COVID-19 subject UPHS-0062

2021-05-05

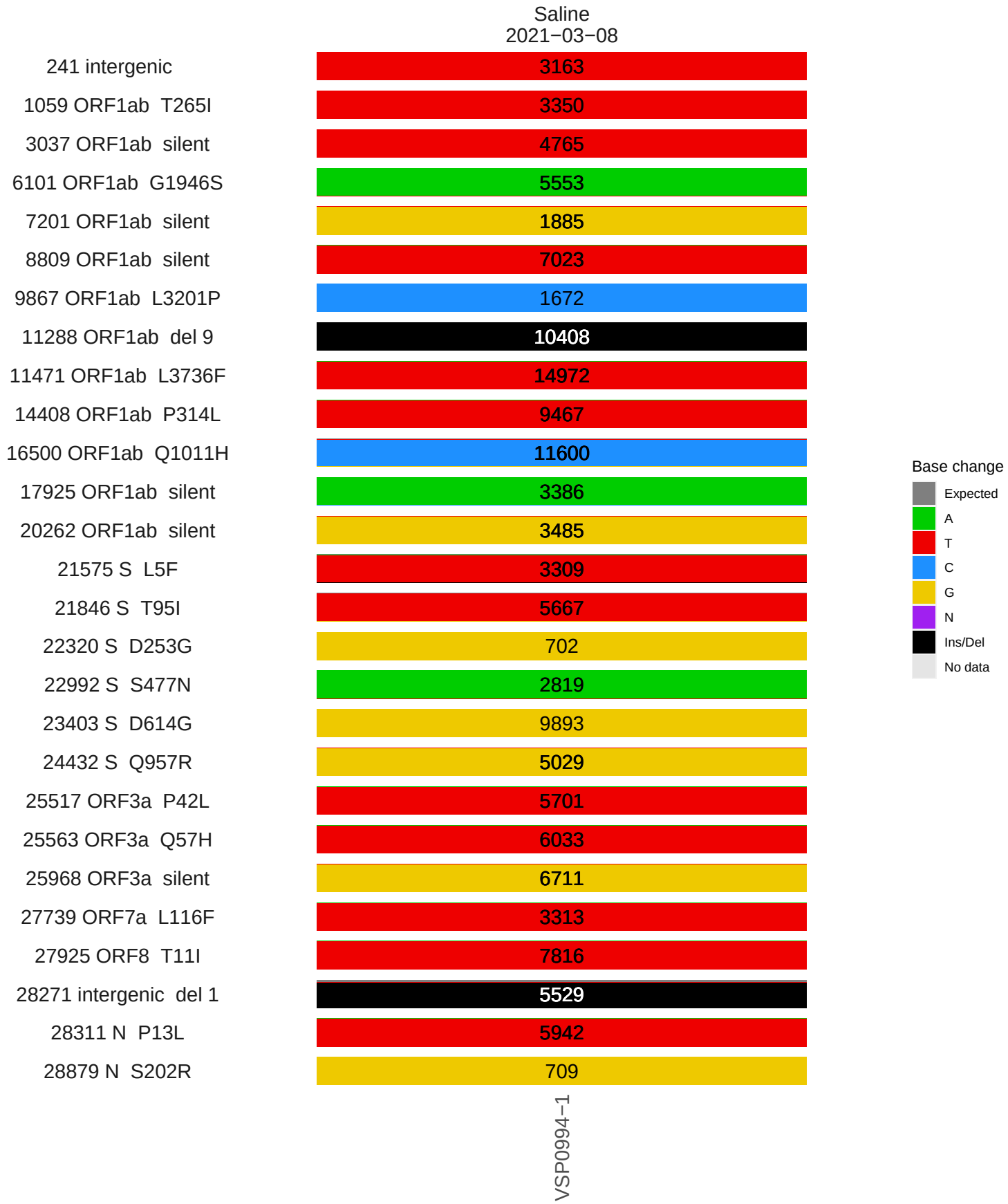
The table below provides a summary of subject samples for which sequencing data is available.
The experiments column shows the number of sequencing experiments performed for each specimen.
Experiment specific analyses are shown at the end of this report. Lineages are called with the Pangolin software 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)
VSP0994-1	single experiment	NA	Saline	2021-03-08	29.81	B.1.526.2	99.9%	99.8%

Variants shared across samples

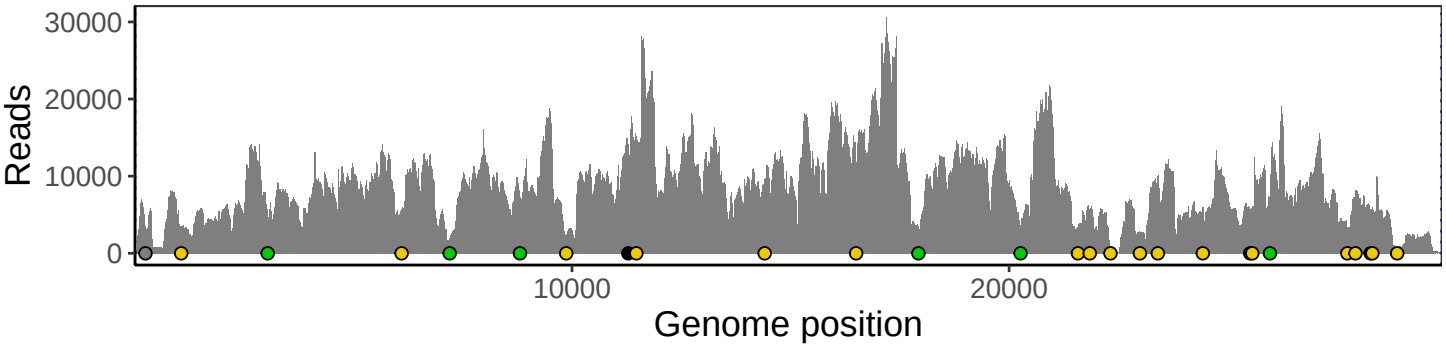
The heat map below shows how variants (reference genome /home/everett/projects/SARS-CoV-2-Philadelphia/Wuhan-Hu-1) are shared across subject samples where the percent variance is colored. Variants are called if a variant position is covered by 5 or more reads, the alternative base is found in > 50% of 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
241 intergenic
3163
1059 ORF1ab T265I
3350
3037 ORF1ab silent
4765
6101 ORF1ab G1946S
5553
7201 ORF1ab silent
1885
8809 ORF1ab silent
7023
9867 ORF1ab L3201P
1672
11288 ORF1ab del 9
10408
11471 ORF1ab L3736F
14972
14408 ORF1ab P314L
9467
16500 ORF1ab Q1011H
11600
17925 ORF1ab silent
3386
20262 ORF1ab silent
3485
21575 S L5F
3309
21846 S T95I
5667
22320 S D253G
702
22992 S S477N
2819
23403 S D614G
9893
24432 S Q957R
5029
25517 ORF3a P42L
5701
25563 ORF3a Q57H
6033
25968 ORF3a silent
6711
27739 ORF7a L116F
3313
27925 ORF8 T11I
7816
28271 intergenic del 1
5529
28311 N P13L
5942
28879 N S202R
709
VSP0994−1
Base change
Expected
A
T
C
G
N
Ins/Del
No data

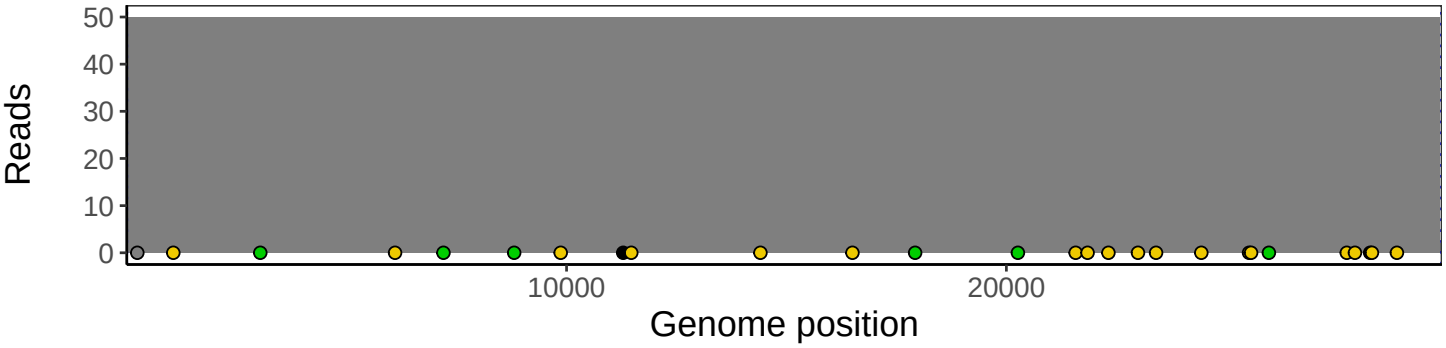
Analyses of individual experiments and composite results

VSP0994-1 | 2021-03-08 | Saline | UPHS-0062 | genomes | single experiment

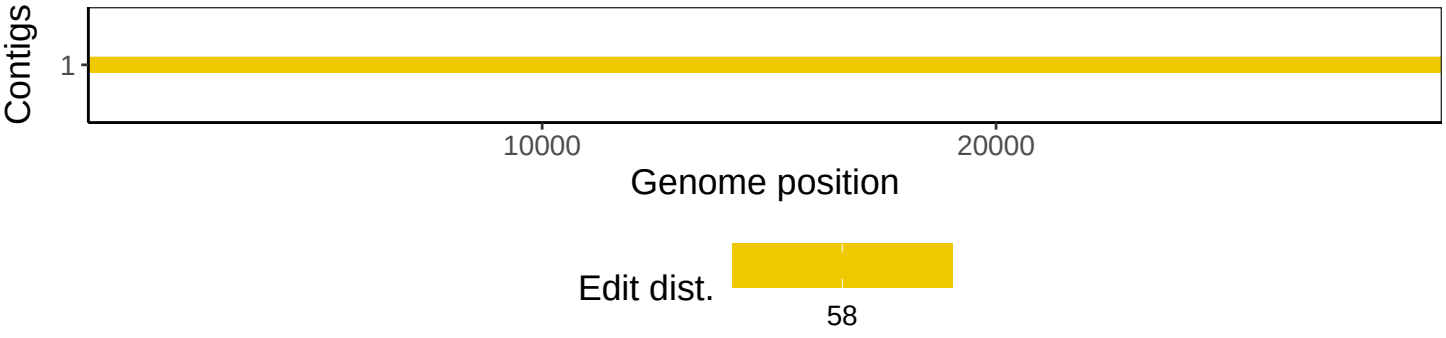
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



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
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