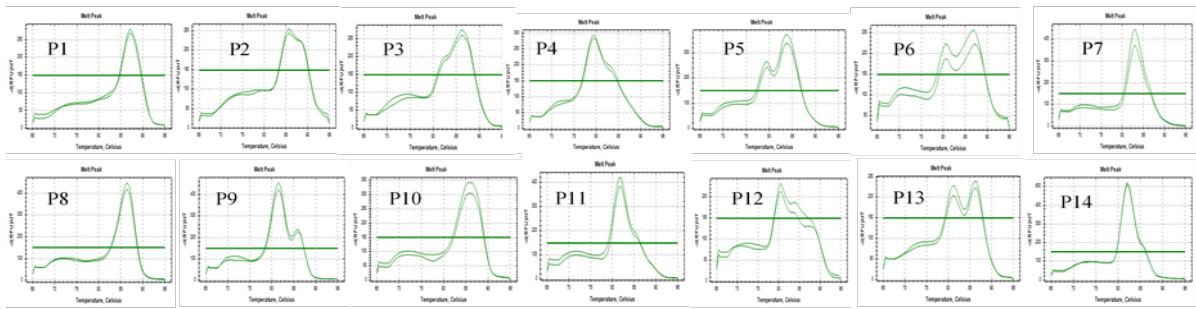


At 56°C



At 60°C

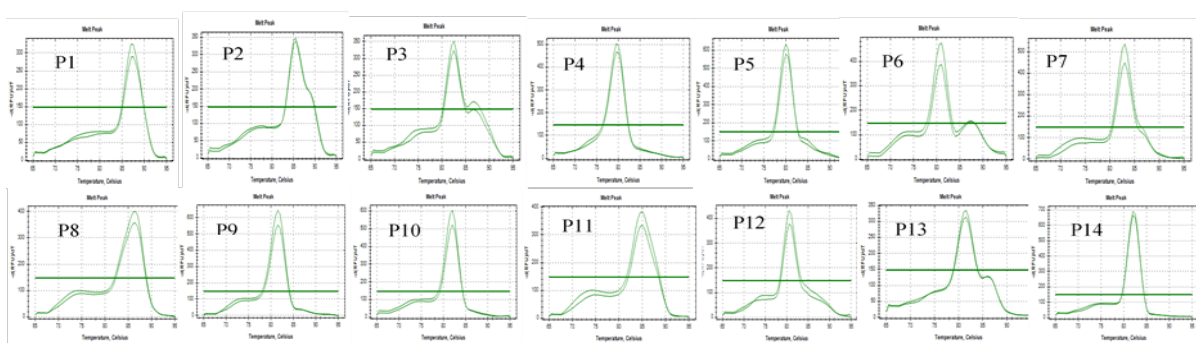


Fig. S1 Melting curves of the genomic DNA extracted from *S. hermonthica* seeds mixed into Dutch agricultural soil. qPCR analysis was performed with 14 different primer sets at two different annealing temperatures (see Figure 1 for details on the primer sets tested).

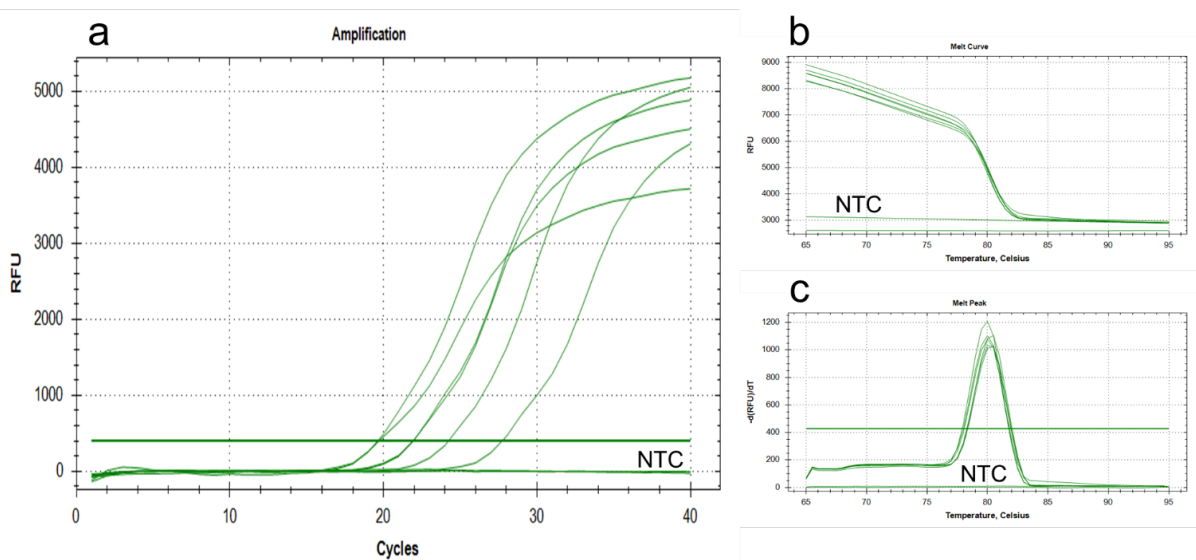


Fig. S2 qPCR detection of *Striga* seeds. (a) Detection of five *S. hermonthica* biotypes and one *S. asiatica* biotype collected from different regions of Ethiopia by qPCR. (b) Melt curve and (c) Melt peak associated with the detection of the samples by qPCR. The single peak corresponds to specific amplification of the target gene. NTC represents non-templated control.