

## Supplementary materials

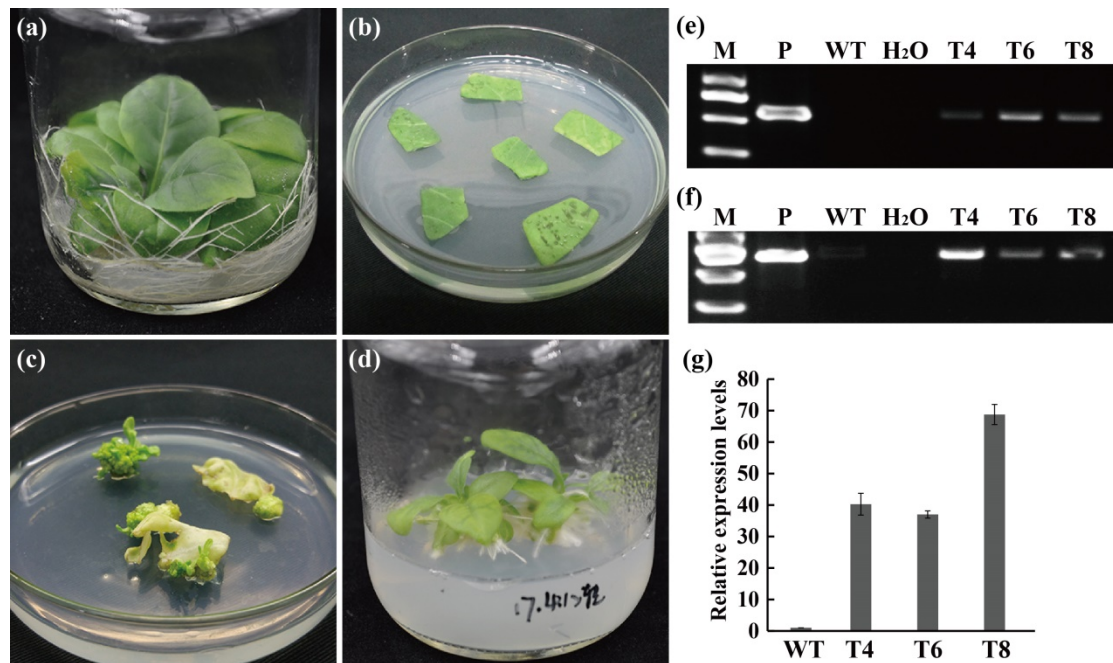
**Table S1** Primer sequences used in this study

Gene	Purpose	Forward primers (5'–3')	Reverse primers (5'–3')
<i>PtrLEA7</i> - pBI121	Overexpression	GCT <u>CTAGA</u> ATGGCATCCCACGA TCAGAG	CGGGATCCTTAACCCT TTGTATAAGTGT
<i>PtrLEA7</i> - 101LYFP	Subcellular localization	CGGAATTCATGGCATCCCACGA TCAGAG	CGGGATCCACCCTTTG TATAAGTGTTC
35S-F <i>NPT II</i>	Transgenic identification	TCCTCGGATTCCATTGCCCAGC CGGCTATGACTGGGCACAACA	CGGCAGGAGCAAGGT GAGATG
<i>PtrLEA7</i> - qPCR <i>CAT</i>	qPCR	GAAAACCCAGGAGACGGGAC CTGCCAGTTCTTTCAACGCC	CCGAAAGTGTCTTCA CGGC AAACCCTTAGCACTGG CTCC
<i>SOD</i>		CAAGCCGCAATAGCAGCCAT	CCGTGGAATGCAGAGT GAAG
<i>POD</i>		TGCTGGACAGCACTAACACC	CCAATGGAAGTGGCCCA TGAC
<i>ACTIN</i>	Internal reference	CCGACCGTATGAGCAAGGAAA	TTCCTGTGGACAATGG ATGGA
<i>Ubiquitin</i>		GGTGTTCAGTGGCGGACG	TCCTCCCCTCAGCTAC GGGGTAT

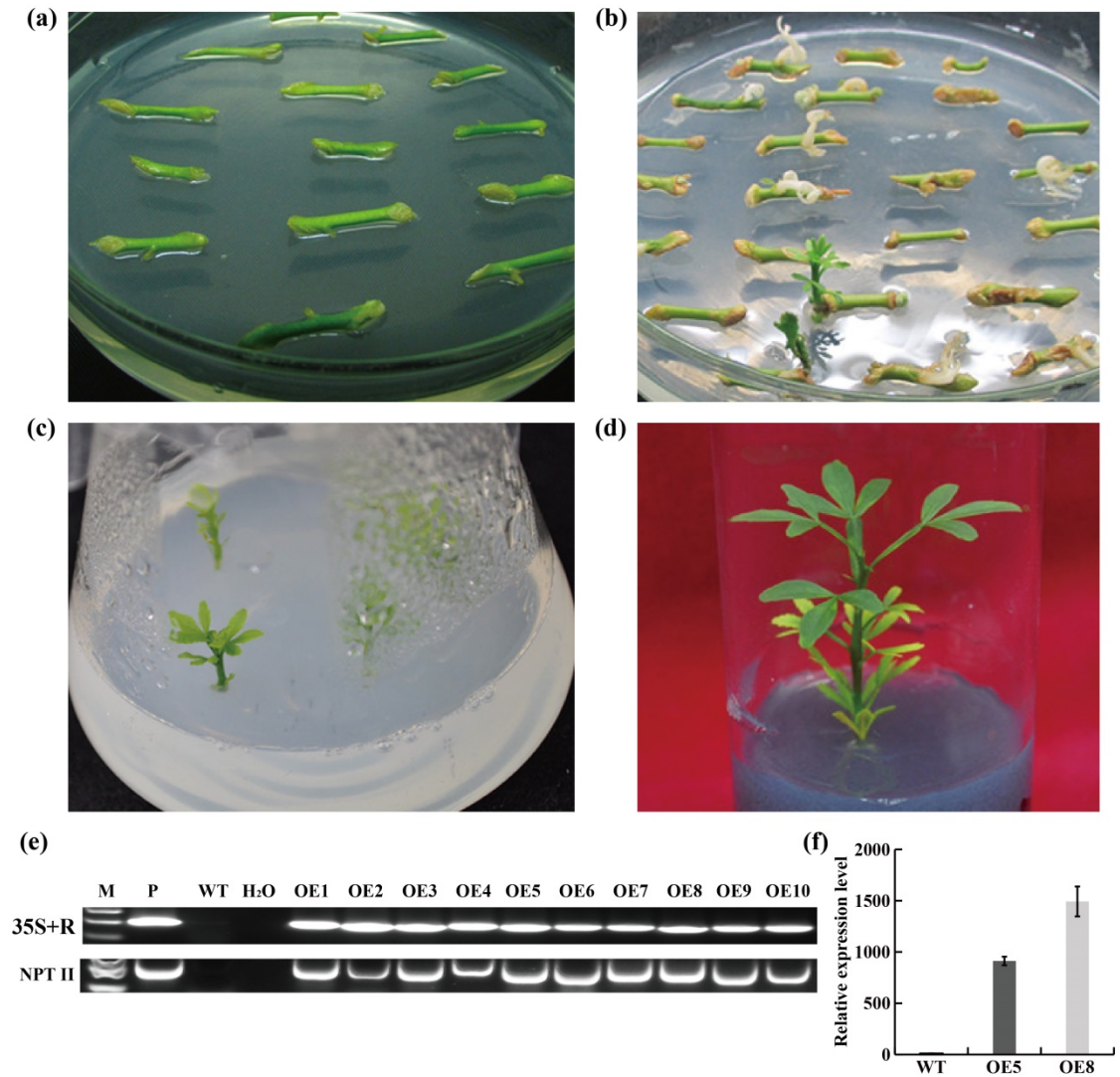
Note: Underlines indicate enzyme site.

**Table S2** Prediction of subcellular localization of *PtrLEA7* by two online tools

Location	Plant-mPLoc	YLoc <sup>+</sup>
Cytoplasm	–	43.3%
Nucleus	Nucleus	51%
Extracellular space	–	5.6%



**Fig. S1** Genetic transformation and molecular identification of transgenic tobacco plants overexpressing *PtrLEA7*. (a) Wild type tobacco plants used for transformation. (b) Leaf discs on the selection medium. (c) Regenerated buds on the selection medium for 40 d. (d) Representative rooted plants. (e,f) Transgenic identification of three transgenic lines (T4, T6, T8) by PCR using 35S-forward + *PtrLEA7*-reverse primers (e) and *NPT II* primers (f). M, marker; P, plasmid; WT, wild type. (g) Gene expression levels of *PtrLEA7* in three transgenic lines and wild type by qPCR.



**Fig. S2** Genetic transformation and molecular identification of transgenic trifoliate orange overexpressing *PtrLEA7*.

(a) Co-culture of the shoot segments. (b) Regenerated shoots on the selection medium. (c) Regenerated plants on the shooting medium. (d) Representative rooted plants. (e) Genetic identification of ten transgenic lines (OE1-OE10) and wild type by PCR using two pairs of primers. M, DNA marker; P, plasmid; WT, wild type. (f) Relative expression levels of *PtrLEA7* in two transgenic lines and wild type by qPCR.