# COVID-19 subject UPHS-0046

2021-05-05

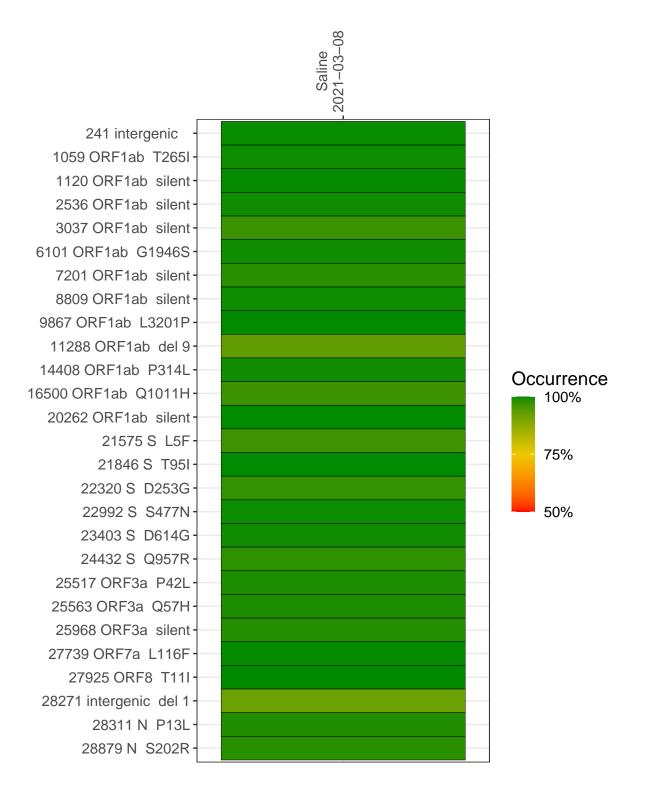
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. Lineages are called with the Pangolin software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

Table 1. Sample summary.

Experiment	Type	Genomes	Sample type	Sample date	Largest contig (KD)	Lineage	Reference read coverage	Reference read coverage (>= 5 reads)
VSP0978-1	single experiment	NA	Saline	2021-03-08	29.80	B.1.526.2	99.7%	99.6%

#### 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 2021-03-08

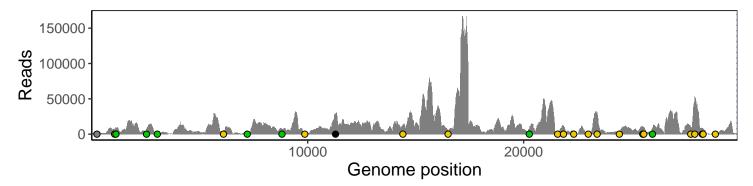
	2021 00 00
241 intergenic	2639
1059 ORF1ab T265I	6809
1120 ORF1ab silent	9641
2536 ORF1ab silent	7178
3037 ORF1ab silent	1924
6101 ORF1ab G1946S	5138
7201 ORF1ab silent	790
8809 ORF1ab silent	7071
9867 ORF1ab L3201P	506
11288 ORF1ab del 9	25383
14408 ORF1ab P314L	13108
16500 ORF1ab Q1011H	4599
20262 ORF1ab silent	7184
21575 S L5F	1588
21846 S T95I	11575
22320 S D253G	1170
22992 S S477N	352
23403 S D614G	30968
24432 S Q957R	3368
25517 ORF3a P42L	10817
25563 ORF3a Q57H	9862
25968 ORF3a silent	8467
27739 ORF7a L116F	10175
27925 ORF8 T11I	52208
28271 intergenic del 1	8294
28311 N P13L	7810
28879 N S202R	636
	0978-1
	760



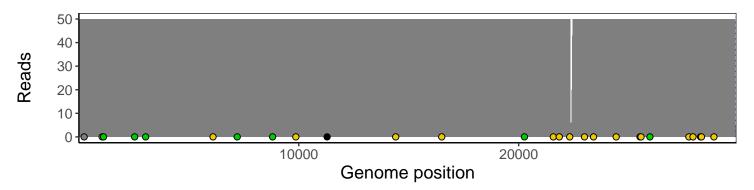
### Analyses of individual experiments and composite results

#### $VSP0978\text{-}1 \mid 2021\text{-}03\text{-}08 \mid Saline \mid UPHS\text{-}0046 \mid genomes \mid single \ experiment$

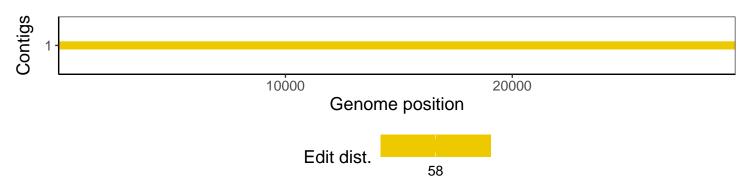
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



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