

Table 6. Primers used in this study (restriction sites underlined)

Primers used to clone <i>dgp</i> genes for expression in <i>B. fragilis</i>	
<i>dgpA</i> forward	AT <u>GGATCC</u> GGCAATCCAATAGAAGAGCTTATTA
<i>dgpA</i> reverse	CAGGATCCACGGAAATTCGATCAGTTTTCTAC
<i>dgpC</i> forward	AC <u>GGATCC</u> TTTAATAGCAAACGACGATTTCA
<i>dgpC</i> reverse	CTGGATCCGATTCTCCCATCTCTTGACCTATC
<i>dgpE</i> forward	TTGGATCCAACCTTTTTCTAAACAACCCCTGTT
<i>dgpE</i> reverse	GAGGATCCGAATCCCGGCGTTAGTATACAG
Primers used to create His-tagged DgpA and DgpC	
<i>dgpA</i> forward	TTGGATCCTATGAACAAAAAGTTTTCTACGATT
<i>dgpA</i> reverse	TGGATCCTTATTTAACGATTGTCTTAACAGCCTCT
<i>dgpC</i> forward	GGGATCCTATGAACAAAAAGTTTTCTACTCTTTTGG
<i>dgpC</i> reverse	CCGGATCCTTATTTAACAACTTTAACAGCTTCC
Primers used to demonstrate promoter inversion	
<i>dgpB</i> forward	ACGGATGAATTATAGGTTTTAGCC
<i>dgpB</i> reverse	GTATTCAATTCATCAGCACCTGAG
<i>dgpB</i> center	AGGTACGCGTTTCTTCGATACTT
<i>dgpC</i> forward	ATAGAGCAAGCTACTGCCGATAAT
<i>dgpC</i> reverse	GACCTTGATAGTTCCGTTATCGTT
<i>dgpC</i> center	AAGGTACGCATTTCTTCGATACTT
<i>dgpD</i> forward	GCTCCCATTTTGATCAATTCCTAT
<i>dgpD</i> reverse	ATACTTGCAGGAATATTTGCTTGG
<i>dgpD</i> center	TATCATTTACCAAACAAGCGATTC
<i>dgpF</i> forward	CAGCAAAGTTCGTGTGTATTTCATT
<i>dgpF</i> reverse	GTCAGCAGTCATGTTTATCTTTGC
<i>dgpF</i> center	AGGTACGCGTTTCTTCGATACTT
<i>dgpG</i> forward	TTGTAACCCGTTTAGAACCCAGTC
<i>dgpG</i> reverse	GTACCATACAAACCGTTGATCTGA
<i>dgpG</i> center	ACCTCTCTTTTCCTCCCTTAAAAA
<i>dgpH</i> forward	TCTCAAGTGAGTTGGTCTGTTGAT
<i>dgpH</i> reverse	AACAAATTTCCAACCAGTTCATTC
<i>dgpH</i> center	CACCCCTTATCCTCTTTCTTATCA
<i>dgpI</i> forward	ACTTATCGGTGCATGCTCTGTG
<i>dgpI</i> reverse	GGAAACCTCAGTCTCACCTAACTC
<i>dgpI</i> center	TCATCCCTTATTCGCTTTCTTATC

Table 6. Primers used in this study (cont.)

Primers used to clone <i>ssrA</i>, <i>ssrB</i>, <i>ssrC</i>, and <i>ssrD</i>	
<i>ssrA</i> forward	TGACGGATCCACTATCAATCAATCCGTTTCCAGT
<i>ssrA</i> reverse	CAAAGGATCCAAACGAGCGATAAGTCTGCTAAAC
<i>ssrB</i> forward	TTTGGGATCCTAAGCCAACCGAATTAAAGTGTTT
<i>ssrB</i> reverse	TTCGGGATCCACTTAGCAGAGGTGAAAGGAGCTT
<i>ssrC</i> forward	TCTAGGATCCAAAGCGTTAAACATCCCATAATCT
<i>ssrC</i> reverse	AGTTGGATCCTCATGATAGCTACCGGAGTATTGA
<i>ssrD</i> forward	GTAAGGATCCTCTTGATAGATCCGGAAATAATG
<i>ssrD</i> reverse	TTTAGGATCCAATTTTGCGCATTTACTCGTATTT
Primers used to clone the promoter, IRs, and flanking regions	
<i>dgpB</i> forward	CATTCTGCAGCGATTCTCAAGGGCTTTAATTTAC
<i>dgpB</i> reverse	TCAACTGCAGCACCAGCCAAAAGAGTAGAAAAC
<i>dgpC</i> forward	AAGACTGCAGAGTCGATTCTCTGGGAGTTAAGAA
<i>dgpC</i> reverse	TCAACTGCAGCACCTGCCAAAAGAGTAGAAAAC
<i>dgpF</i> forward	AGTTCTGCAGAAGGAATATTTGTTGATTGCTTGA
<i>dgpF</i> reverse	TTCCCTGCAGGTAACCTTTGCCATTTTCAACCTT
Primers used to demonstrate inversion of promoters by SsrA-SsrD	
<i>dgpB</i> C1	AGGTACGCGTTTCTTCGATACTT
<i>dgpB</i> C2	AATCGCTTGTTTCGTAAAGGATAA
<i>dgpC</i> C1	AAGGTACGCATTTCTTCGATACTT
<i>dgpC</i> C2	GTCTGGAAGGGTTGCAAGTATC
<i>dgpF</i> C1	AGGTACGCGTTTCTTCGATACTT
<i>dgpF</i> C2	GTTTGTATGGAATCGGTTGTTT
C7	GGGGACATTGTCTCTCTTC