

Supplemental Figure S1. Detection of nonspecific protein band in α -GFP western blot. The indicated proteins were expressed in PtoR tomato protoplasts for 16 hrs, total protein extracted, and 40 µg of total protein analyzed by α -GFP western blot. A nonspecific protein band indicated by the asterisk can be detected in every sample including GFP alone and the empty vector. Filled triangle = GFP-Adi3 proteins; open triangle = GFP-Adi3^{\Delta T-loop} proteins; gray triangle = GFP; asterisk = nonspecific protein; open diamond = RuBisCo loading control.



Supplemental Figure S2. HA-Adi3 subcellular localization and reduction of GFP-Adi3 nuclear localization when expressed with GBP. *A*, Western blot of cellular fractionation of HA-Adi3 expression in tomato protoplasts. Western blots were carried out with the α -HA antibody and are representative of three independent experiments. *B*, Reduction of nuclear GFP-Adi3 by GBP. The indicated proteins were expressed in tomato protoplasts for 16 hrs, nuclei isolated from each sample, and an α -GFP western blot carried out. The GBP protein is not capable of reducing GFP nuclear localization but does show a strong reduction of nuclear localized GFP-Adi3. The GBP protein does not have a GFP tag and thus is not detected in the western blot. Filled triangle = GFP-Adi3; gray triangle = GFP.



Supplemental Figure S3. The number of punctate structures from deletion of the Adi3 T-loop extension. The indicated *GFP-Adi3* constructs were expressed in tomato protoplasts for 16 hrs., viewed by confocal microscopy, and the number of GFP positive punctate structures counted in a minimum of three cells for each construct. The data from five independent experiments were averaged and shown with standard error.



Supplemental Figure S4. Western blot detection of AvrPto expression after transformation of protoplasts transiently expressing Adi3. PtoR protoplasts transiently expressing GFP-Adi3 for 14 hrs were transformed with a *BFP-avrPto* construct, proteins expressed for an additional 3 hrs, and the presence of Adi3 and AvrPto proteins analyzed by α -GFP western blot. Filled triangle = GFP-Adi3; open triangle = BFP-AvrPto, gray triangle = RuBisCo loading control.



Supplemental Figure S5. Overexpression of Adi3 can suppress H_2O_2 cell death induction in yeast. *A*, *GFP-Adi3* was expressed from the galactose inducible yeast expression vector pPS808 overnight at 30°C in the presence of glucose, switched to galactose media for 6 hrs, switched back to glucose media in the presence of 0 or 5 mM H_2O_2 for 2 hrs, and plated on YPD plates in serial dilutions. Galactose induced expression of the *Pst* effector protein GFP-AvrPtoB was used as a positive CDS control (48). *B*, Western blot detection (α -GFP) of Adi3 and AvrPtoB galactose induced yeast expression. Filled triangle, GFP-Adi3; open triangle, GFP-AvrPtoB.