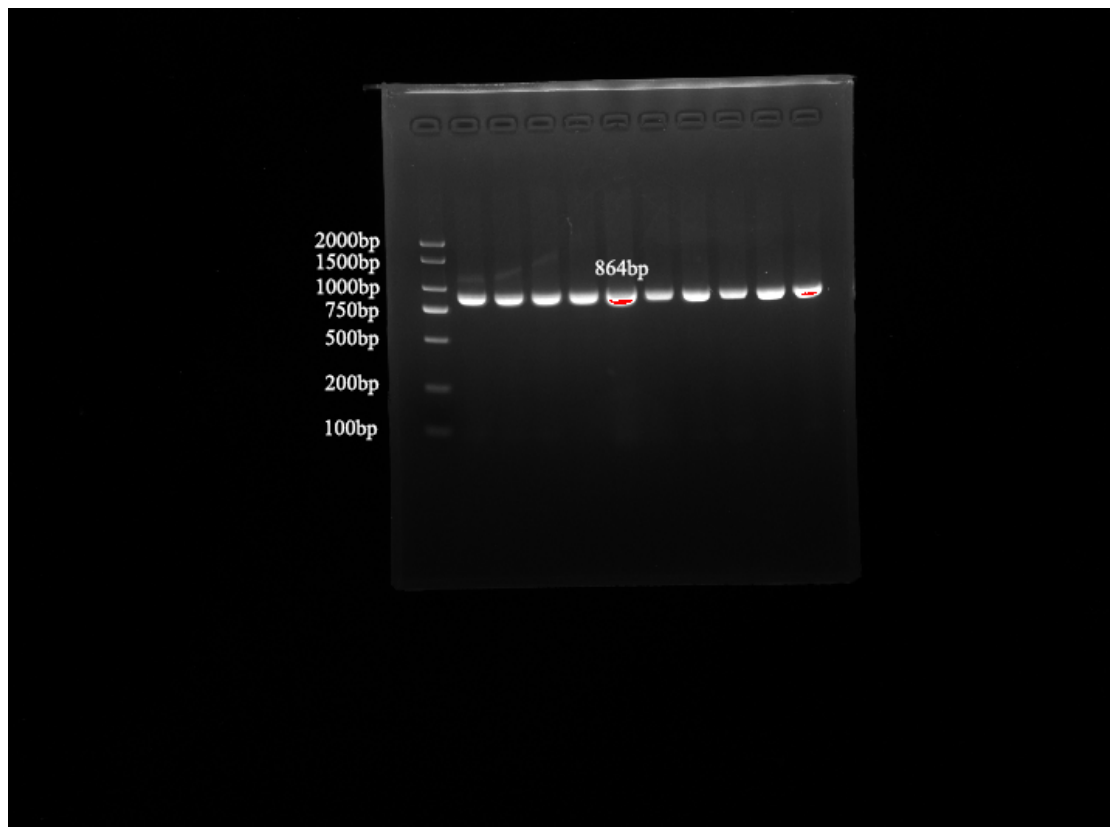


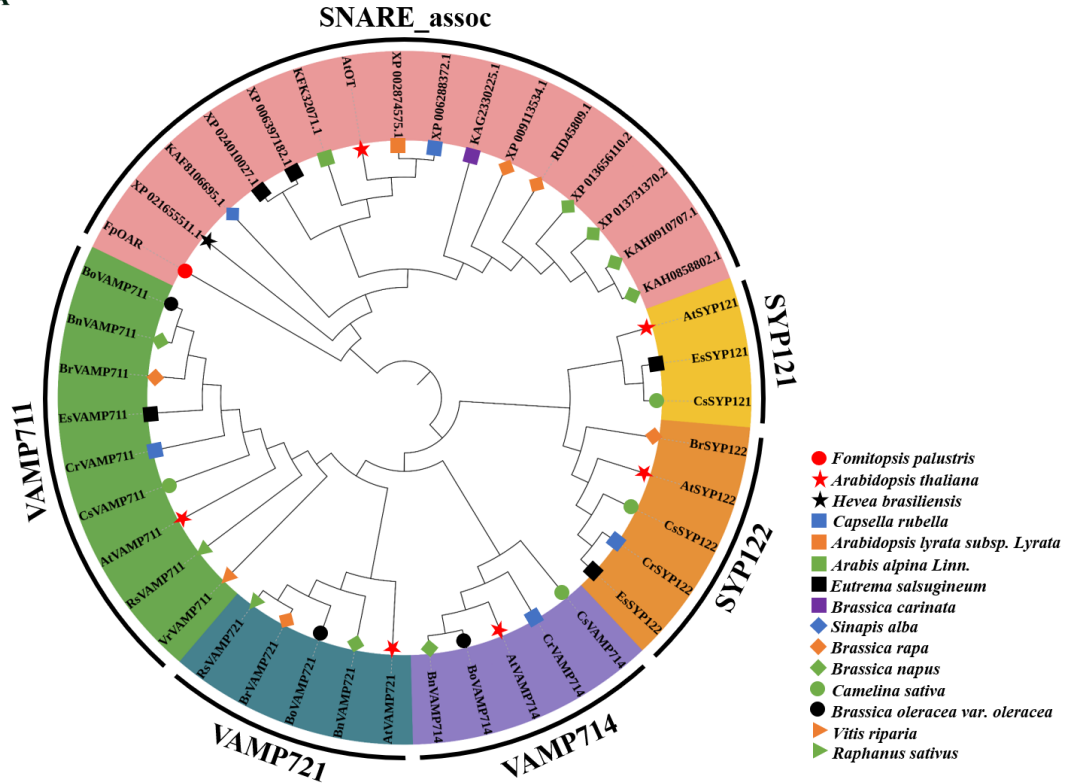
Supplementary Table S1 Primers used in the experiments.

Primer name	Primer sequence (5'-3')	Description
<i>AtOT</i>	F: ATGGCGGCTCCTCGG R: TCATTCATATACTCTCTTTCTCT	For ORF sequence cloning
<i>AtOT</i> (pCAMBIA1300)	F: TCATTTGGAGAGAACACGGGGGACTATGGCGGCTCCTCGG R: CCTCGCCCTTGCTCACCATGTCATTCATATACTCTCTTTC	For subcellular localizaton
<i>AtOT</i> (pDR196)	F: TTGGGTACCGGGCCCCCCTCGAGGATGGCGGCTCCTCGGAATTT R: CTAGTGGATCCCCCGGGCTGCAGGTCATTCATATACTCTCTTTCTCT	For yeast heterologous expression vector construction
<i>FpOAR</i> (pDR196)	F: GGTACCGGGCCCCCCTCGAGGATGACCGACCTGCATCGAAG R: CTAGTGGATCCCCCGGGCTGCAGGTCAGAGAAGATCTTCTTGCC	
q- <i>AtOT</i>	F: GGC GGCTCCTCGGAAT R: GGCAGGCATGGTGAGGT	For qRT-PCR
q- <i>AtActin2</i>	F: TGCCAATCTACGAGGGTTTC R: TTCTCGATGGAAGAGCTGGT	
<i>atot</i>	LP: ATGTTAAGTTCTTCAGTTTC RP: TCAGAATCACCCATATTTGGTCAAA LBb1.3: ATTTTGCCGATTTTCGGAAC	For homozygous identification

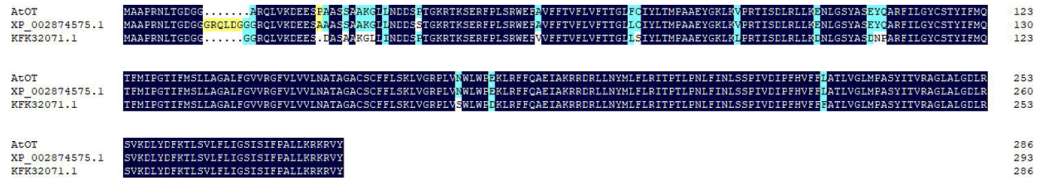


Supplementary Figure S1. The PCR amplification products of the ORF sequences of

A



B



Supplementary Figure S2. Phylogenetic analysis of FpOAR, AtOT and SNARE protein members in other plants (A) and multiple sequence alignment of AtOT, XP_002874575.1 and KFK32071.1 (B).