COVID-19 subject UPHS-0392

2021-06-23

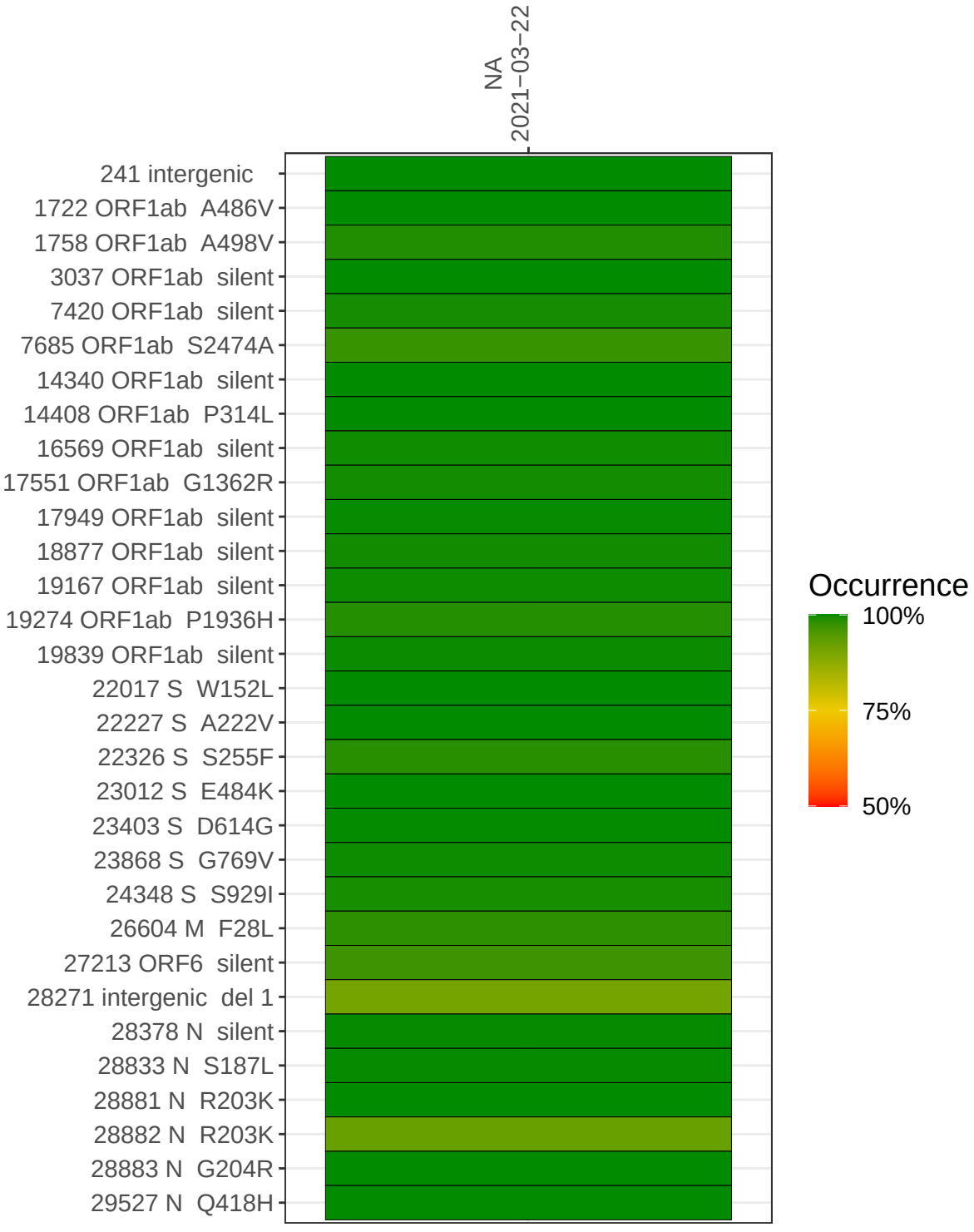
The table below provides a summary of subject samples for which sequencing data is available.
The experiments column shows the number of sequencing experiments performed for each specimen.
Experiment specific analyses are shown at the end of this report. Lineages are called with the Pangolin software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

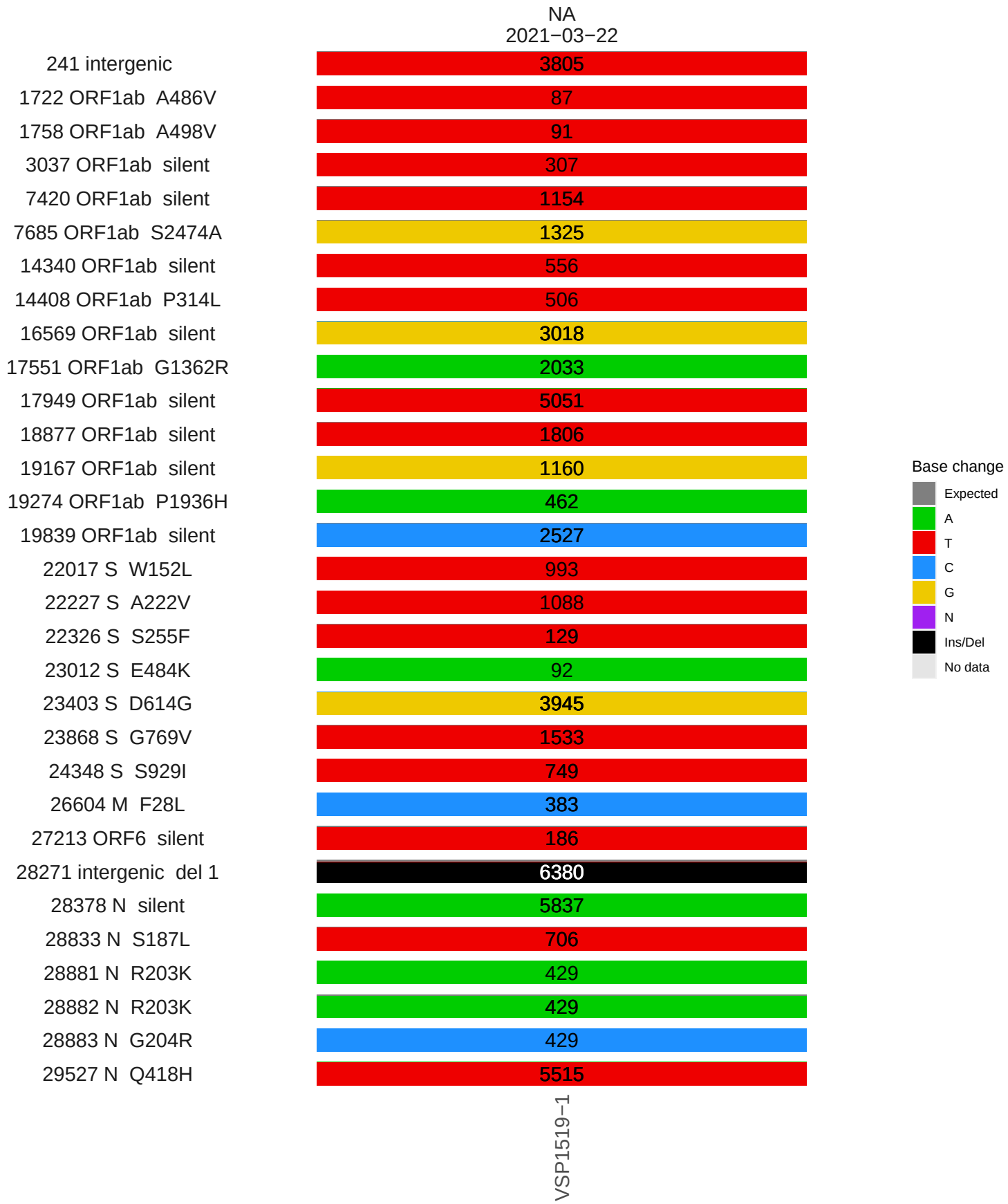
Table 1. Sample summary.

Experiment	Type	Genomes	Sample type	Sample date	Largest contig (KD)	Lineage	Reference read coverage	Reference read coverage (>= 5 reads)
VSP1519-1	single experiment	NA	NA	2021-03-22	21.74	R.1	99.2%	98.2%

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.

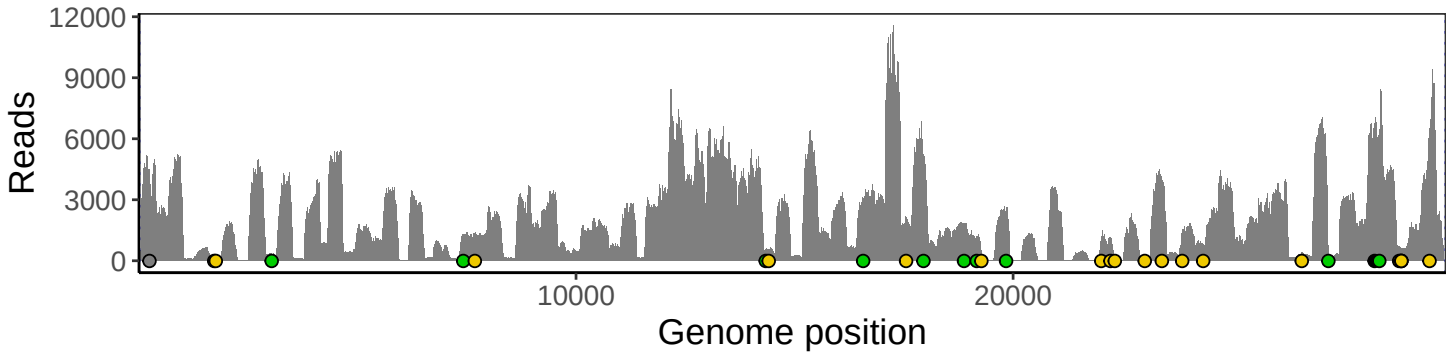



NA
2021−03−22
241 intergenic
3805
1722 ORF1ab A486V
87
1758 ORF1ab A498V
91
3037 ORF1ab silent
307
7420 ORF1ab silent
1154
7685 ORF1ab S2474A
1325
14340 ORF1ab silent
556
14408 ORF1ab P314L
506
16569 ORF1ab silent
3018
17551 ORF1ab G1362R
2033
17949 ORF1ab silent
5051
18877 ORF1ab silent
1806
19167 ORF1ab silent
1160
19274 ORF1ab P1936H
462
19839 ORF1ab silent
2527
22017 S W152L
993
22227 S A222V
1088
22326 S S255F
129
23012 S E484K
92
23403 S D614G
3945
23868 S G769V
1533
24348 S S929I
749
26604 M F28L
383
27213 ORF6 silent
186
28271 intergenic del 1
6380
28378 N silent
5837
28833 N S187L
706
28881 N R203K
429
28882 N R203K
429
28883 N G204R
429
29527 N Q418H
5515
VSP1519−1
Base change
Expected
A
T
C
G
N
Ins/Del
No data

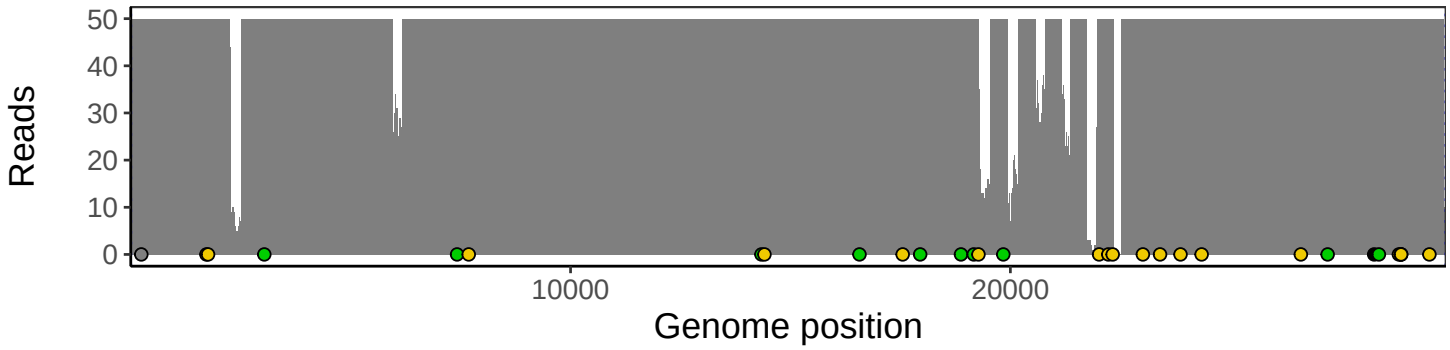
Analyses of individual experiments and composite results

VSP1519-1 | 2021-03-22 | NA | UPHS-0392 | genomes | single experiment

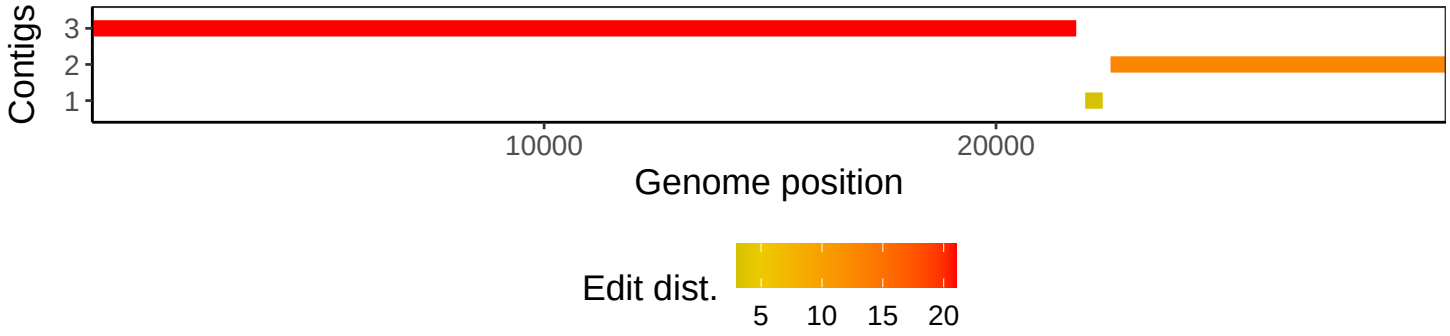
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



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	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