# COVID-19 subject HUP-Q-0004

2021-03-01

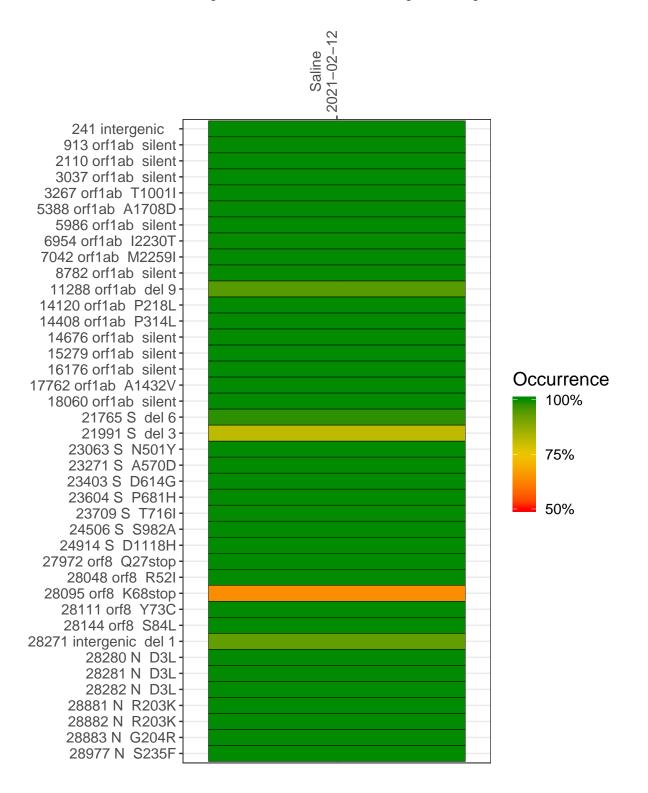
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. Lineages are called with the Pangolin software 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)
VSP0815-1	single experiment	NA	Saline	2021-02-12	29.80	B.1.1.7	99.8%	99.7%

#### Variants shared across samples

The heat map below shows how variants (reference genome USA-WA1-2020) 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.



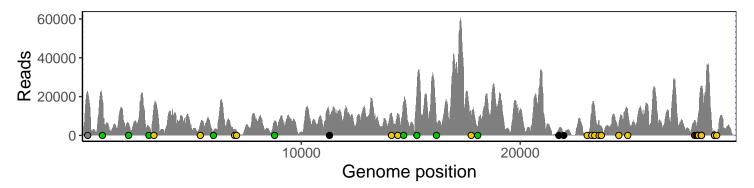
#### Saline 2021-02-12

	2021-02-12
241 intergenic	21873
913 orf1ab silent	18984
2110 orf1ab silent	6192
3037 orf1ab silent	4589
3267 orf1ab T1001I	15389
5388 orf1ab A1708D	6058
5986 orf1ab silent	2229
6954 orf1ab I2230T	3240
7042 orf1ab M2259I	2517
8782 orf1ab silent	7432
11288 orf1ab del 9	10165
14120 orf1ab P218L	11488
14408 orf1ab P314L	11046
14676 orf1ab silent	17260
15279 orf1ab silent	23532
16176 orf1ab silent	6934
17762 orf1ab A1432V	10080
18060 orf1ab silent	4259
21765 S del 6	2931
21991 S del 3	1016
23063 S N501Y	859
23271 S A570D	12796
23403 S D614G	14863
23604 S P681H	7836
23709 S T716I	4472
24506 S S982A	6518
24914 S D1118H	13490
27972 orf8 Q27stop	7686
28048 orf8 R52I	5447
28095 orf8 K68stop	5875
28111 orf8 Y73C	8466
28144 orf8 S84L	11292
28271 intergenic del 1	25088
28280 N D3L	23324
28281 N D3L	23324
28282 N D3L	23391
28881 N R203K	2330
28882 N R203K	2330
28883 N G204R	2330
28977 N S235F	778
	5–1
	5

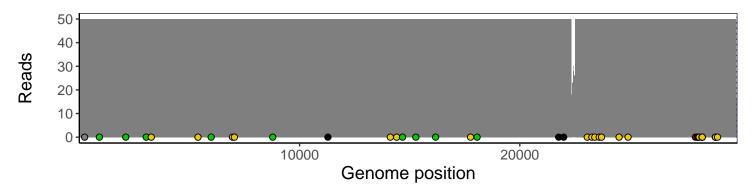
#### Analyses of individual experiments and composite results

### $VSP0815\text{-}1 \mid 2021\text{-}02\text{-}12 \mid Saline \mid HUP\text{-}Q\text{-}0004 \mid genomes \mid single \ experiment}$

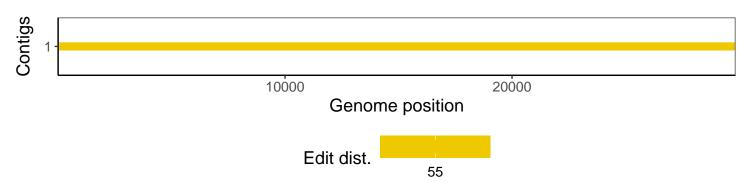
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



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	2.3.3
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	3.0.0
tidyverse	1.2.1
ShortRead	1.34.2
$\operatorname{GenomicAlignments}$	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
$\operatorname{GenomeInfoDb}$	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