## **Supplementary material**

 Table S1 Extraction of proline and total protein procedure.

Test name	Procedure	Reference
Proline	After separately homogenizing 0.1 g of root and shoot tissues	Bates et al.
	in 2 ml of 3% sulphosalicyclic acid solution and centrifuging	(1973)
	for 5 min, 1 ml of supernatant was mixed with an equal	
	volume of acid ninhydrin solution and glacial acetic acid and	
	finally incubated at 100 °C for 1 h. The reaction was	
	terminated by cooling in an ice bath, and toluene was added	
	to the reaction mixture and vortexed for 15 sec. The intensity	
	of the chromophore product containing toluene was	
	measured at 520 nm. Proline in fresh tissue was determined	
	according to the standard curve.	
Total protein	Total proteins from harvested tomato plant leaf samples were	(Bradford,
	measured with the Bradford method. For the estimation of	1976)
	proteins in an unknown sample, 100 µl of sample extract was	
	mixed with 5 ml of Bradford solution and incubated for 10	
	min. Finally, absorbance was read at 595 nm. Protein	
	contents were interpreted from the standard calibration curve	
	for proteins	