

COVID-19 subject UPHS-1359

2021-06-23

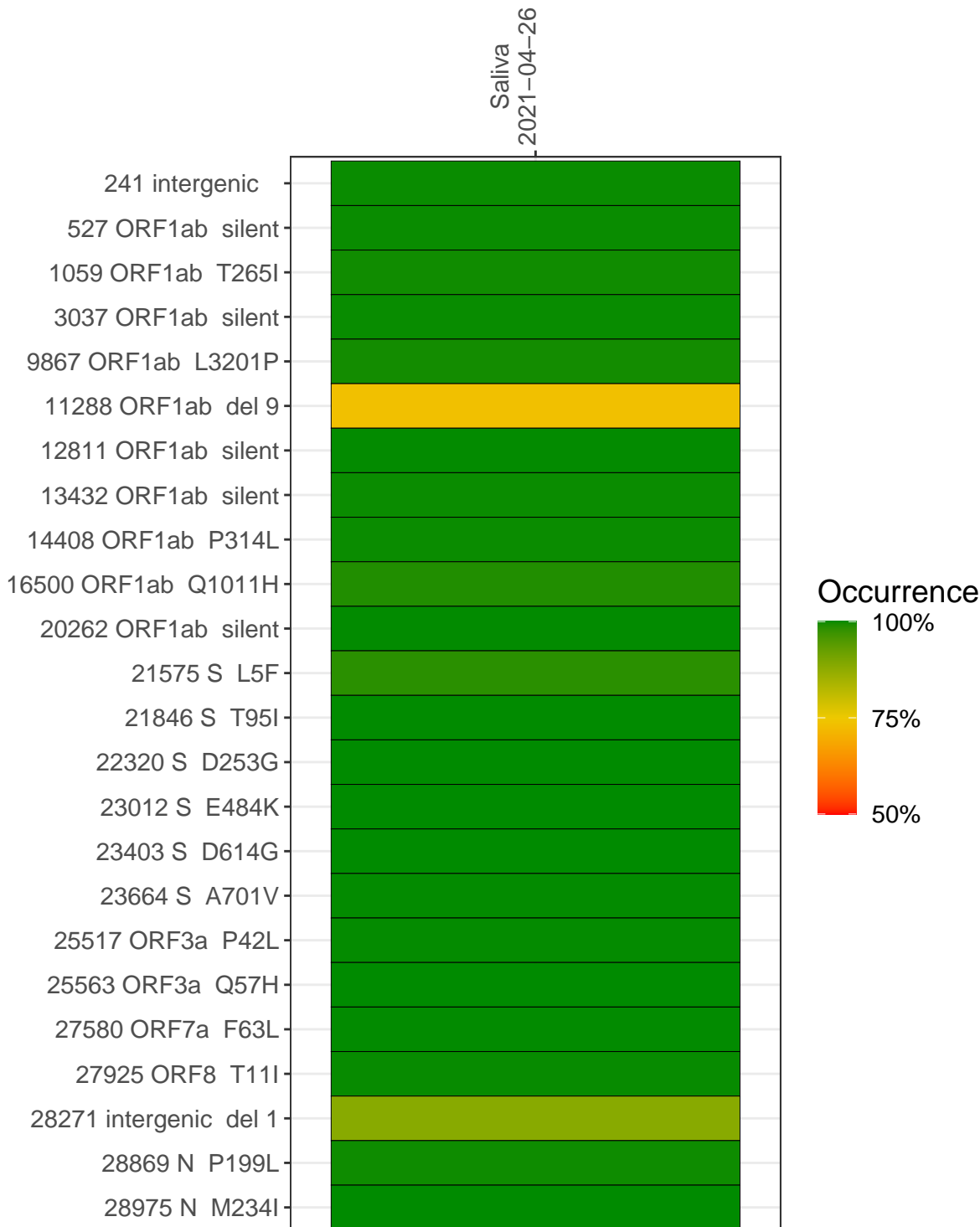
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. Lineages are called with the Pangolin software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

Table 1. Sample summary.

Experiment	Type	Genomes	Sample type	Sample date	Largest contig (KD)	Lineage	Reference read coverage	Reference read coverage (≥ 5 reads)
VSP2614-1	single experiment	NA	Saliva	2021-04-26	29.79	B.1.526	99.8%	99.8%

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.



Saliva
2021-04-26

241 intergenic	2783
527 ORF1ab silent	884
1059 ORF1ab T265I	2880
3037 ORF1ab silent	3429
9867 ORF1ab L3201P	963
11288 ORF1ab del 9	3713
12811 ORF1ab silent	4727
13432 ORF1ab silent	2917
14408 ORF1ab P314L	5145
16500 ORF1ab Q1011H	4170
20262 ORF1ab silent	2192
21575 S L5F	1855
21846 S T95I	5048
22320 S D253G	772
23012 S E484K	782
23403 S D614G	6263
23664 S A701V	5267
25517 ORF3a P42L	3722
25563 ORF3a Q57H	4818
27580 ORF7a F63L	4693
27925 ORF8 T11I	7516
28271 intergenic del 1	5689
28869 N P199L	1128
28975 N M234I	945

Base change

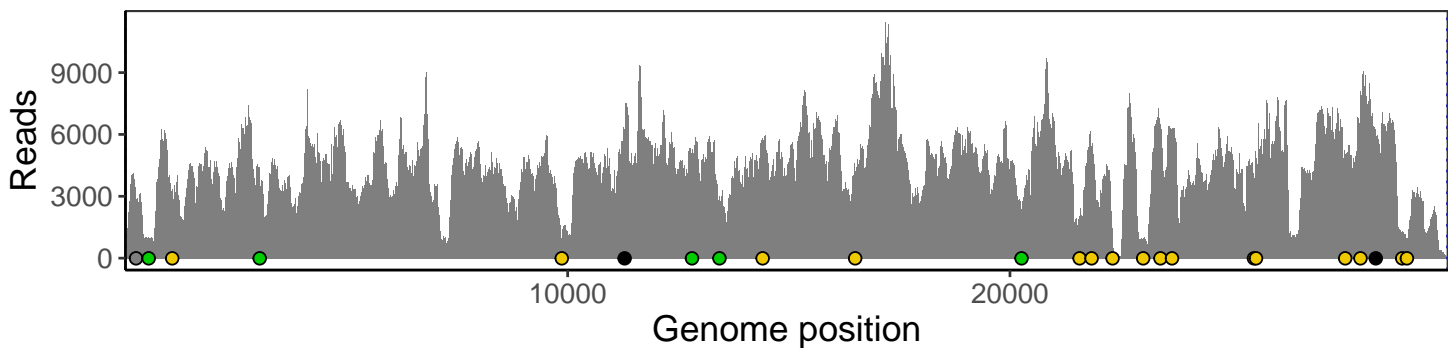
- Expected
- A
- T
- C
- G
- N
- Ins/Del
- No data

VSP2614-1

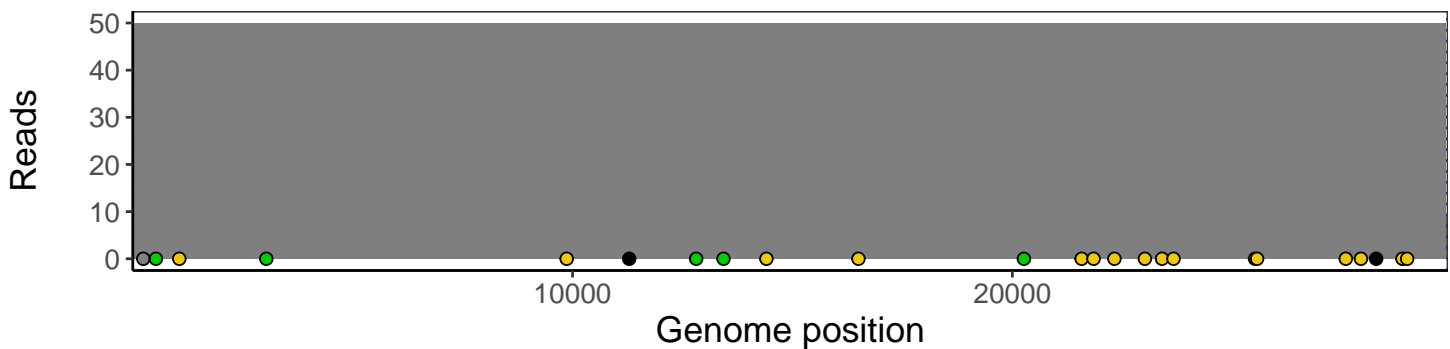
Analyses of individual experiments and composite results

VSP2614-1 | 2021-04-26 | Saliva | UPHS-1359 | genomes | single experiment

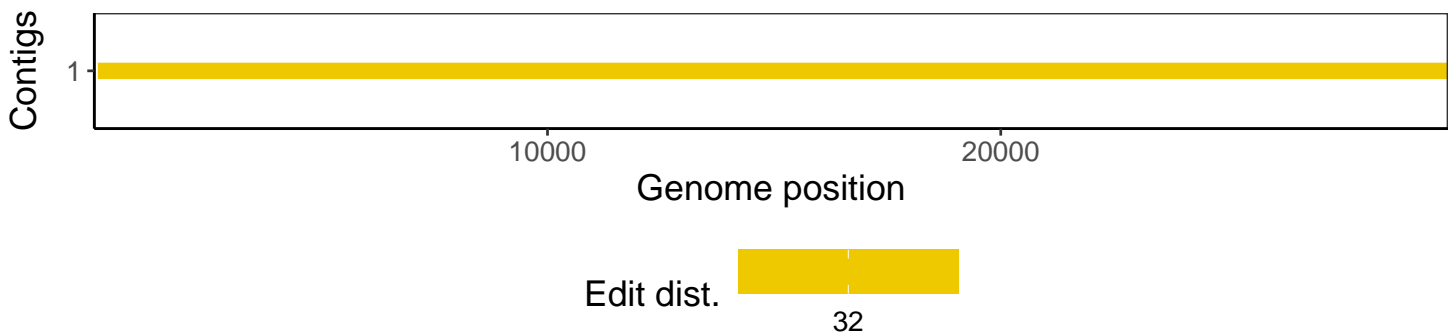
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according to variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



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	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
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