

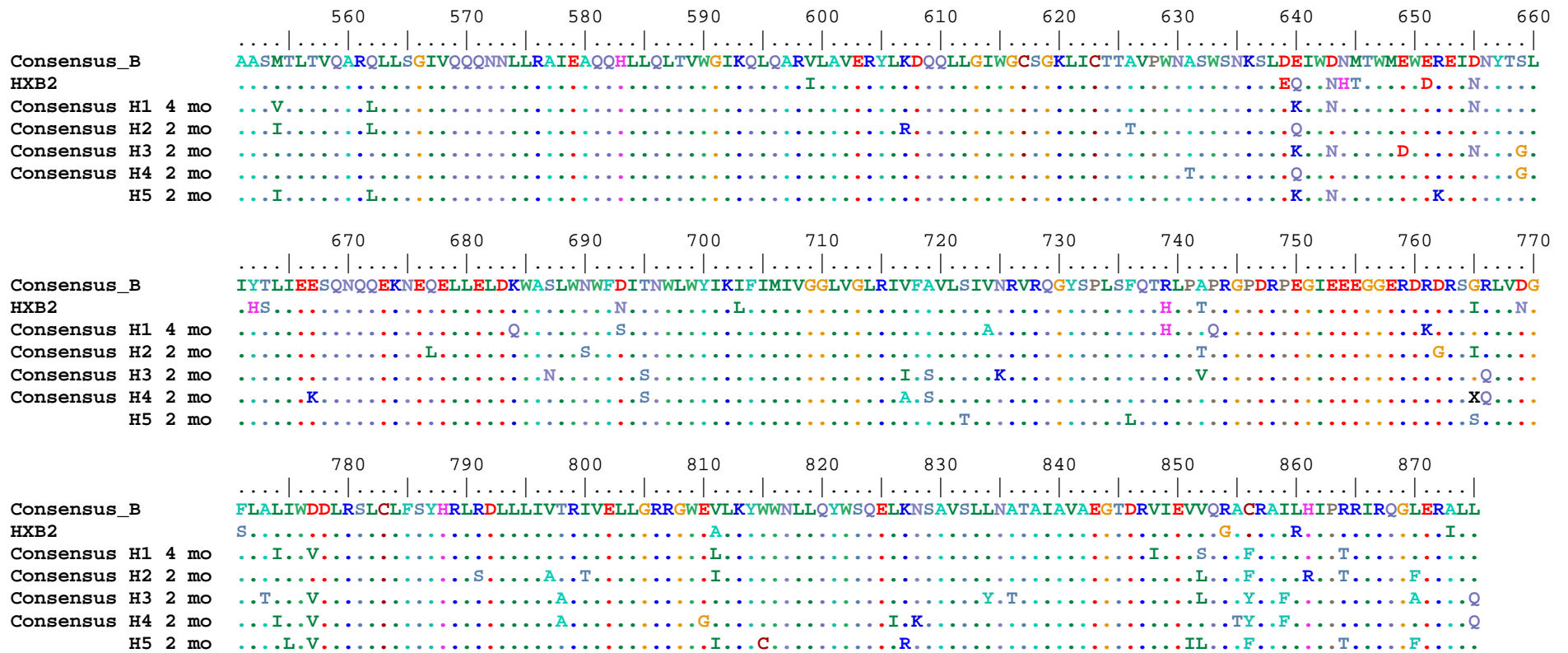
10 20 30 40 50 60 70 80 90 100 110
Consensus_B
HXB2
Consensus H1 4 mo
Consensus H2 2 mo
Consensus H3 2 mo
Consensus H4 2 mo
H5 2 mo

120 130 140 150 160 170 180 190 200 210 220
Consensus_B
HXB2
Consensus H1 4 mo
Consensus H2 2 mo
Consensus H3 2 mo
Consensus H4 2 mo
H5 2 mo

230 240 250 260 270 280 290 300 310 320 330
Consensus_B
HXB2
Consensus H1 4 mo
Consensus H2 2 mo
Consensus H3 2 mo
Consensus H4 2 mo
H5 2 mo

340 350 360 370 380 390 400 410 420 430 440
Consensus_B
HXB2
Consensus H1 4 mo
Consensus H2 2 mo
Consensus H3 2 mo
Consensus H4 2 mo
H5 2 mo

450 460 470 480 490 500 510 520 530 540 550
Consensus_B
HXB2
Consensus H1 4 mo
Consensus H2 2 mo
Consensus H3 2 mo
Consensus H4 2 mo
H5 2 mo



Supplementary Figure 1: Consensus gp160 sequences of the earliest virus variants from each patient aligned to the consensus B and HXB2 Env sequences. Since isolation of clonal virus variants was not successful for the earliest time point in individual H5, a single *env* sequence was generated from proviral DNA in patient PBMC from this time point. In those cases where no single residue was present in at least 50 % of the sequences from one patient, an X is shown.