COVID-19 subject UPHS-0357

2021-06-23

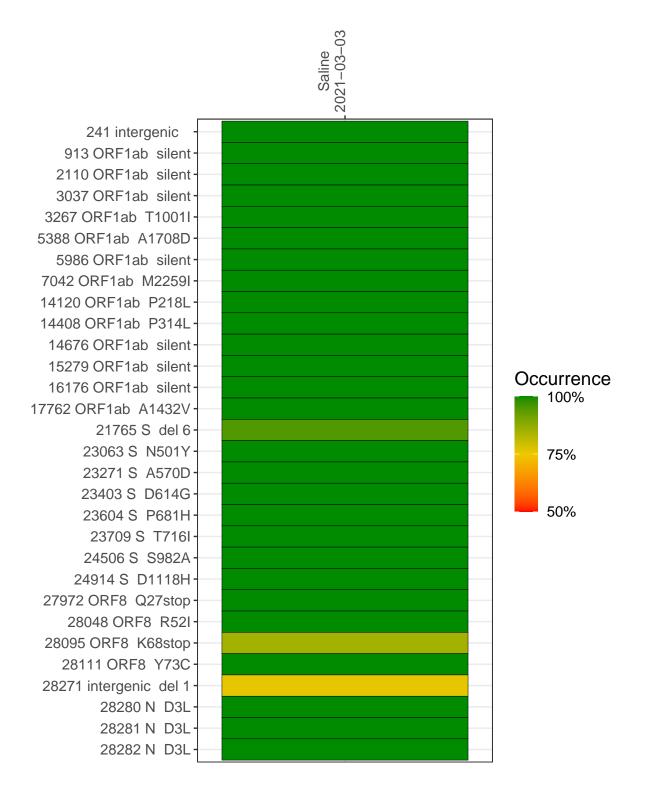
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. Lineages are called with the Pangolin software 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)
VSP1402-1	single experiment	NA	Saline	2021-03-03	21.68	B.1.1.7	99.6%	97.6%

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 2021-03-03

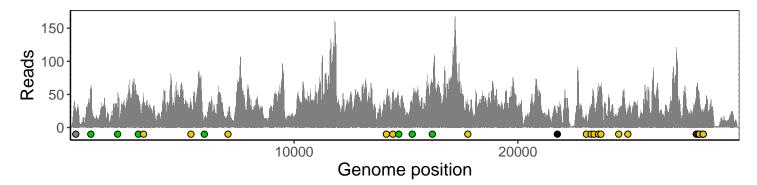
	2021-03-03
241 intergenic	27
913 ORF1ab silent	58
2110 ORF1ab silent	29
3037 ORF1ab silent	37
3267 ORF1ab T1001I	45
5388 ORF1ab A1708D	37
5986 ORF1ab silent	14
7042 ORF1ab M2259I	22
14120 ORF1ab P218L	38
14408 ORF1ab P314L	52
14676 ORF1ab silent	36
15279 ORF1ab silent	31
16176 ORF1ab silent	63
17762 ORF1ab A1432V	35
21765 S del 6	20
23063 S N501Y	17
23271 S A570D	29
23403 S D614G	44
23604 S P681H	52
23709 S T716I	60
24506 S S982A	30
24914 S D1118H	31
27972 ORF8 Q27stop	59
28048 ORF8 R52I	50
28095 ORF8 K68stop	53
28111 ORF8 Y73C	42
28271 intergenic del 1	21
28280 N D3L	16
28281 N D3L	16
28282 N D3L	17
	VSP1402-1
	Š/



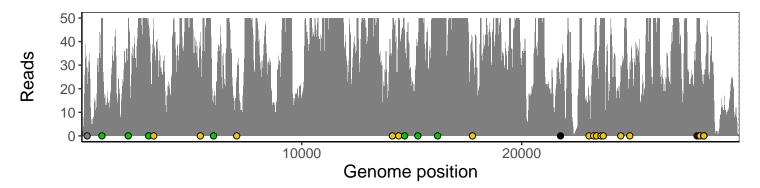
Analyses of individual experiments and composite results

$VSP1402\text{-}1 \mid 2021\text{-}03\text{-}03 \mid Saline \mid UPHS\text{-}0357 \mid genomes \mid single \ experiment$

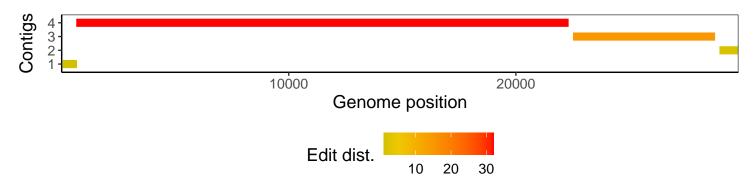
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



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	3.1.3				
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				
GenomicAlignments	1.12.2				
${\bf Summarized Experiment}$	1.6.5				
DelayedArray	0.2.7				
matrixStats	0.54.0				
Biobase	2.36.2				
Rsamtools	1.28.0				
GenomicRanges	1.28.6				
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				