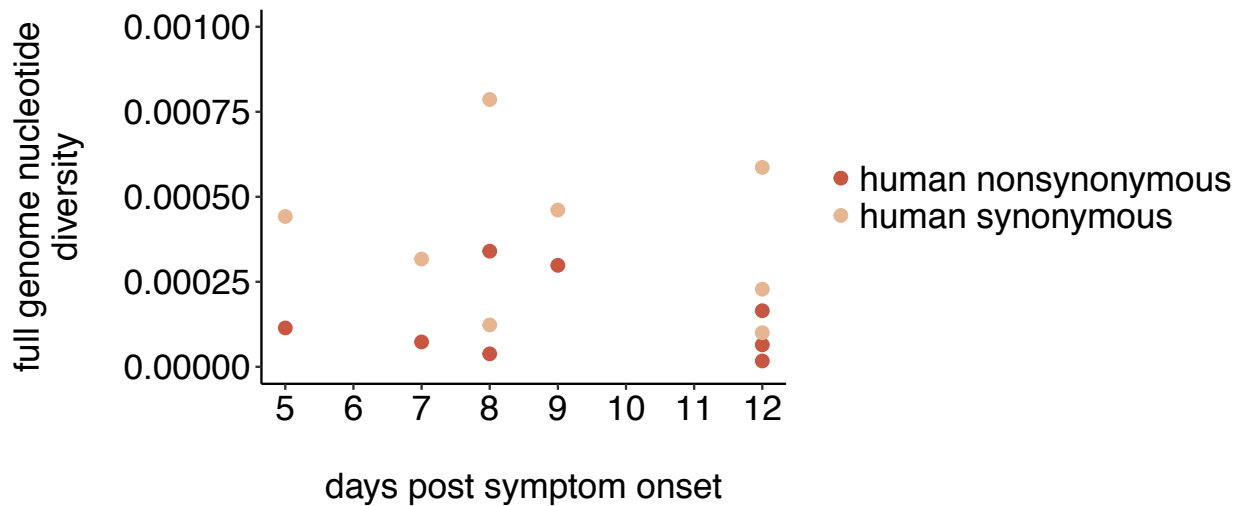


a**b**

| | “early infection” (5-8 days) | “late infection” (9-12 days) |
|-----------------------------|------------------------------|------------------------------|
| host-specific mutations | 18 | 22 |
| non-host specific mutations | 82 | 84 |

Figure S4: Neither diversity nor host-specific mutations increase over time

(a) For each human sample, the full genome nucleotide diversity (π_N or π_S) is plotted vs. the days post-symptom onset. Dark red dots represent the mean, full-genome nonsynonymous diversity for a given sample (π_N), and light red dots represent the mean, full-genome synonymous diversity for that same sample (π_S). Neither nonsynonymous nor synonymous diversity are correlated with days post symptom onset (nonsynonymous: $r^2 = -0.17$, $p = 0.69$; synonymous: $r^2 = -0.22$, $p = -0.61$). **(b)** To compare whether the number of putative host-adapting mutations increased over time in humans, we compared the number of host-specific and non-host specific mutations in humans sampled either in “early infection” (5-8 days post symptom onset), or in “late infection” (9-12 days post symptom onset). We divided the data into these categories by splitting on the mean days post symptom onset for human samples, which was 8 days. We then compared the proportion of host-specific variants during early and late infections with a Fisher’s exact test. The proportion of variants that are host-specific is not different in early vs. late infections ($p = 0.72$).