

Table S1. Oligodeoxyribonucleotides used in this study

Name	Sequence (5'- 3')	Used for
ML-296	ATATAGTTTGGTTATTCTGTTGAATTTATGATGGATGG GGGACATAAAAAACAAAATAGAT	SDM ^a
ML-297	ATCTATTTTGTTTTTATGTCCCCATCCATCATAAATTC AACAGAATAACCAAATATAT	SDM
ML-394	CCTTGACAGAAATTTATTATTAAGCATTATTAGAATAC TGGAACCAATTTATTCATTTCA	SDM
ML-395	TGAAATGAATAAATTGGTTCAGTATTCTAATAATGCT TAATAATAAATTTCTGTCAAGG	SDM
ML-397	CCTTGACAGAAATTTATTATTAAGCATTATTAGAATAA TCGAACCAATTTATTCATTTCA	SDM
ML-398	TGAAATGAATAAATTGGTTCGATTATTCTAATAATGCT TAATAATAAATTTCTGTCAAGG	SDM
ML-331	AAGGGCAAATTGCTTTTATAGAGGCGAAATATACAAA TAAACGTCTC	SDM
ML-332	GAGACGTTTATTGTATATTTTCGCTCTATAAAAGCAA TTTGCCCTT	SDM ^a
ML-1344	TGAATAAATTGGTTCATTATTCTAATAACGCTTAATAATAA ATTTCTGTCAAGGCTTA	SDM ^a
ML-1345	TAAGCCTTGACAGAAATTTATTATTAAGCGTTATTAGAATAA TGGAACCAATTTATTTCA	SDM
virBQF	CCAAGTTCTCGGATGCTATGC	qRT-PCR
virBQR	CTCTTGATGCCAGAAAAGTAGCAA	qRT-PCR
icsAQF	TGATGGACTTTCTCCCTTGGG	qRT-PCR
icsAQR	TACCACGCATCCATTCCATCT	qRT-PCR
groELQF	TCTGTGGCTGGCCTGATGA	qRT-PCR
groELQR	TTTTTCGGCAGGTCGGTAAC	qRT-PCR
htpGQF	CAACCGCGTTGAAGGTAAGC	qRT-PCR
htpGQR	GACGGGATGTACAGCAGGCT	qRT-PCR
lacZQF	CCCATCTACACCAACGTGACC	qRT-PCR
lacZQR	AACAAACGGCGGATGACC	qRT-PCR
mdhQF	CCTGGTACAGCAAGTTGCG	qRT-PCR
mdhQR	CGTAGGCACATTCGACAACG	qRT-PCR
ML-512	TCAACAATCTTCTTATCTGATC	Primer Extension
ML-1314	TTAACCCCTTTTCTCTC	Primer Extension
ML-U25	GTTTCATTGCCTGAGCTA	Toeprinting
ML-U26	CAAATCGATAGAAACCTC	Toeprinting
GN20	TACGGATCCATCTTAATAACGGAAAG	Northern
FsinR	GGGATCCCCATCTGGC	Northern
ML-333	NNNGGATCCATGGGACATAAAAAACAAAATAG	<i>virF</i> ₃₀ pAC-30 cloning
ML-335	NNNGGATCCATGGAACCAATTTATTCATTTTC	<i>virF</i> ₂₁ pAC-21 cloning
ML-336	NNNGGATCCGAAAACCCATCTGGCAA	<i>virF</i> ₃₀ pAC30 and <i>virF</i> ₂₁ pAC21 cloning
ML-438	NNNGGATCCTTAATACGACTCACTATAGACAGAAATT TCTTAGTACTCT	T7- <i>virF</i> ₃₀ pAC-30- FT cloning
ML-566	TTACTATTTATCGTCGTCATCTTTGTAGTCAAATTTTT ATGATATAAGTAAAATTTCA	T7- <i>virF</i> ₃₀ pAC-30- FT cloning
ML-U6 ^b	<u>TCGACTTAATACGACTCACTATAGGGTCCATTACTGATGAGT</u> <u>CCGTGAGGACGAAACGGTACTCGGTACCGTCCG</u>	T7-HH pAC-T7-HH cloning
ML-U13 ^b	<u>GATCCGACGGTACCGAGTACCGTTTCGTCCTCACGGACTCAT</u> <u>CAGTAATGGACCCTATAGTGAGTCGTATTAAG</u>	T7-HH pAC-T7-HH cloning
ML-U7	NNNGGTACCGTCTAATGGAACCAATTTATTCATTTTC	<i>virF</i> +309-FT pAC- T7-HH-21-FT cloning

ML-U8	NNNGGATCCCAGTCATAGCCGAATAGCCT	<i>virF</i> +309-FT pAC-T7-HH-21-FT cloning
FustradFL	TTAGGATCCATATCCATCATAAATTC	<i>virF</i> pFL-4A fusion cloning
FustradFS	ATAGGATCCATAAATTCAACAGAATA	<i>virF</i> pFL-1A fusion cloning
ML-496	CTGGATCCTCCGCTTCCTTTAGCAGC	<i>virF-lacZ</i> fusion cloning
ML-330	NNNGGATCCATAGAAACCTCCTCCTCAGA	<i>virF-lacZ</i> fusion cloning
ML-334	NNNGGATCCATGACGGTTAGCTCAGGCAATG	Transcriptional <i>virF-lacZ</i> fusion cloning
ML-1342	NNNGGATCCGAATAATGGAACCAATTTATTC	Transcriptional <i>virF-lacZ</i> fusion cloning
ML-1343	NNNGATCCGAGCGAAATATACAAATAAACG	Transcriptional <i>virF-lacZ</i> fusion cloning
bx5	GGAGCTCTCACATCAG	<i>virB-lacZ</i> p _{virB} -lacZ fusion cloning
bx6	GTCGTTGCACAAATCC	<i>virB-lacZ</i> p _{virB} -lacZ fusion cloning
ML-U1	GAAATTAATACGACTCACTATAGACAGAAATTTCTTAGTTACT	<i>virF</i> R1 mRNA template for <i>in vitro</i> transcription
ML-U20	TAATACGACTCACTATAGGG	<i>virF</i> R2 mRNA template for <i>in vitro</i> transcription
ML-982	NNNGGATCCTGAGGGGATCTTGAAGTTCC	<i>virF</i> R1 and R2 mRNA template for <i>in vitro</i> transcription
QH7-	TAGGGATCCAAGCGAACCTTTATATC	<i>virF-lacZ</i> p <i>virF-lacZ</i> cloning
ML-U3	GAAATTAATACGACTCACTATAGGGAGATTATAAGCCTTGACAGA	VirF ₂₁ PCR template for <i>in vitro</i> translation
ML-U28	GGGATCCCCATCTGGC	VirF ₂₁ PCR template for <i>in vitro</i> translation
ML-U30	GGTTATAGTCCCTTTCAGTGCA	DNase I footprinting
ML-U29	CAAGCGAACCTTTATATC	DNase I footprinting

^aSDM: Site Directed Mutagenesis

^bThe underlined nucleotides specify the hammerhead sequence used for mRNA cleavage