

RNA-Mediated Gene Silencing Signals Are Not Graft Transmissible from the Rootstock to the Scion in Greenhouse-Grown Apple Plants *Malus* sp.

Supplementary Information

Figure S1. Molecular evaluation of *hrp-gusA* transgenic apple clones by PCR (A) and RT-PCR; (B) on genomic DNA and cDNA, respectively. H₂O, blank probe; plasmid, positive control pHELLSGATE::*hrp-gusA* (the vector PCR 2.1 Topo containing a genomic clone of EF1alpha was used as control for the EF1a RT-PCR), PinS, descendent of the cultivar “inova”. Transgenic clone T667 was separately tested by RT-PCR (data not presented).

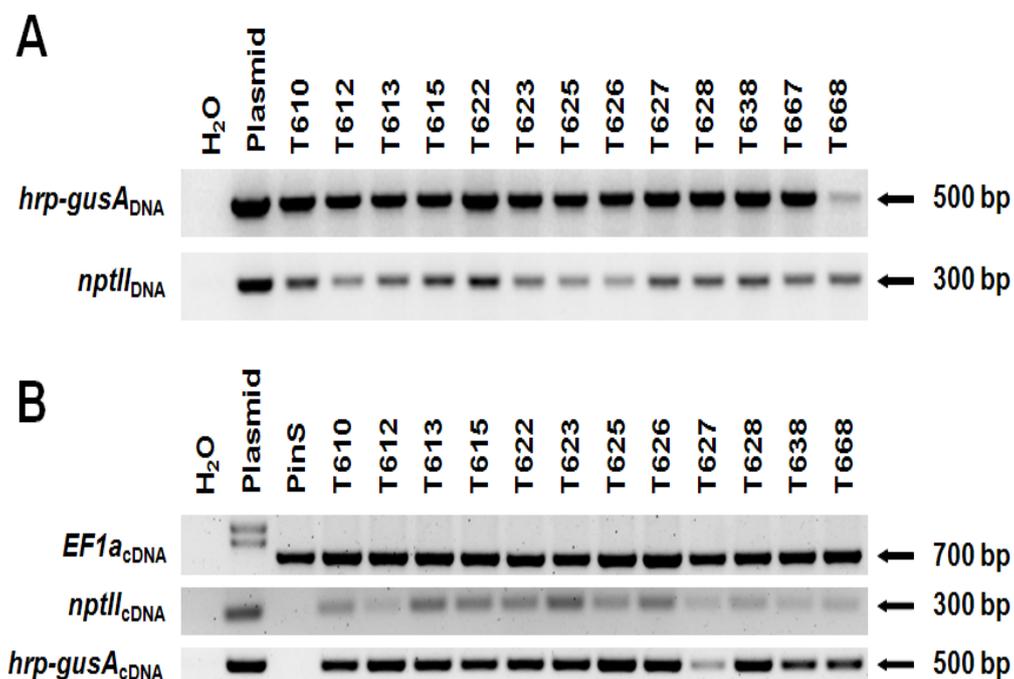
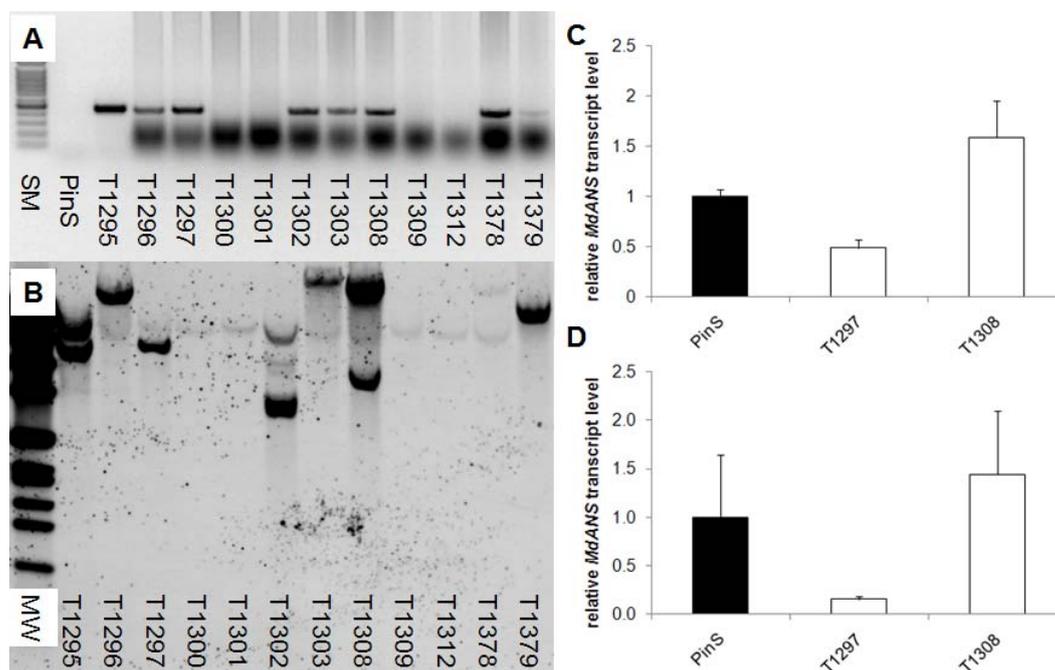


Figure S2. Molecular evaluation of *hrp-Mdans* transgenic plants. (A) PCR based detection of the *hrp-Mdans* hairpin gene construct on genomic DNA using the primers HG1 and HG3n; (B) Southern blot hybridization of *Sac*I digested DNA using a DIG-labeled probe of the *nptII*; (C) Relative mRNA expression of the *Mdans* gene measured on *in vitro* grown plants (mean of three biological replicates, each with three technical replicates); (D) Relative mRNA expression of the *Mdans* gene measured on greenhouse grown plants (mean of three biological replicates, each with three technical replicates). SM, molecular size marker (100 bp DNA ladder, MBI Fermentas); MW, 1 kbp, molecular weight marker II, DIG labelled (Roche); “PinS”, descendant of the apple cultivar “Pinova”.



© 2012 by the authors; licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution license (<http://creativecommons.org/licenses/by/3.0/>).