

Figure S1. Construction and validation of the recombinant viral vector pTRV2-CsPOR1 and pTRV2-CsTCS1. (A,C) Construction and validation of the recombinant viral vector pTRV2-CsPOR1. (B,D) Construction and validation of the recombinant viral vector pTRV2-CsTCS1.

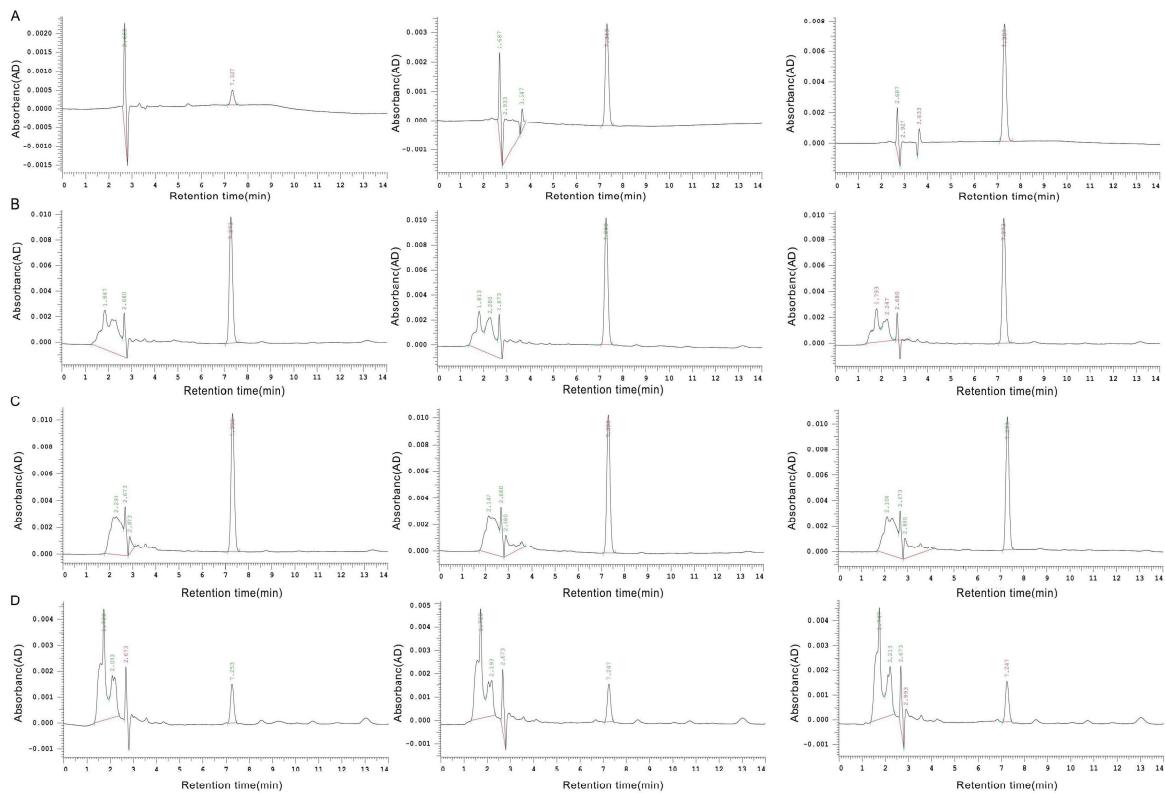


Figure S2. HPLC analysis of caffeine content in the leaves of CsTCS1 silent plants. (A) Peak diagram of caffeine standard ( $10 \mu\text{g}\cdot\text{mL}^{-1}$ ,  $50 \mu\text{g}\cdot\text{mL}^{-1}$ ,  $100 \mu\text{g}\cdot\text{mL}^{-1}$ ); (B) untreated tea plants; (C) infection with pTRV1 + pTRV2; (D) infection with pTRV1 + pTRV2-CsTCS1. HPLC, high-performance liquid chromatography.

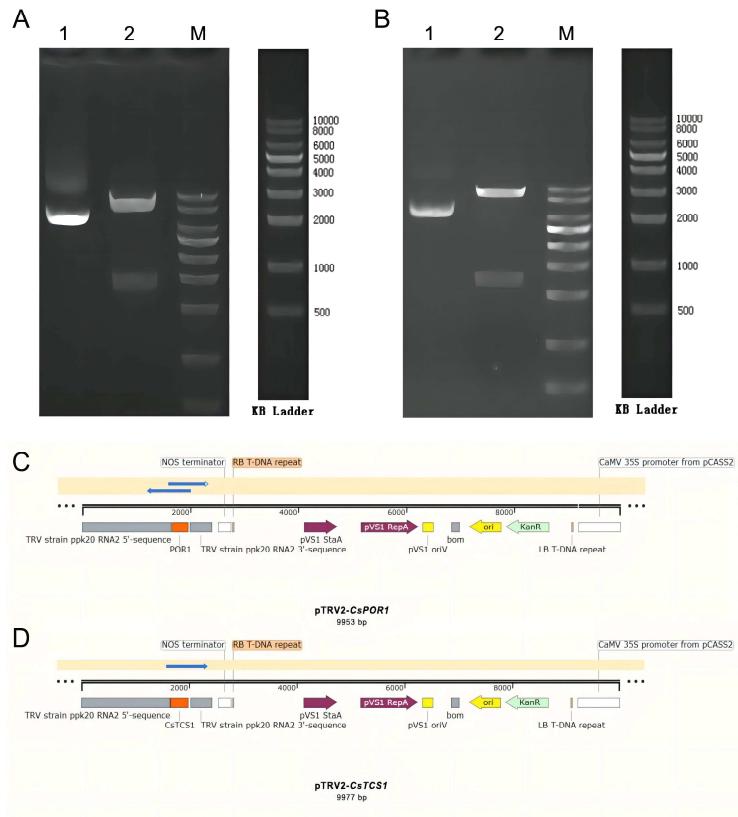


Figure S3. Comparison of tea plants before and after infection with pTRV1 + pTRV2-CsPOR1 *Agrobacterium*. (A,B) Image of tea plants before the pTRV1 + pTRV2-CsPOR1 *Agrobacterium* infection. (C,D) Image of tea plants after the pTRV1 + pTRV2-CsPOR1 *Agrobacterium* infection.

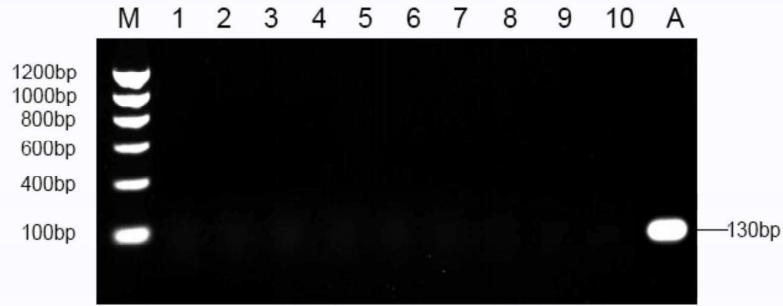


Figure S4. Detection of Agrobacterium contamination. 1, uninfected seedlings; 6, uninfected cuttings; 2 and 7, seedlings infected with pTRV1 + pTRV2; 3 and 8, cuttings infected with pTRV1 + pTRV2; 4, seedlings infected with pTRV1 + pTRV2-CsPOR1; 5, cuttings infected with pTRV1 + pTRV2-CsPOR1; 9, seedlings infected with pTRV1 + pTRV2-CsTCS1; 10, cuttings infected with pTRV1 + pTRV2-CsTCS1. A, positive control. M: DL1200 marker.