

# COVID-19 subject UPHS-1366

*2021-05-21*

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 ( $\geq 5$ reads)
VSP2621-1	single experiment	NA	Saline	2021-04-27	29.81	B.1.526.2	99.8%	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 2021-04-27	
241 intergenic	2773	
1059 ORF1ab T265I	3390	
1387 ORF1ab silent	3373	
3037 ORF1ab silent	4776	
6101 ORF1ab G1946S	4043	
6470 ORF1ab T2069A	4640	
7201 ORF1ab silent	1350	
8809 ORF1ab silent	3168	
9430 ORF1ab silent	6426	
9867 ORF1ab L3201P	1277	
11288 ORF1ab del 9	4170	
14408 ORF1ab P314L	7847	
16500 ORF1ab Q1011H	4651	
18131 ORF1ab T1555I	8840	
20262 ORF1ab silent	1727	
21575 S L5F	1692	
21846 S T95I	5881	
22320 S D253G	465	
22992 S S477N	840	
23403 S D614G	7270	
24432 S Q957R	3074	
25517 ORF3a P42L	5231	
25563 ORF3a Q57H	6998	
25968 ORF3a silent	5906	
26885 M silent	3959	
27739 ORF7a L116F	2411	
27925 ORF8 T11I	6630	
28271 intergenic del 1	4710	
28311 N P13L	4803	
28879 N S202R	861	
	VSP2621-1	

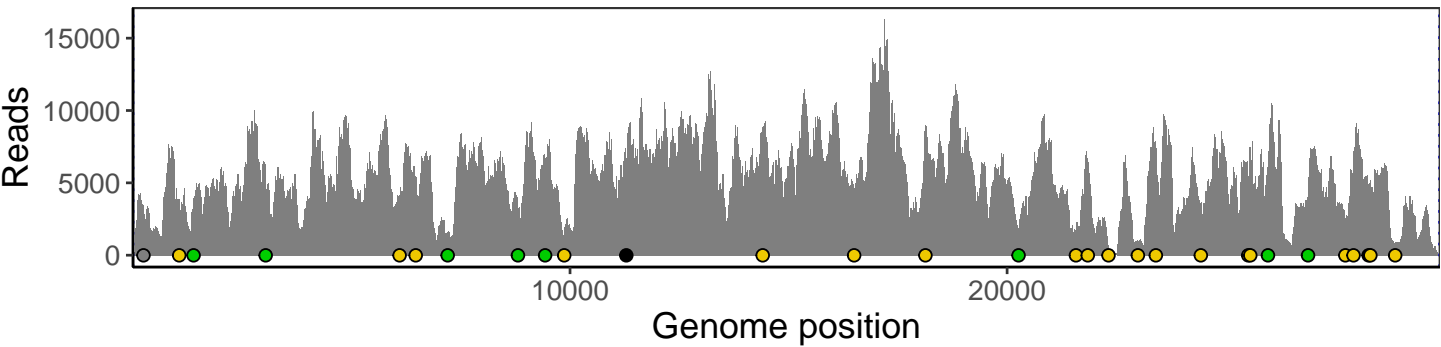
Base change

- Expected
- A
- T
- C
- G
- N
- Ins/Del
- No data

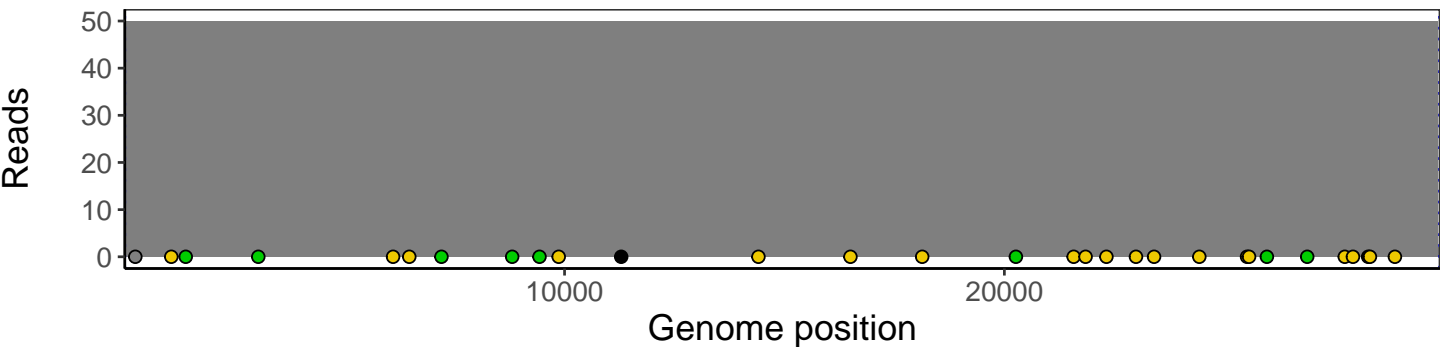
# Analyses of individual experiments and composite results

VSP2621-1 | 2021-04-27 | Saline | UPHS-1366 | genomes | single experiment

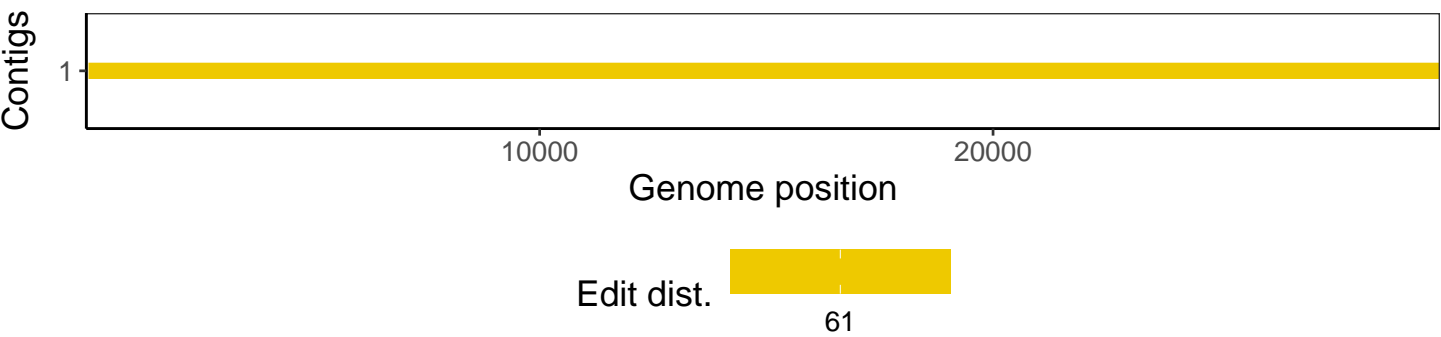
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according to variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



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 htlib 1.10
bcftools	1.10.2-34-g1a12af0-dirty Using htlib 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
GenomicAlignments	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
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