# COVID-19 subject UPHS-1647

2021-06-03

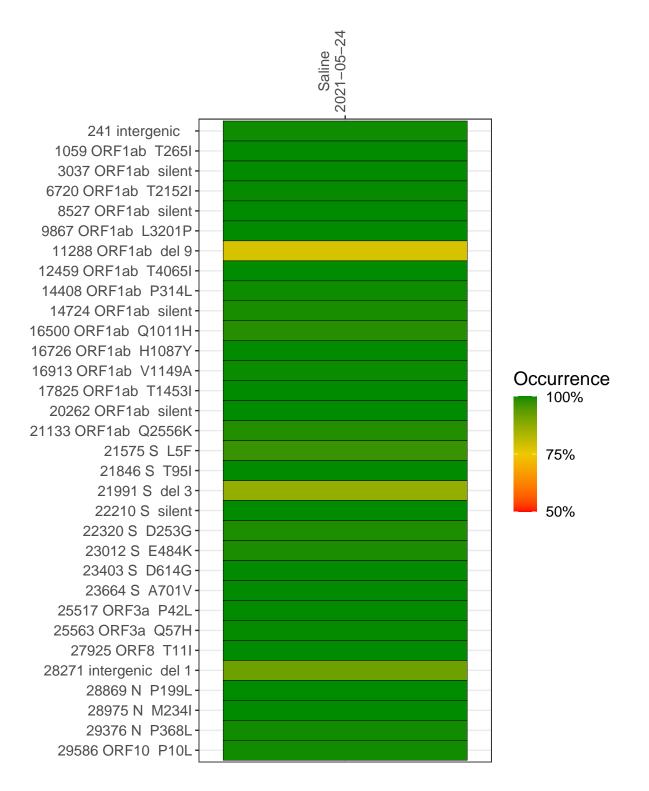
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. Lineages are called with the Pangolin software 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)
VSP2948-1	single experiment	NA	Saline	2021-05-24	29.87	B.1.526	99.7%	99.7%

#### Variants shared across samples

The heat map below shows how variants (reference genome /home/common/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 2021-05-24

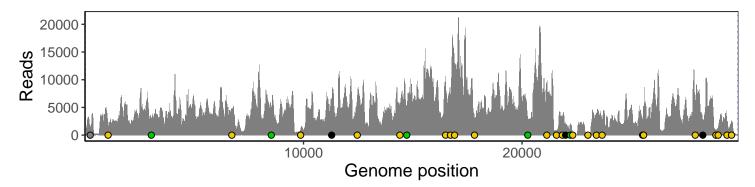
	2021-05-24
241 intergenic	1422
1059 ORF1ab T265I	1877
3037 ORF1ab silent	2801
6720 ORF1ab T2152I	4758
8527 ORF1ab silent	4828
9867 ORF1ab L3201P	901
11288 ORF1ab del 9	4283
12459 ORF1ab T4065I	7209
14408 ORF1ab P314L	4764
14724 ORF1ab silent	4160
16500 ORF1ab Q1011H	8885
16726 ORF1ab H1087Y	7459
16913 ORF1ab V1149A	11986
17825 ORF1ab T1453I	2896
20262 ORF1ab silent	3035
21133 ORF1ab Q2556K	8866
21575 S L5F	750
21846 S T95I	3043
21991 S del 3	1842
22210 S silent	1866
22320 S D253G	415
23012 S E484K	123
23403 S D614G	3906
23664 S A701V	2823
25517 ORF3a P42L	2662
25563 ORF3a Q57H	5157
27925 ORF8 T11I	6209
28271 intergenic del 1	2802
28869 N P199L	878
28975 N M234I	1564
29376 N P368L	2490
29586 ORF10 P10L	2500
	7-



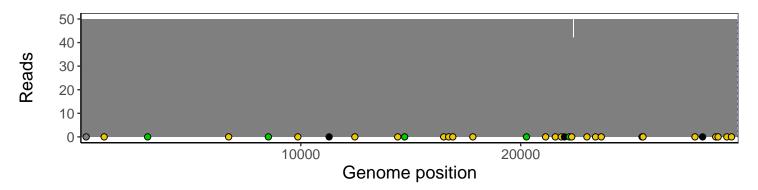
## Analyses of individual experiments and composite results

### $VSP2948-1 \mid 2021-05-24 \mid Saline \mid UPHS-1647 \mid genomes \mid single \; experiment$

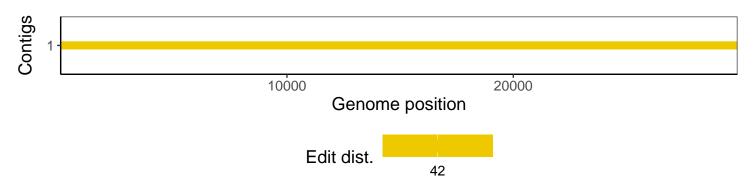
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



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	3.3.3
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