

Supplemental Material**A. VP8*-P[4], 159 amino acid (aa) residues**

LDGPYQPTTFKPPNDYWLLISSNTDGVVYESTNNSDFWTAVIAVEPHVSQTNRQYVLFGENKQFNVENSSDKWKF-FEMFKGSSQSDFSNRRTLTSNNRLVGMKYGGRVWTFHGETPRATTDSSNTADLNNISIIHSEFYIIPRSQESKCNEY-INNGL

B. VP8*-P[6], 159 aa residues

LDGPYQPTNFKPPNDYWILLNPTNQVVLLEGTNNTDIWVALLLVEPNVTNQSRQYTLFGEMKQITVENNTNKWKF-FEMFRSNVSAEFQHKRTLTSDTKLAGFMKFYNSVWTFHGETPHATTDYSSTSNLSEVQTVIHVEFYIIPRSQESKCSEY-INTGL

C. VP8*-P[8], 159 aa residues

LDGPYQPTTFTPPTDYWILINSNTNGVVYESTNNSDFWTAVIAVEPHVDPVDRQYNVFGENKQFNVRNDSKWK-FLEMRGSSQNDFYNRRTLTSDTRLVGIKYGGRVWTFHGETPRATTDSSNTANLNGISITIHSEFYIIPRSQESKCNEY-INNGL

D. VP5*, 232 aa residues

AQVNEDIIVSKTSLWKEMQYNRDIIRFKFGNSIVKMGGGLGYKWSEISYKAANYQNYLRDGEQVTAHTTCSVNGVN-NFSYNGGSLPTDFGISRYEVIKENSIVYVDYWDSDSKAFRNMVYVRSALANLNSVKCTGGSYDFSIPVGAWPVMNG-GAVSLHFAGVTLSTQFTDFVSLNSLRFRLTVDPEFSILRTRTVNLYGLPAANPNNGNEYEISGRFSLISLVPTNDD

Supplement 1. Consensus amino acid sequences of antigens VP8* (three genotypes) and VP5* that were employed for “reverse translation” *in silico* with subsequent cloning of synthetic genes into a vector for expression in *Escherichia coli*. (A) VP8*-P[4], (B) VP8*-P[6], (C) VP8*-P[8], (D) VP5*.