

Table 1 Suppl. List of RT-qPCR primers used in this study. Protein ID according to the *Phytozome* database or Genbank accession number (\*).

Gene name	Protein ID	Sequence	
<i>S</i> -adenosylmethionine synthetase family	Glyma15g21890.1	forward	5'-GTGCTTCTGGAAGTTAAATGG-3'
		reverse	5'-TGATCTCTCCGAAAACCATC-3'
<i>S</i> -adenosylmethionine synthetase 1	Glyma03g38190.3	forward	5'-GACTGGCAGTATATCCAGTTACAG-3'
		reverse	5'-GATGCAAAAGAAGGGTGAT-3'
<i>S</i> -adenosylmethionine synthetase 2	Glyma19g40810.1	forward	5'-CTGCTTCTTCAGCTTGAGAAATG-3'
		reverse	5'-CAAAGACCATGACCATGTTGG-3'
ACC synthase	Glyma18g47280.1	forward	5'-CCTTAGACTCAGCTTCTCCAAC-3'
		reverse	5'-TCCTGTATATGTGGACCGCA-3'
ACC oxidase	Glyma05g36310.1	forward	5'-GGAAATTCGGTTGACCATT-3'
		reverse	5'-CAAAGATGGAGATCCCTGTGATA-3'
18S rRNA	X02623.1*	forward	5'-TGATTAACAGGGACAGTCGG-3'
		reverse	5'-ACG GTA TCT GAT CGT CTT CG-3'

Table 2 Suppl. List of 3 proteins commonly identified in 2-d-flooded soybean and 2-d-drought stressed soybean compared to control 2-d-old soybean. Protein ID, according to the *Phytozome* database; M.P. - number of matched peptide; Ratio - relative abundance of protein; 4(2) F - 4-d-old soybean treated with flooding for 2 d; 4(2) D - 4-day-old soybean treated with drought for 2 d. These proteins were identified by Oh and Komatsu (2015).

Protein ID	Description	M.P.	Ratio 4(2) F	<i>P</i> -value 4(2) F	Ratio 4(2) D	<i>P</i> -value 4(2) D
Glyma15g21890.1	<i>S</i> -adenosylmethionine synthetase family	4	0.4175	0.0127	9.0403	0.0383
Glyma03g38190.3	<i>S</i> -adenosylmethionine synthetase 1	3	0.4185	0.0213	10.6780	0.0101
Glyma19g40810.1	<i>S</i> -adenosylmethionine synthetase 2	3	0.4185	0.0213	10.6780	0.0101

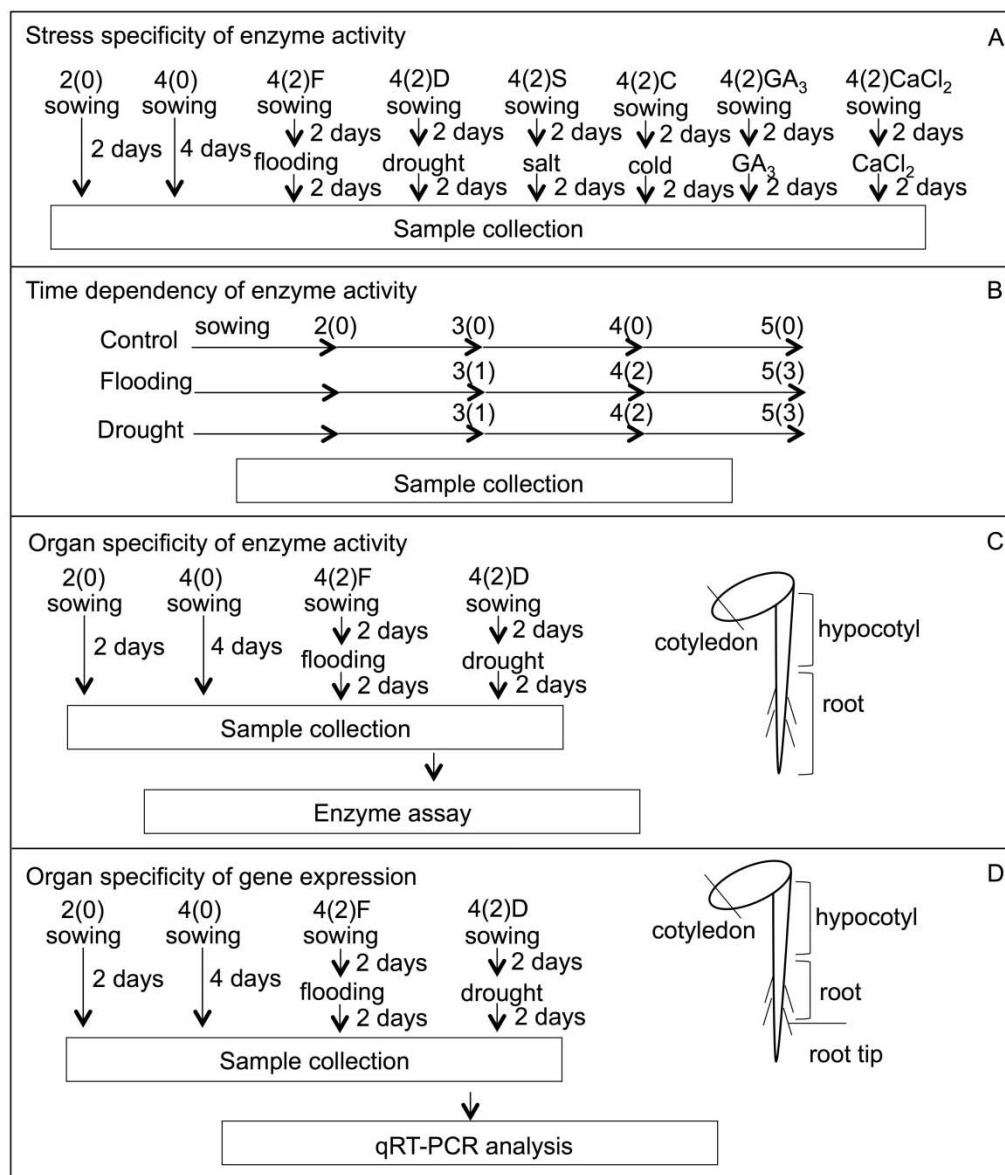


Fig. 1 Suppl. Experimental workflow of this study. To examine stress-specific enzyme activity, 2-d-old soybeans were treated with flooding, drought, salt, cold, GA<sub>3</sub>, or CaCl<sub>2</sub> for 2 d, and then root was collected (A). To investigate temporal changes in enzyme activity, 2-d-old soybeans were treated with flooding or drought for 1, 2, and 3 d, and then root was collected (B). To examine organ specificity of enzyme activity, 2-d-old soybeans were treated with flooding or drought for 2 d, and then root, hypocotyl, and cotyledon were collected (C). To examine organ specificity of gene expression, 2-d-old soybeans were treated with flooding or drought for 2 d, and root tip, root, hypocotyl, and cotyledon were collected (D). Three independent experiments were performed as biological replicates.

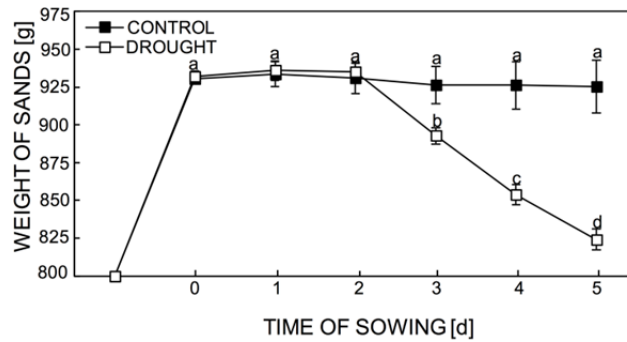


Fig. 2 Suppl. Reduction of the sand mass under drought stress. In this study, silica sands (800 g) was used in a plastic case (180 × 140 × 45 mm) and 150 cm<sup>3</sup> of water was added (0 day). Additional 50 cm<sup>3</sup> of water was supplied to maintain the normal condition for soybean for next 2 d. For control, 50 cm<sup>3</sup> water was supplied on days 3, 4 and 5; for drought, water was withheld for 3 d. All the plastic cases were kept in growth chamber under irradiance of 160 μmol m<sup>-2</sup> s<sup>-1</sup>, 16-h photoperiod, and a temperature of 25 °C. Data are shown as means ± SD from 3 independent biological replicates. Means with different letters indicate significant changes according to Duncan's multiple comparison test ( $P < 0.05$ ).

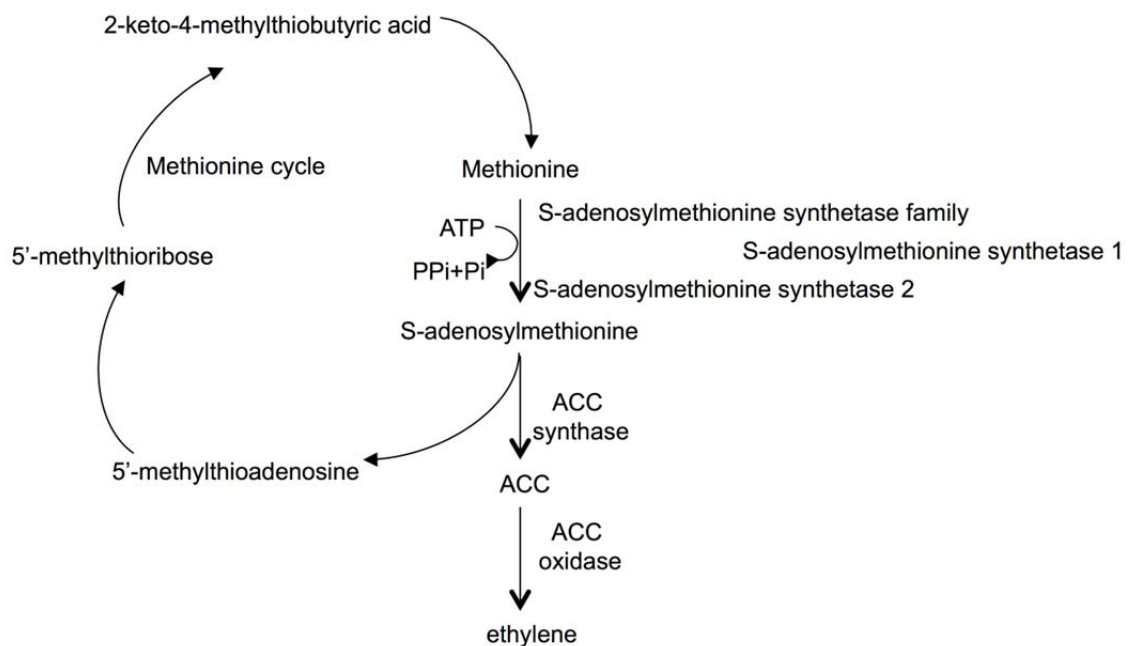


Fig. 3 Suppl. Pathway of ethylene biosynthesis. *S*-adenosylmethionine synthetase converts methionine to *S*-adenosylmethionine which is further converted into ACC by ACC synthase. Ethylene is converted from ACC by ACC oxidase (Wang *et al.* 2002).

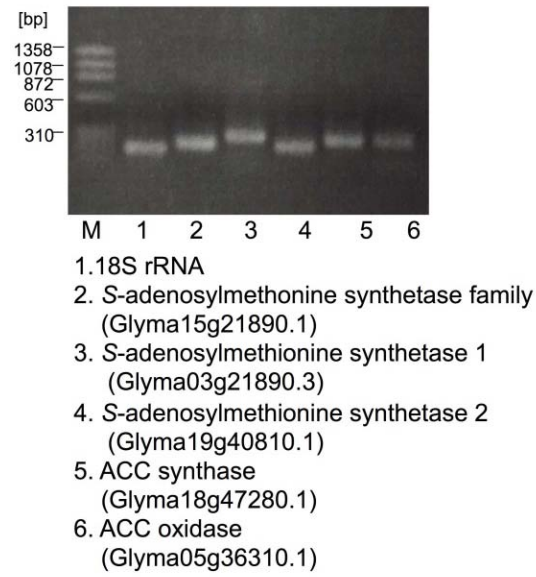


Fig. 4 Suppl. Agarose gel electrophoresis of RT-qPCR products obtained with the primers listed in Table 1 Suppl. They were visualized in agarose gel stained by ethidium bromide. M indicates marker.