

**FIGURE S1.** PCR amplifications of *R. etli* pSym markers in the TER strains.

To determine the presence of markers of the *R. etli* pSym in the putative transconjugants of endophytic recipients, DNA was isolated from the different strains and used as template to produce PCR products for nodD1 (A), nodA (B), fixNd (C), nifH (D), and GFP (E), from the pSym of R. etli CFNX182-1. The lane labeled 182-1 corresponds to strain CFNX182-1, MWM – molecular weight markers.