

COVID-19 subject 205

2020-09-04

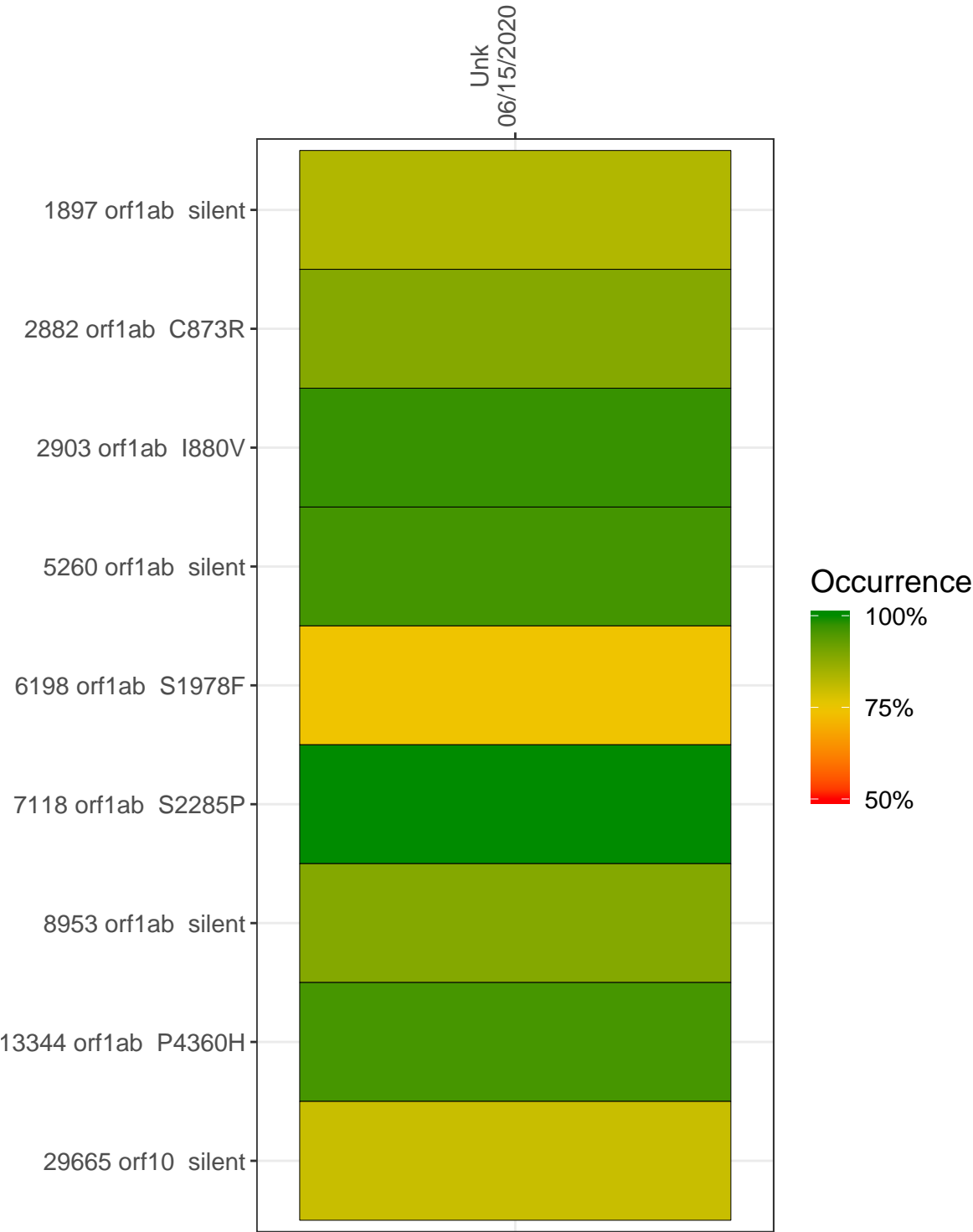
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. The code base for this analysis can be found ([here](#)).

Table 1. Sample summary.

Experiment	Type	Input genomes	Sample type	Sample date	Largest contig (KD)	Reference read coverage	Reference read coverage (>= 5 reads)
VSP0209-1	single experiment	NA	Unk	06/15/2020	NA	4.8%	2.2%

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.



Unk
06/15/2020

1897 orf1ab silent

59

2882 orf1ab C873R

859

2903 orf1ab I880V

2409

5260 orf1ab silent

52

6198 orf1ab S1978F

27

7118 orf1ab S2285P

7

8953 orf1ab silent

18

13344 orf1ab P4360H

573

29665 orf10 silent

15

Base change

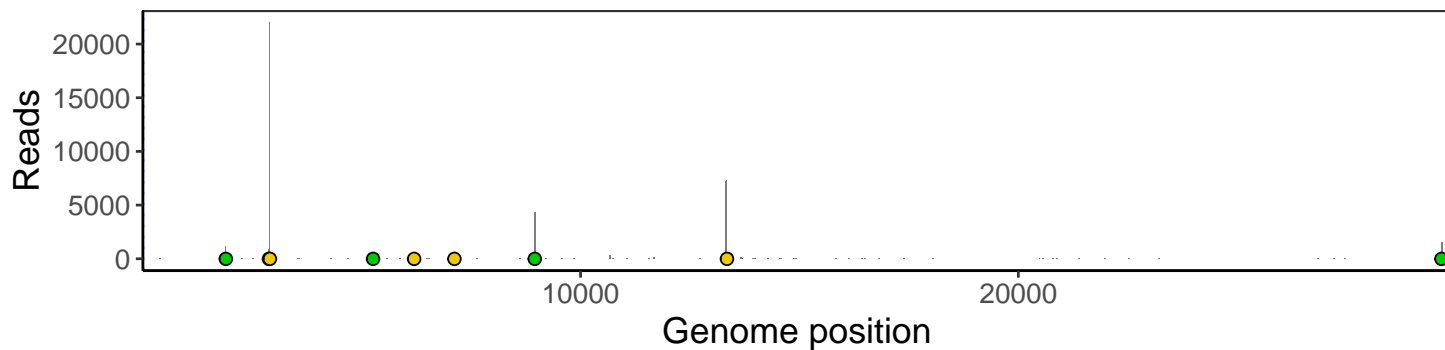


VSP0209-1

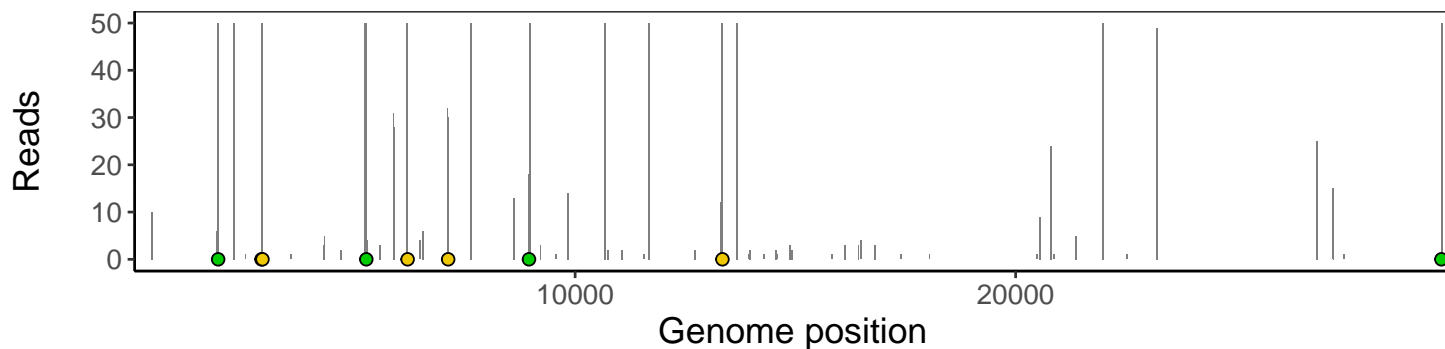
Analyses of individual experiments and composite results.

VSP0209-1 | 06/15/2020 | Unk | 205 | NA 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.



No contig data available.