

1 Supporting Information

2 Table S1. Primers for mutagenesis^a.

Mutation	Primer
l ₀	FW: GCTCTGGCTTGTGCTCTGGCTATGGACGCTCTGGCTCTGGCTGGCACAACCTATGGCATG RV: AGCCAGAGCCAGAGCGTCCATAGCCAGAGCACAAGCCAGAGCGCCTGATGTAACTTCAC
N154A	FW: TCGGAAAACCATGGGGCTTATTCAGCG RV: GCCCCATGGTTTTCCGAAGTGGTGGTT
BC	FW: ACGCTGCTGCTGCTGCTCCCTGCAAAATTCCGATT RV: GGAGCAGCAGCAGCAGCGTAGGAGAGTTCAATGAC
DE	FW: CGGCTGCTGCTGCCGCTTCAAAGGTGCTGGTCGAG RV: GAAGCGGCAGCAGCAGCCGCGACGAAGGGGTTCCAC
FG	FW: TTGCTGCTGCTGCTGCTCAGATCAACCACCATTGG RV: TGAGCAGCAGCAGCAGCAACTACGATGTAGGAGTC
kl	FW: TCATCATCAGAGTACTCAAGCTCATCAAAGTCAACATCAGGCCACCTGAAA RV: TGACTTTGATGAGCTTGAGTACTCTGATGATGAGGCTCCTGCCAACGCCTG
ij	FW: GGCGGCTGCCACAGCTCAGTCCGTTGTTGCTCTT RV: ACTGAGCTGTGGCAGCCGCCTCTTCAAACCTCCAT
R9A	FW: CTGGGAATGGCAATGCTGACTTCATA RV: GCATTGCCATTCCCAGACAATTAATA
E373A	FW: CAAAGGTGCTGGTCGCGATGGAACCC RV: GCGACCAGCACCTTTGAATTGGCACT
H144A	FW: GTTGGCATTGTTGTGGCTGGAACCACC RV: GCCACAAAAATGCCAACTTCGTATTTG
H319A	FW: CCGGCGGACACTGGTGCCGGAACAGTT RV: GCACCAGTGTCCGCCGATTTTTTCGCG
F407A	FW: ACGCTGGCAAAGCCGCTTCAACAACCT RV: GCGGCTTTGCCAGCGTGCTTCCAGCT
T410A	FW: AAAGCCTTTTCAACAGCTTTGAAGGG RV: CTGTTGAAAAGGCTTTGCCAGCGTG
L221S	FW: GAATGGTTTTCATGACTCCGCTCTCCCC RV: GAGTCATGAAACCATTCCCTATGGACC
W217A	FW: CTGGTCCATAGGGAAGCGTTTCATGAC RV: GCTTCCCTATGGACCAGAAATGACTTT
Q258A	FW: GTTGCTCTTGGGTCAGCGGAAGGAGGC RV: GCTGACCCAAGAGCAACAACGGACTGT

3 ^a Pairs of forward (FW) and reverse (RV) primers was used to generate mutations in

4 infectious clone, CprME and prME plasmids.