

**Table 1.** Oligonucleotide primers used in this study.

Number/name	Sequence (5'-3') <sup>a,b</sup>
1 1085 (+)	GTAAGCTTTTGCACCTGGAGGGCGACTG
2 1120 (+)	GTAAGCTTTTGAATCGCTTCTCAGCCC
3 1146 (+)	GTAAGCTTAGCTTGAGCCGAGAACGAGGC
4 1162 (+)	GTAAGCTTGAGCGGAGTGACCGGG
5 1177 (+)	GTAAGCTTCGTGCGCGGACACCTGCTAC
6 1192 (+)	GTAAGCTTGAGGAGCGGAGTAGAGAACGGC
7 1200 (+)	GTAAGCTTTCCCAGCCTGAGGGGCTGC
8 1222 (+)	GTAAGCTTGTGAAGCGAGAAGCTGCC
9 1235 (+)	GTAAGCTTGCGGCTCTCTCTGAGGAGG
10 1245 (+)	GTAAGCTTAAATCCAAGGCTATCATTGAGG
11 1412 (-)	TTCTCGAGCTGGCCTTCAGGTAGAAATTCC
12 1600 (-)	TTCTCGAGGTTGTGGTCAGACTCCTCCTC
13 S1232A (+)	CCTACCCCCAGTGG <b>CC</b> CCCCCTGAAGGCG
14 S1232A (-)	CGCCTTCAGGGGG <b>GGC</b> ACTGGGGGTAGG
15 S1239A (+)	GCTCTC <b>GCT</b> GAGGAGGAGTTAGAG
16 S1239A (-)	CCTCCTC <b>AGCG</b> AGAGCCGCC
17 KK1245/6EE (+)	AG <b>GAGGAAT</b> CCAAGGCTATCATTGAGG
18 KK1245/6EE (-)	CCTTGGA <b>TTCTCCT</b> CCTCTAACTCCTCCTC
19 LL1255/7AA (+)	T <b>GCC</b> CATGCCAATGACATGAAAGAGGCAG
20 LL1255/7AA (-)	GTCATT <b>GGC</b> AT <b>GGC</b> ATATTCCTCAATGATAGCC
21 L1243G (+)	GGAGGAG <b>GG</b> AGAGAAGAAATCCAAGG
22 L1243G (-)	CTTCTCT <b>CCCTCCTCCT</b> CAGAGAGAGC

<sup>a</sup> restriction sites used for cloning are underlined

<sup>b</sup> mutated nucleotides are in bold Italic