



Supplemental Figure 1. RT-PCR and expression of wild-type (WT) Col-0 and *vip1-2* RNA. A. A 138 bp PCR product was amplified from the 3' end of the *VIP1* and *vip1-2* transcripts and visualized by ethidium bromide staining after electrophoresis through a 1.5% agarose gel. M, size marker; **B.** Quantitative RT-PCR of *vip1-2* gene expression relative to that of *VIP1* in wild-type plants. Results show the average of three technical replicates \pm SE. **C.** Expression and localization of VIP-Venus (left) and *vip1-2*-Venus (right) in tobacco BY-2 protoplasts. Constructs comprised of cDNAs encoding VIP-Venus or *vip1-2*-Venus (*vip1-2* out of frame with Venus) and a nuclear marker mRFP-NLS were co-transfected into tobacco BY-2. Transfected cells were imaged by confocal microscopy 16 hours later. Four images of each cell are presented (clockwise from top left: merged YFP, mRFP, and DIC; YFP; YFP + mRFP; mRFP). Bars indicate 20 μ m.