

Supplemental Table S1. List of transformation vectors.

Vector names	Descriptions
pPZP211- <i>pCER6::TSF</i>	<i>CaMV35S</i> in pPZP211 was replaced by a genomic DNA fragment containing the region from 1,951 bp upstream to 1 bp upstream of the start codon of <i>CER6</i> using <i>HindIII</i> and <i>XbaI</i> sites to obtain pPZP211- <i>pCER6</i> ; then the full length coding sequence (528 bp) of <i>TSF</i> was cloned into pPZP211- <i>pCER6</i> at <i>XbaI</i> sites.
pPZP211- <i>pCER6::CO</i>	The full-length coding sequence (1,122 bp) of <i>CO</i> was cloned into pPZP211- <i>pCER6</i> at <i>XbaI</i> sites.
pPZP211- <i>pCER6::GI</i>	The full-length coding sequence (3,522 bp) of <i>GI</i> was cloned into pPZP211- <i>pCER6</i> at <i>XbaI</i> sites.
pPZP211- <i>CaMV35S::CRY2-GFP</i>	Full-length cDNA of <i>CRY2</i> (1,836 bp) was amplified by PCR and the <i>PHYB</i> moiety of pPZP211/35S- <i>PBG-nosT</i> (Matsushita et al., 2003) was replaced using <i>XbaI</i> and <i>ClaI</i> sites to obtain pPZP211- <i>CaMV35S::CRY2-GFP</i> .