

**Table 1**  
**Primer Sequences for some Commonly Used Promoters and Reporter Genes**

Gene/promoter	Primer sequences	Size of PCR fragment, bp
<i>npt-II</i> gene	5' TCT CAC CTT GCT CCT GCC 3' (forward primer) 5' AGG CGA TAG AAG GCG ATG C 3' (reverse primer)	464
<i>gusA</i> gene	5' AGC ATC TCT TCA GCG TAA GG 3' (forward primer) 5' TGA ACA ACG AAC TGA ACT GG 3' (reverse primer)	611
<i>nos</i> promoter	5' GAC AAG CCG TTT TAC CGT TTG GAA CTG 3' (forward primer) 5' CTG CAG ATT ATT TGG ATT GAG AGT G 3' (reverse primer)	270
CaMV35S promoter	5' CTA CTC CAA AAA TGT CAA AGA TAC AGT C 3' (forward primer) 5' GGG CTG TCC TCT CCA AAT G 3' (reverse primer)	370 <sup>a</sup>
non-tDNA RB <sup>b</sup>	5' CGC TCT TTT CTC TTA GGT TTA 3' (forward primer)	
<i>npt-II</i> 5'	5' GTC ATA GCC GAA TAG CCT C 3' (reverse primer)	

<sup>a</sup>Expected size for a single 35S promoter. If a double 35S promoter is present within the construct, two PCR products will be obtained, one of 370 bp and the other 775 bp.

<sup>b</sup>This primer can be used in conjunction with the *npt-II* 5' primer or a construct-specific primer to check for *Agrobacterium* contamination.