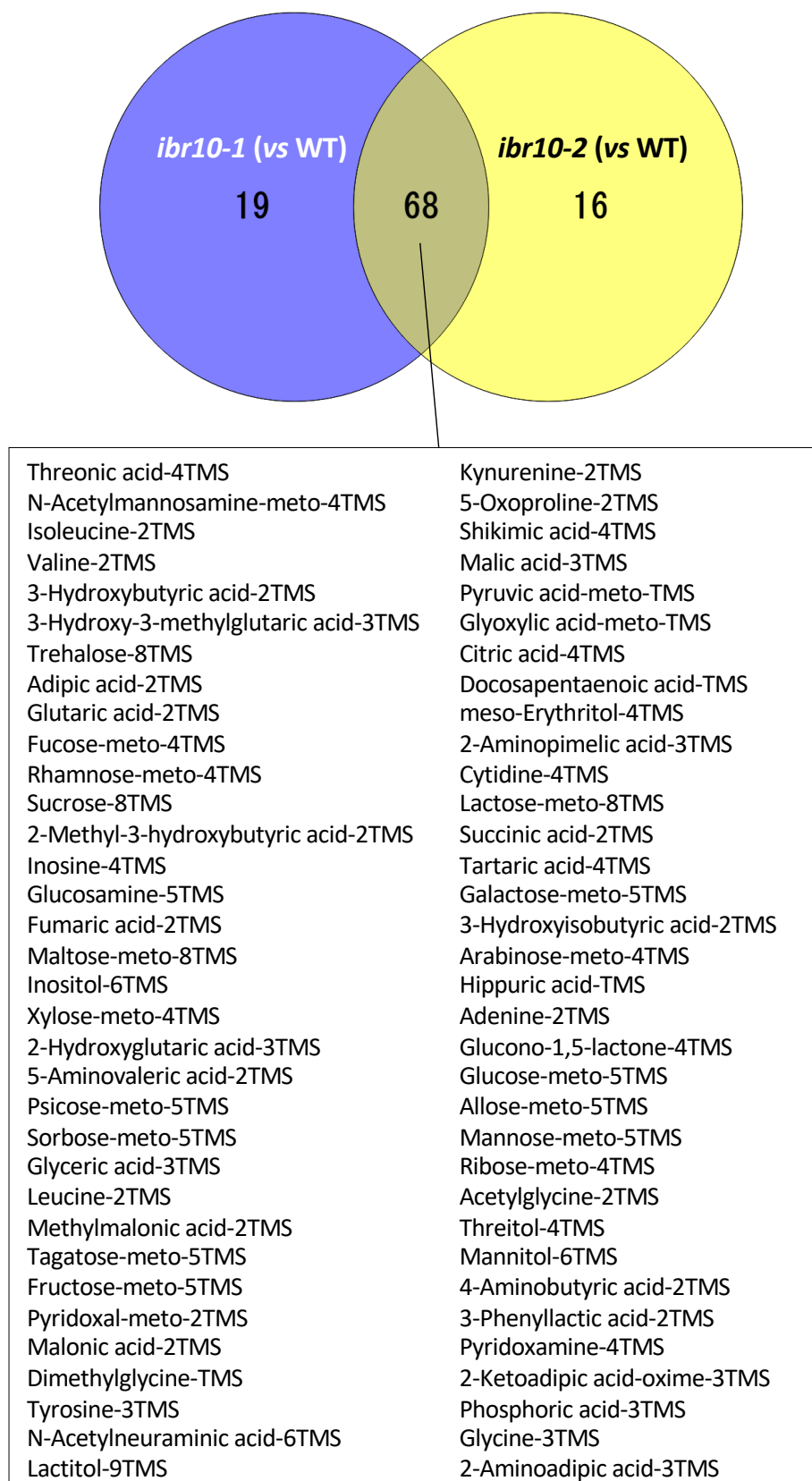


Supplementary Material

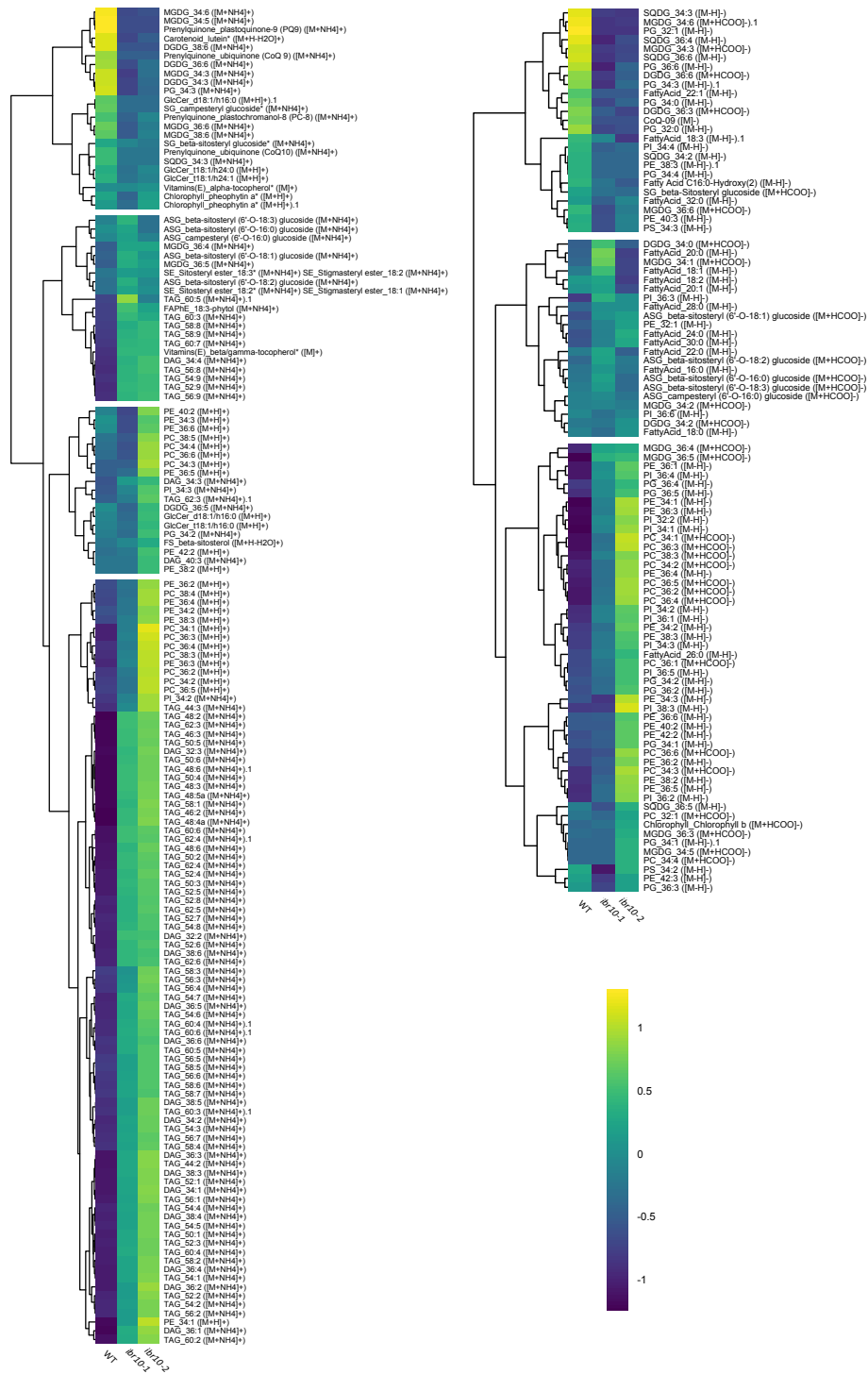
Skotomorphogenesis exploits threonine to promote hypocotyl elongation

Hiromitsu Tabeta, Higashi Yasuhiro, Okazaki Yozo, Kouji Takano, Kiminori Toyooka, Mayumi Wakazaki, Mayuko Sato, Kazuki Saito, Masami Y Hirai, Ali Ferjani



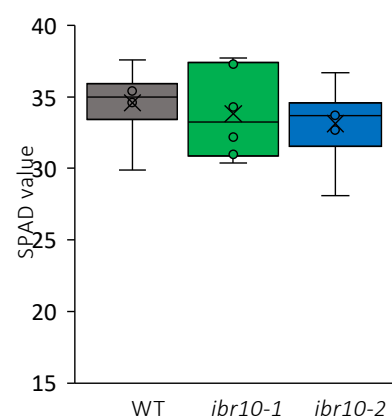
Supplementary Figure S1. Metabolic changes in *ibr10* mutants. (In support of Figure 2)

Venn diagram for each mutant line (3 DAI). Numbers represent the metabolites whose contents were significantly changed compared with the WT as determined by Student's *t*-test ($n = 6$, $P < 0.05$). Factors shown in the black box are significantly different in all mutants compared with the WT. DAI, days after induction of seed germination.



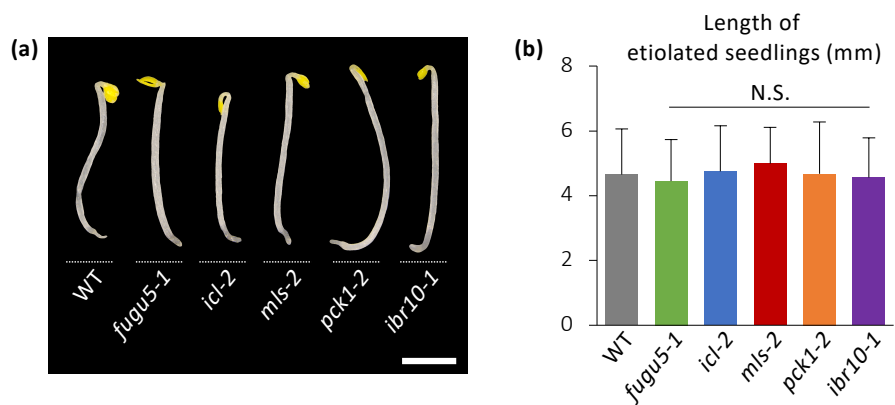
Supplementary Figure S2. LC-MS/MS lipidome analysis of *ibr10* mutants. (In support of Figure 3)

Hierarchical clustering analysis of *ibr10-1* and *ibr10-2* is shown. All 146 metabolites detected by LC-qTOF-MS were categorized into several clusters based on their behavior. The right and left sides show metabolites detected in positive and negative modes, respectively. Data are Z-scores of the average relative contents ($n = 6$) using 3 DAI etiolated seedlings. DAI, days after induction of seed germination.



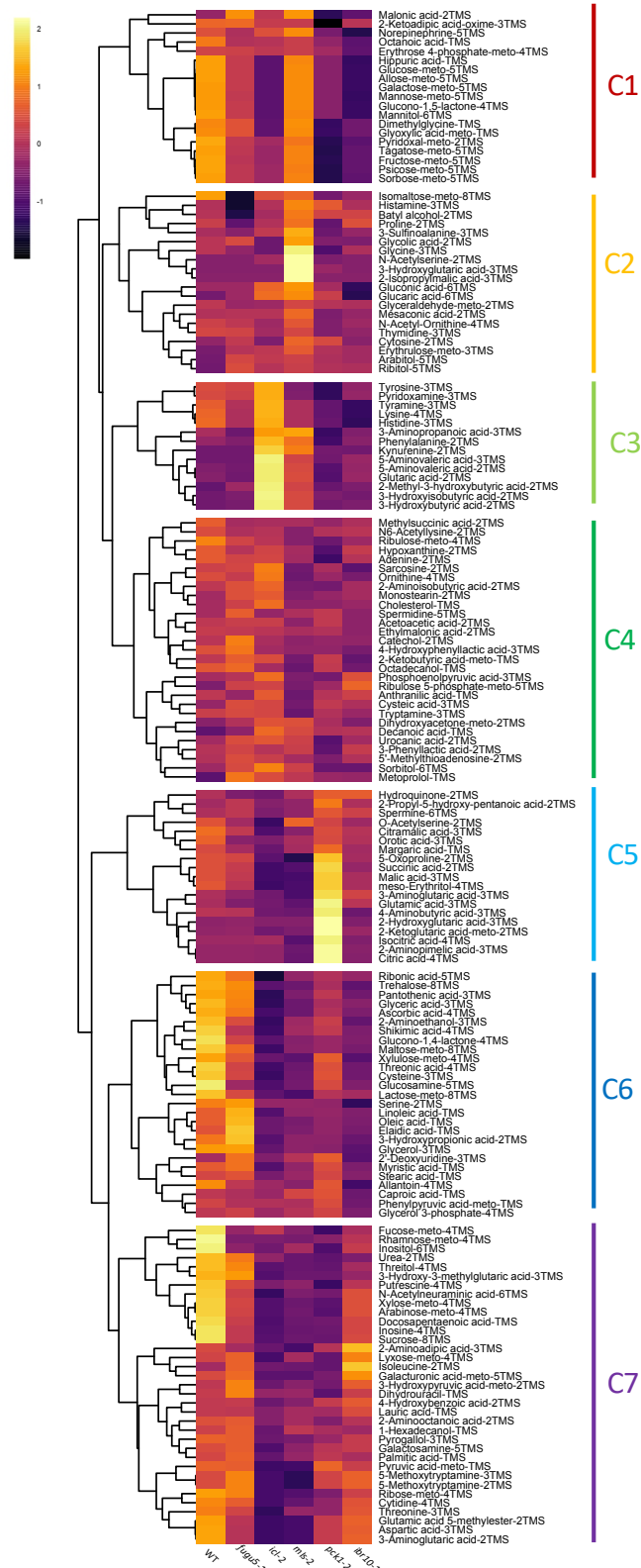
Supplementary Figure S3. SPAD values during the vegetative growth phase. (In support of Figure 4)

The SPAD value of a rosette leaf of *ibr10*. SPAD values were measured at 36 DAS ($n = 6$, biological replicates). Lower and upper boxes represent the first and third quartiles, respectively, and the line in the boxes represents the median. Each single biological replicate contains 10 technical replicates. DAS, days after seed sowing.



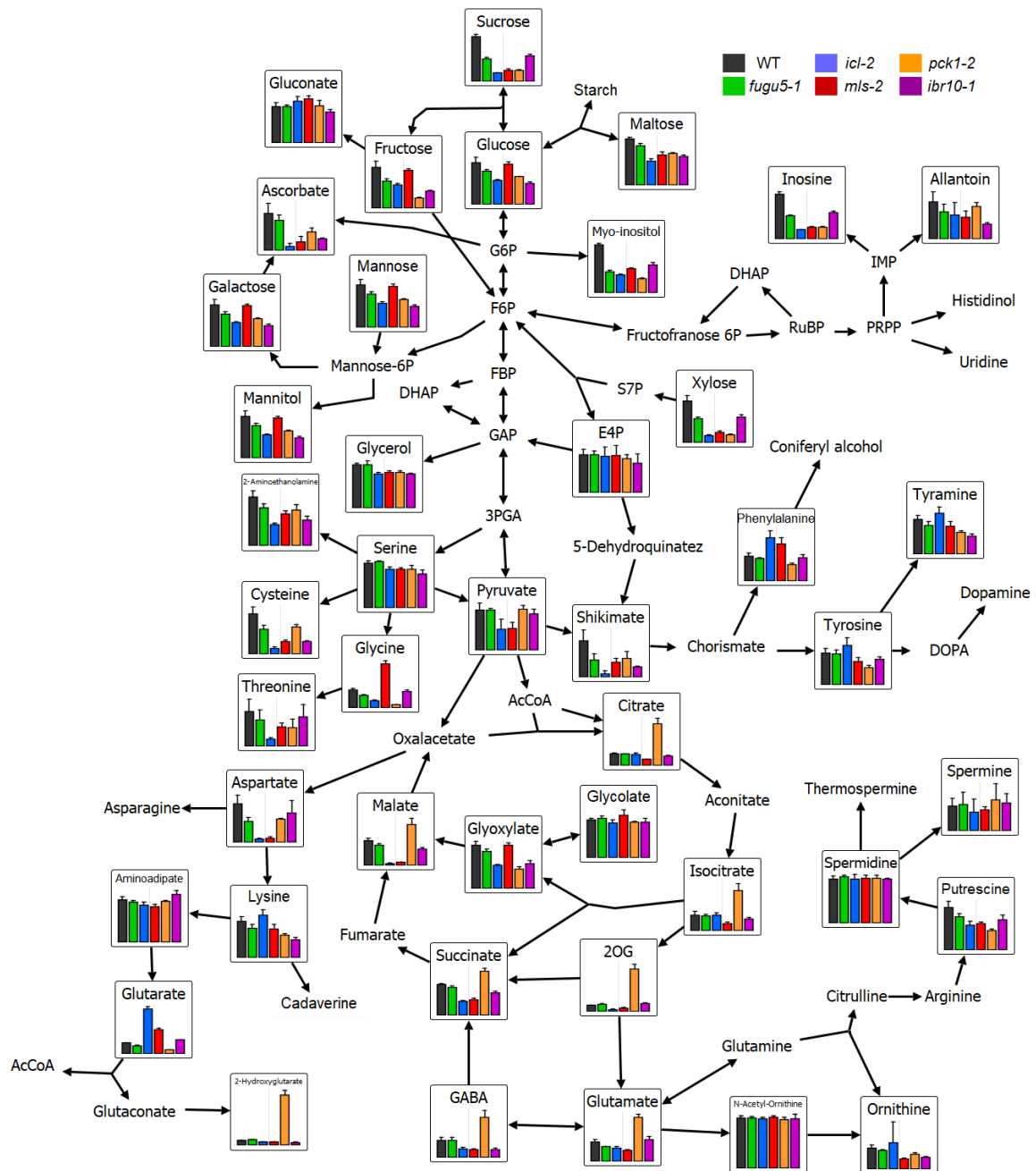
Supplemental Figure S4. Hypocotyl length of etiolated seedlings grown on MS medium with 2% sucrose. (In support of Figure 5A)

Phenotypes of etiolated seedlings grown on MS medium with 2% sucrose. (a) Photographs were taken at 3 DAI. Bar = 2 mm. (b) Data of bar plot are means + SD of 3 DAI etiolated seedlings ($n = 20$). N.S. indicates that mutants have no significant differences compared to the WT (Dunnett's test at $p < 0.05$; R ver. 3.5.1). DAI, day after induction of seed germination.



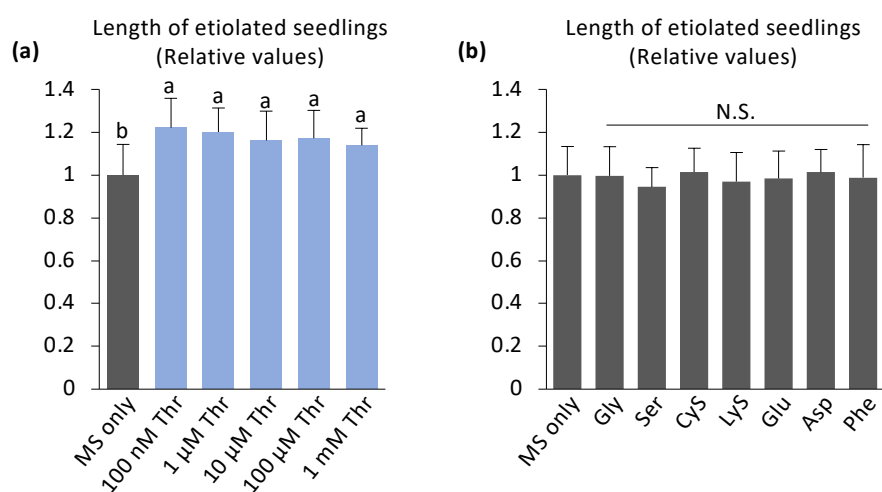
Supplementary Figure S5. Metabolic changes in Class II compensation mutants. (In supports Figure 5)

Hierarchical clustering analysis of Class II compensation-exhibiting mutants is shown. All 163 metabolites detected by GC-Qq-MS were categorized into seven clusters based on behavior. Data are Z-scores of the average contents relative to that of the quality control sample ($n = 4$). Three DAI etiolated seedlings. DAI, days after induction of seed germination.



Supplementary Figure S6. Pathway analysis of Class II compensation mutants. (In support of Figure 5)

Pathway analysis of Class II compensation-exhibiting mutants. Primary metabolites related to TAG-to-Suc conversion were plotted on the corresponding metabolic pathway. Data are means + SD ($n = 4$). Three DAI etiolated seedlings. DAI, days after induction of seed germination.



Supplemental Figure S7. Length of etiolated seedlings grown with exogenous amino acids (In support of Figure 5E)

(a) Length of etiolated seedlings grown for 3 DAI on Suc-free MS medium supplied with various concentrations of Thr. Data of bar plot are means + SD ($n = 20$). Letters indicate groups (Tukey's HSD test at $p < 0.01$; R ver. 3.5.1). **(b)** Length of etiolated seedlings grown for 3 DAI on Suc-free MS medium supplied with 100 μ M of the indicated amino acids. Data of bar plot are means + SD ($n = 20$). N.S. indicates that mutants did not show statistically significant difference compared to the WT (Dunnett's test at $p < 0.5$; R ver. 3.5.1). DAI, days after induction of seed germination.