# COVID-19 subject HUP Q-0083

2021-05-05

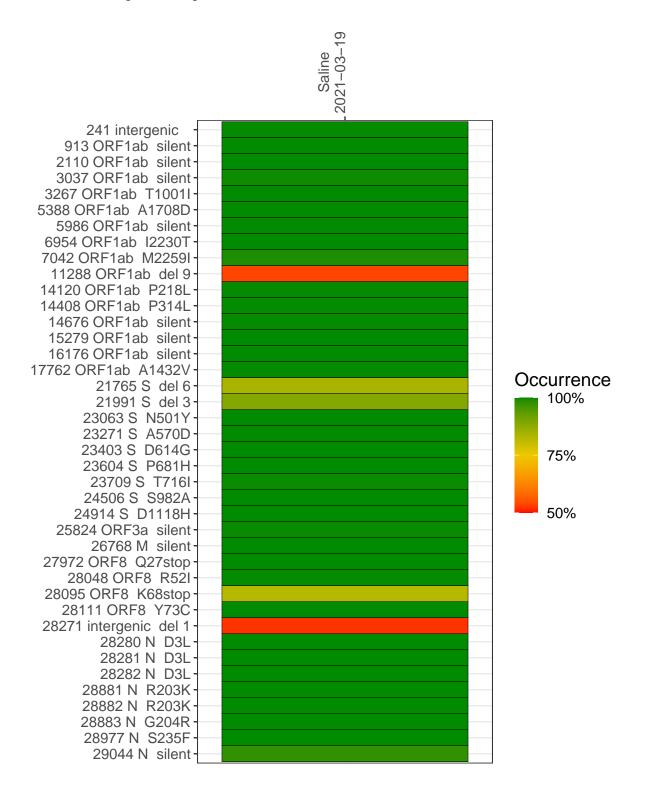
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. Lineages are called with the Pangolin software tool (Rambaut et al 2020) for genomes with > 90% sequence coverage.

Table 1. Sample summary.

Experiment	Туре	Genomes	Sample type	Sample date	Largest contig (KD)	Lineage	Reference read coverage	Reference read coverage (>= 5 reads)
VSP1250-1	single experiment	NA	Saline	2021-03-19	29.86	B.1.1.7	99.7%	99.7%

#### Variants shared across samples

The heat map below shows how variants (reference genome /home/everett/projects/SARS-CoV-2-Philadelphia/Wuhan-Hu-1) are shared across subject samples where the percent variance is colored. Variants are called if a variant position is covered by 5 or more reads, the alternative base is found in > 50% of read pairs and the variant yields a PHRED score > 20. Gray tiles denote positions where the variant was not the major variant or no variants were found. The relative base compositions of each experiment used to calculate tiles are shown in the following plot where the total number of position reads are shown atop of each plot.



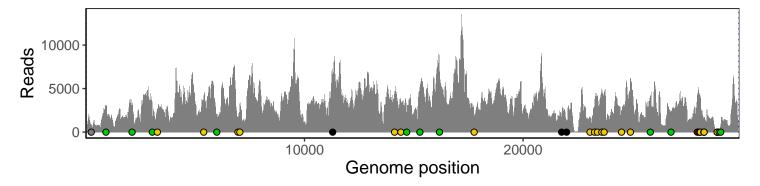
#### Saline 2021-03-19

	2021-03-19
241 intergenic	707
913 ORF1ab silent	2966
2110 ORF1ab silent	2849
3037 ORF1ab silent	2177
3267 ORF1ab T1001I	2732
5388 ORF1ab A1708D	3632
5986 ORF1ab silent	2070
6954 ORF1ab I2230T	1674
7042 ORF1ab M2259I	2521
11288 ORF1ab del 9	3255
14120 ORF1ab P218L	4205
14408 ORF1ab P314L	3287
14676 ORF1ab silent	1749
15279 ORF1ab silent	4556
16176 ORF1ab silent	6861
17762 ORF1ab A1432V	955
21765 S del 6	1673
21991 S del 3	1073
23063 S N501Y	1771
23271 S A570D	2811
23403 S D614G	4070
23604 S P681H	4283
23709 S T716I	3974
24506 S S982A	2460
24914 S D1118H	6005
25824 ORF3a silent	3925
26768 M silent	2246
27972 ORF8 Q27stop	4050
28048 ORF8 R52I	4394
28095 ORF8 K68stop	4094
28111 ORF8 Y73C	3485
28271 intergenic del 1	1416
28280 N D3L	716
28281 N D3L	716
28282 N D3L	782
28881 N R203K	20
28882 N R203K	20
28883 N G204R	20
28977 N S235F	50
29044 N silent	692
	7
	220
	7
	VSP1250-1
	/

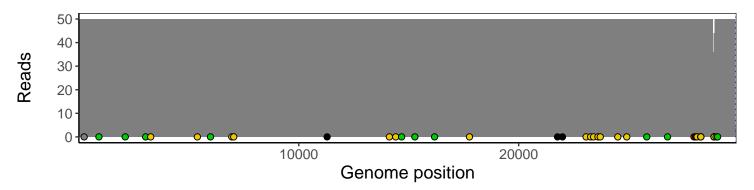
### Analyses of individual experiments and composite results

#### VSP1250-1 | 2021-03-19 | Saline | HUP Q-0083 | genomes | single experiment

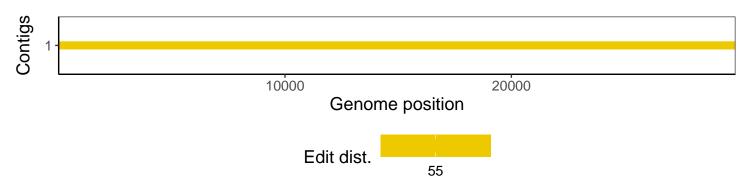
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



The longest five assembled contigs are shown below colored by their edit distance to the reference genome.



## Software environment

Software/R package	Version
R	3.4.0
bwa	0.7.17-r1198-dirty
samtools	1.10 Using htslib 1.10
bcftools	1.10.2-34-g1a12af0-dirty Using htslib 1.10.2-57-gf58a6f3
pangolin	2.3.8
genbankr	1.4.0
optparse	1.6.0
forcats	0.3.0
stringr	1.4.0
dplyr	0.8.1
purrr	0.2.5
readr	1.1.1
tidyr	0.8.1
tibble	2.1.2
ggplot2	3.0.0
tidyverse	1.2.1
ShortRead	1.34.2
${\it Genomic Alignments}$	1.12.2
SummarizedExperiment	1.6.5
DelayedArray	0.2.7
matrixStats	0.54.0
Biobase	2.36.2
Rsamtools	1.28.0
GenomicRanges	1.28.6
$\operatorname{GenomeInfoDb}$	1.12.3
Biostrings	2.44.2
XVector	0.16.0
IRanges	2.10.5
S4Vectors	0.14.7
BiocParallel	1.10.1
BiocGenerics	0.22.1