COVID-19 subject 2753

2021-01-08

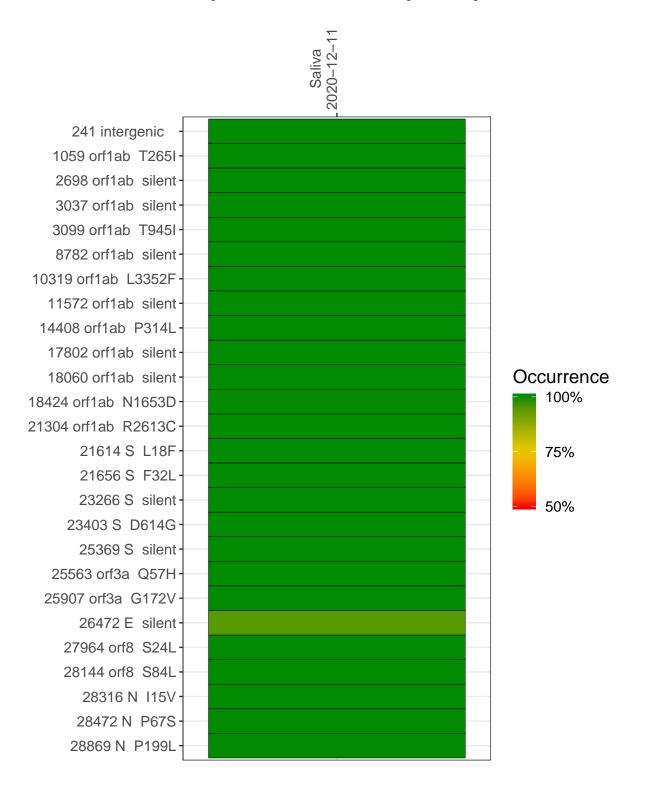
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. The code base for this analysis can be found (here).

Table 1. Sample summary.

Experiment	Туре	Input genomes	Sample type	Sample date	Largest contig (KD)	Reference read coverage	Reference read coverage (>= 5 reads)
VSP0534-1	single experiment	NA	Saliva	2020-12-11	18.02	100.0%	98.9%

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 for 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 2020–12–11

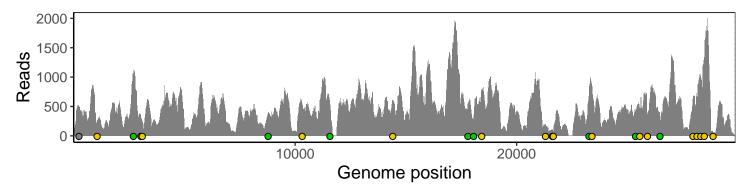
	2020-12-11				
241 intergenic	493				
1059 orf1ab T265I	223				
2698 orf1ab silent	1044				
3037 orf1ab silent	314				
3099 orf1ab T945I	251				
8782 orf1ab silent	386				
10319 orf1ab L3352F	538				
11572 orf1ab silent	520				
14408 orf1ab P314L	582				
17802 orf1ab silent	539				
18060 orf1ab silent	310				
18424 orf1ab N1653D	794				
21304 orf1ab R2613C	190				
21614 S L18F	90				
21656 S F32L	65				
23266 S silent	663				
23403 S D614G	859				
25369 S silent	346				
25563 orf3a Q57H	480				
25907 orf3a G172V	345				
26472 E silent	359				
27964 orf8 S24L	619				
28144 orf8 S84L	645				
28316 N I15V	988				
28472 N P67S	1240				
28869 N P199L	67				
	4 1				



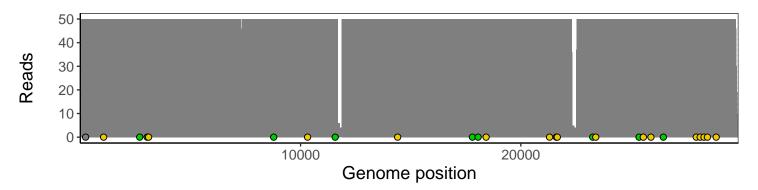
Analyses of individual experiments and composite results.

VSP0534-1 | 2020-12-11 | Saliva | 2753 | 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.

