# COVID-19 subject 228

2020-08-28

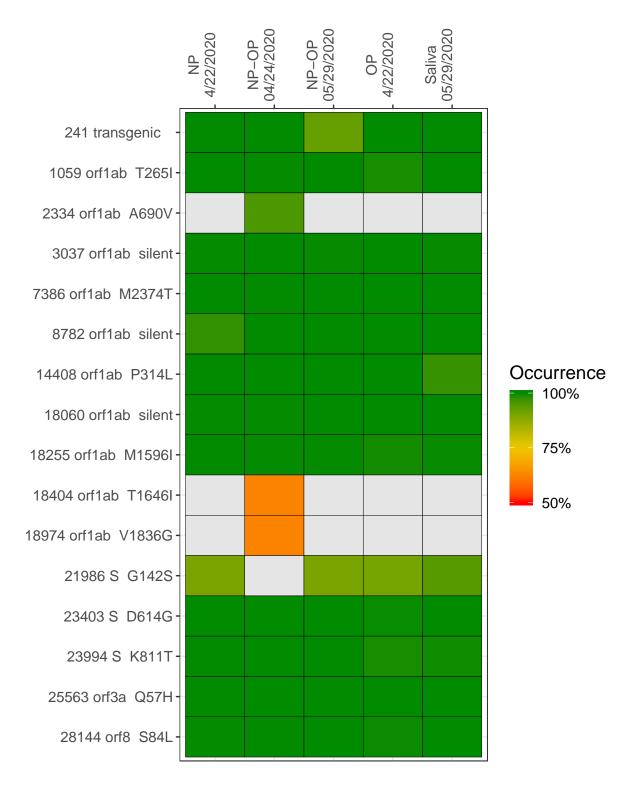
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. The code base for this analysis can be found (here).

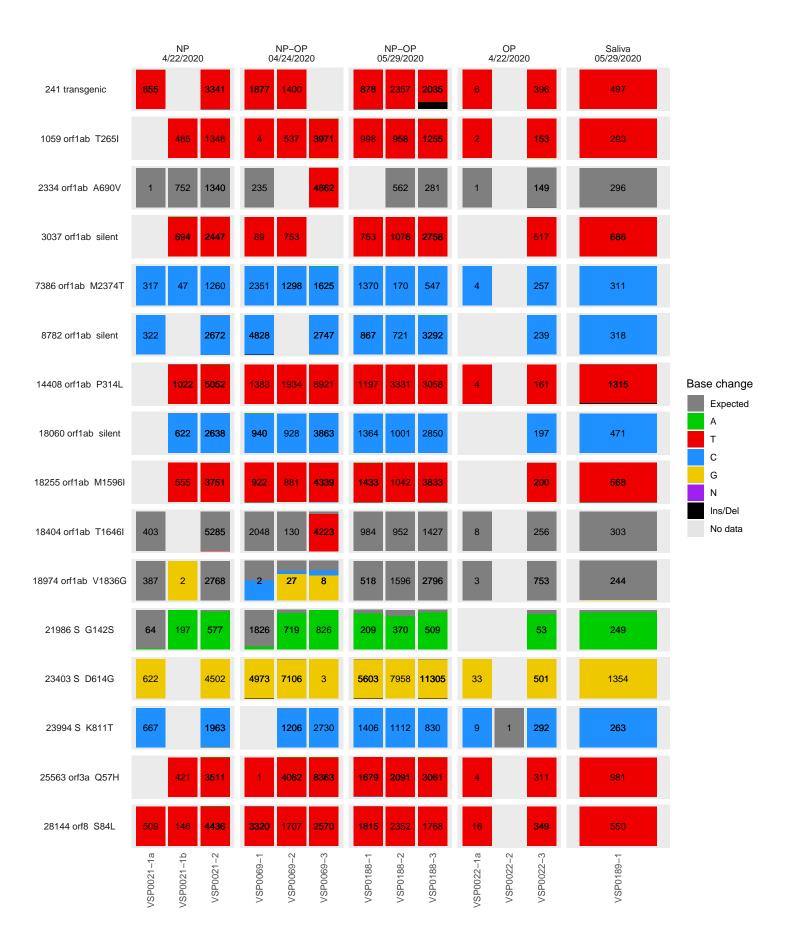
Table 1. Sample summary.

Experiment	Type	Input genomes	Sample type	Sample date	Largest contig (KD)	Reference read coverage	Reference read coverage $(>= 5 \text{ reads})$
VSP0021	composite	NA	NP	4/22/2020	29.82	99.8%	99.8%
VSP0022	composite	NA	OP	4/22/2020	29.68	99.8%	98.8%
VSP0069	composite	NA	NP-OP	04/24/2020	18.75	98.5%	98.5%
VSP0187	composite	NA	ETA	05/29/2020	NA	NA	NA
VSP0188	composite	NA	NP-OP	05/29/2020	29.90	99.8%	99.8%
VSP0021-1a	single experiment	2.20e+03	NP	4/22/2020	0.52	64.8%	61.9%
VSP0021-1b	single experiment	2.20e+03	NP	4/22/2020	0.57	63.0%	62.1%
VSP0021-2	single experiment	1.10e+04	NP	4/22/2020	29.82	99.8%	99.8%
VSP0022-1a	single experiment	2.68e + 05	OP	4/22/2020	1.75	78.3%	37.7%
VSP0022-2	single experiment	1.34e + 06	OP	4/22/2020	0.24	6.1%	0.0%
VSP0022-3	single experiment	1.34e + 06	OP	4/22/2020	29.68	99.8%	98.8%
VSP0069-1	single experiment	9.75e + 01	NP-OP	04/24/2020	6.54	92.2%	88.2%
VSP0069-2	single experiment	9.75e + 01	NP-OP	04/24/2020	9.29	90.5%	90.0%
VSP0069-3	single experiment	9.75e + 01	NP-OP	04/24/2020	6.37	87.1%	85.5%
VSP0187-1	single experiment	6.02e+01	ETA	05/29/2020	NA	NA	NA
VSP0187-2	single experiment	3.01e+02	ETA	05/29/2020	NA	NA	NA
VSP0188-1	single experiment	2.04e+03	NP-OP	05/29/2020	9.60	93.7%	93.5%
VSP0188-2	single experiment	1.02e+04	NP-OP	05/29/2020	22.61	99.1%	99.1%
VSP0188-3	single experiment	1.02e+04	NP-OP	05/29/2020	29.87	99.8%	99.8%
VSP0189-1	single experiment	8.51e + 04	Saliva	05/29/2020	29.82	99.8%	99.8%

#### 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.

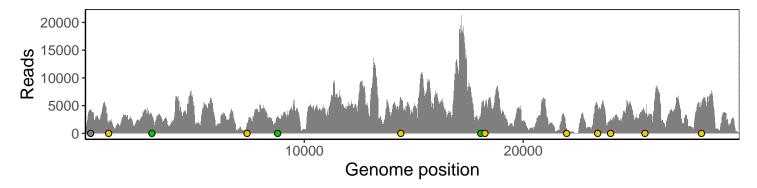




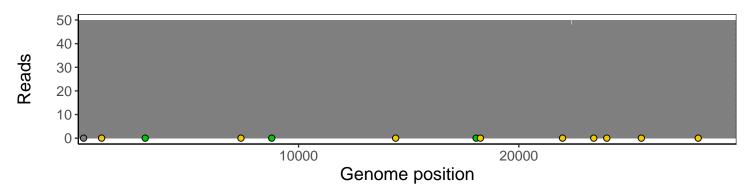
## Analyses of individual experiments and composite results.

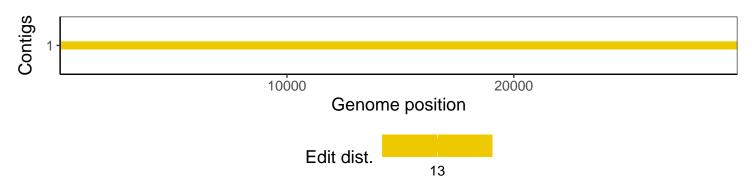
#### $VSP0021 \mid 4/22/2020 \mid NP \mid 228n \mid composite result$

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.



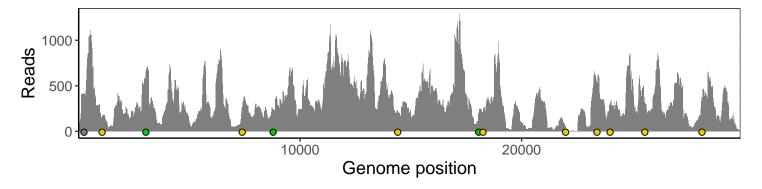
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



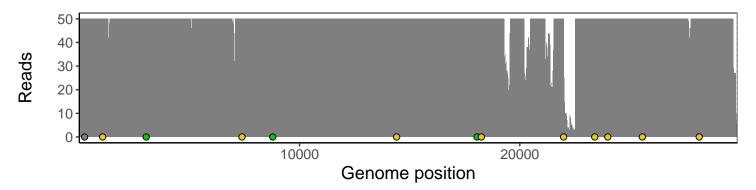


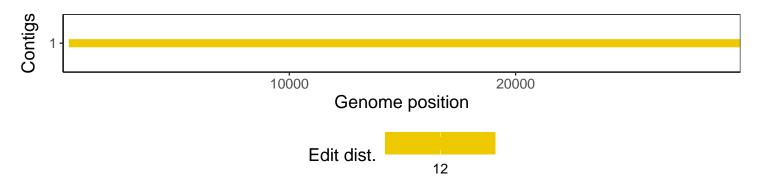
#### $VSP0022 \mid 4/22/2020 \mid OP \mid 2280 \mid composite result$

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.



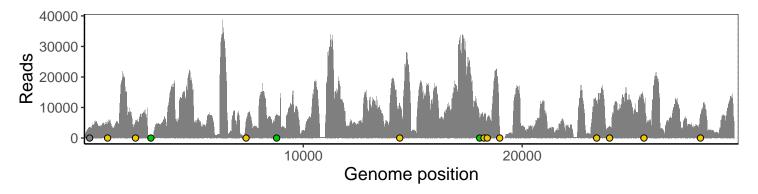
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



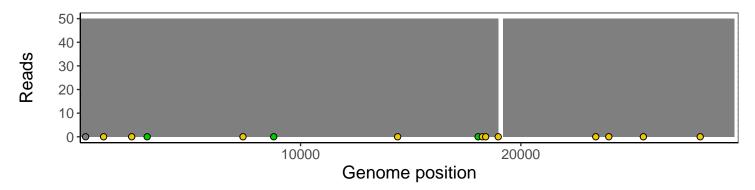


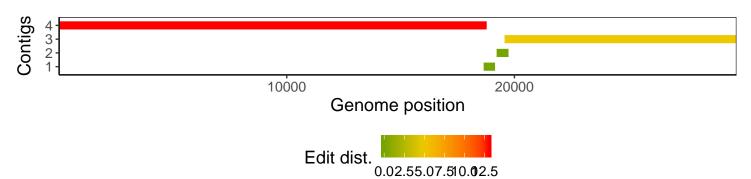
#### $VSP0069 \mid 04/24/2020 \mid NP-OP \mid 228no-t \mid composite result$

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.





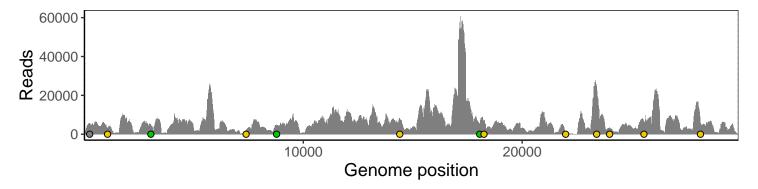
VSP0187 | 05/29/2020 | ETA | 228e-q | composite result

No pileup data available.

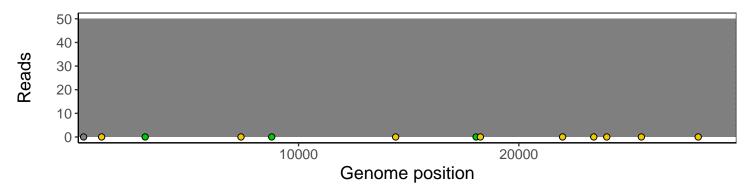
No contig data available.

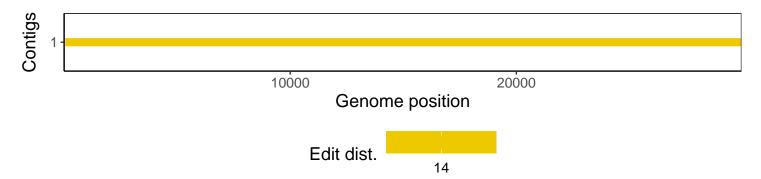
#### VSP0188 | 05/29/2020 | NP-OP | 228no-q | composite result

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.



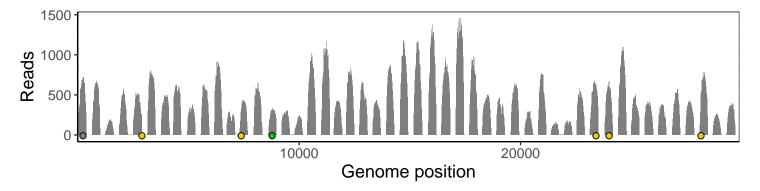
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



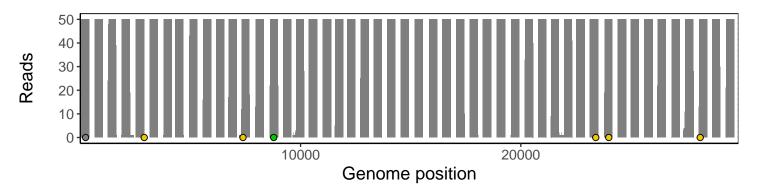


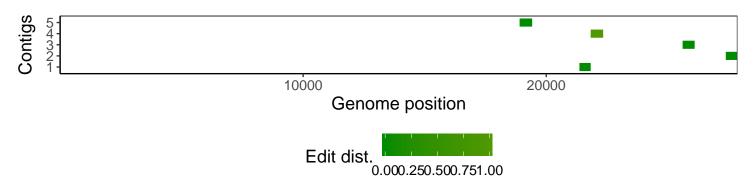
#### VSP0021-1a | 4/22/2020 | NP | 228n | 2200 genomes | single experiment

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.



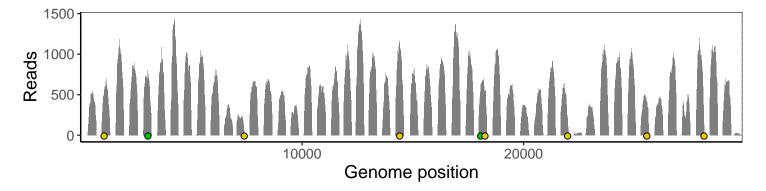
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



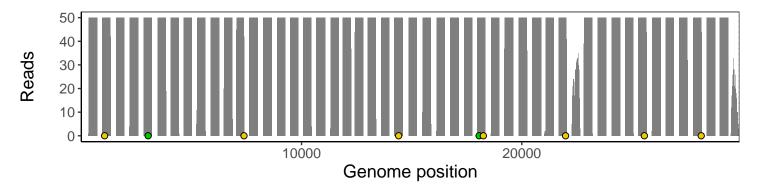


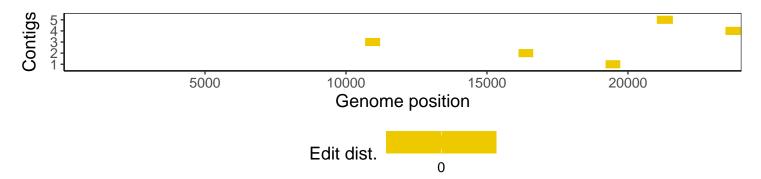
#### VSP0021-1b | 4/22/2020 | NP | 228n | 2200 genomes | single experiment

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.



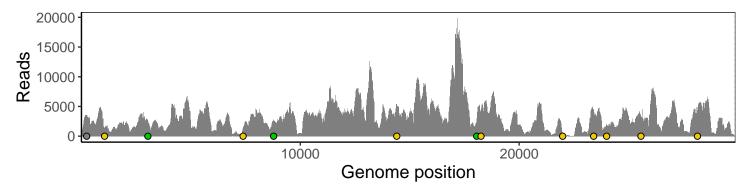
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



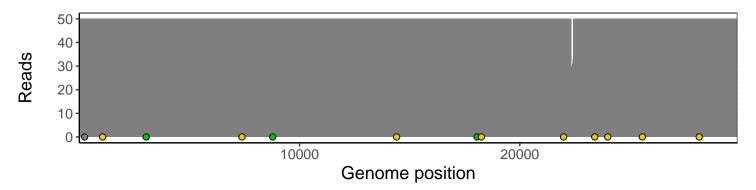


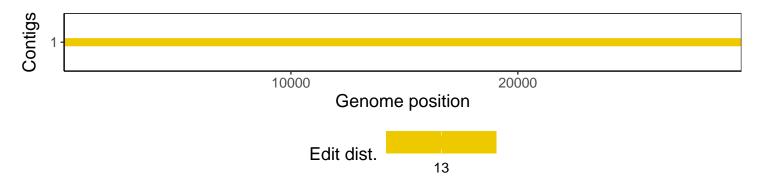
# VSP0021-2 | 4/22/2020 | NP | 228n | 11000 genomes | single experiment

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.



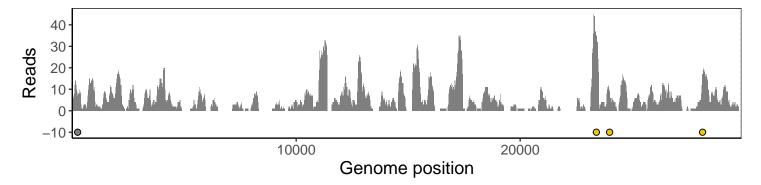
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



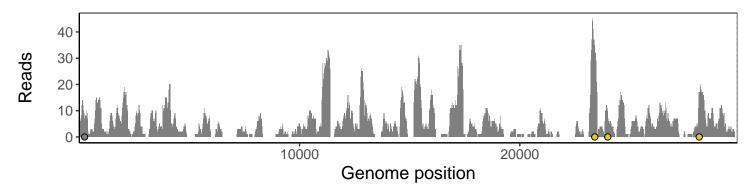


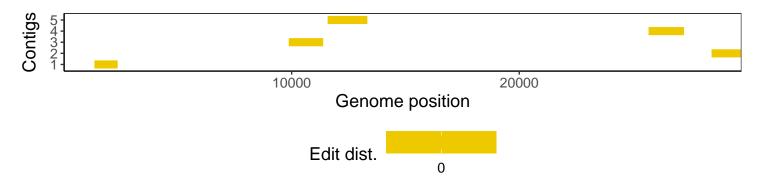
## VSP0022-1a | 4/22/2020 | OP | 228<br/>o | 268000 genomes | single experiment

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.



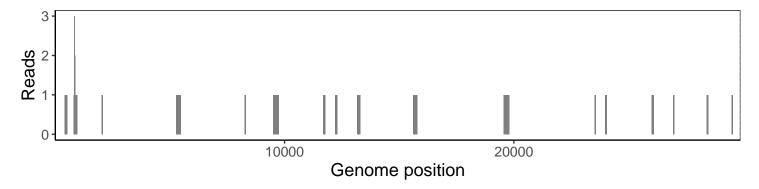
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



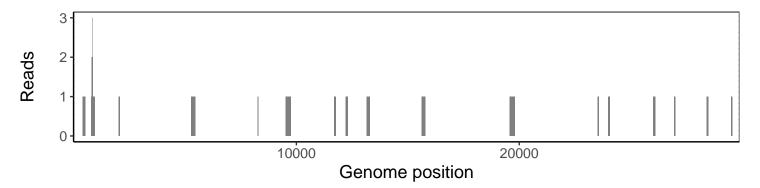


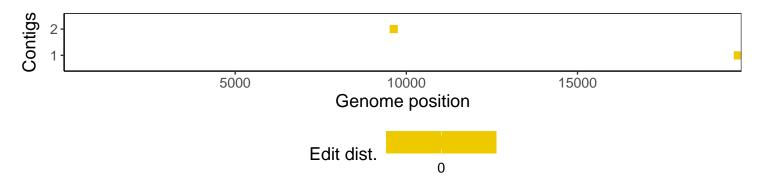
## VSP0022-2 | 4/22/2020 | OP | 228<br/>o | 1340000 genomes | single experiment

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.



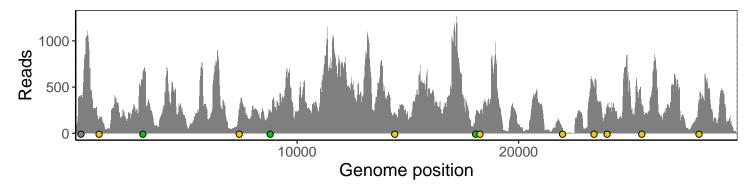
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



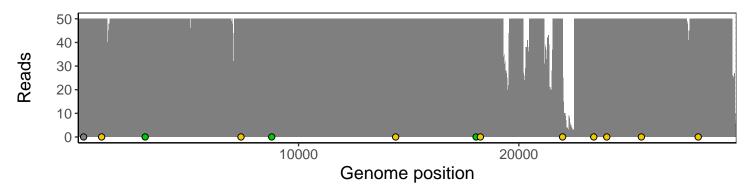


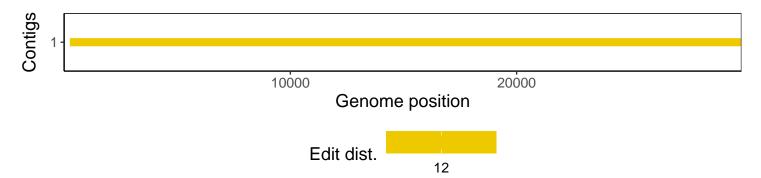
#### VSP0022-3 | 4/22/2020 | OP | 228<br/>o | 1340000 genomes | single experiment

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.



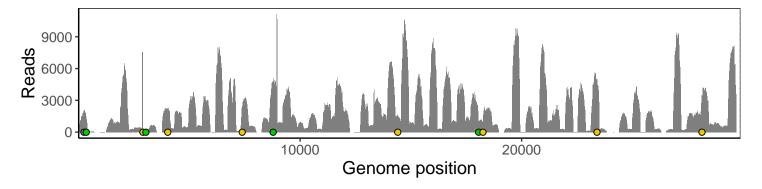
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



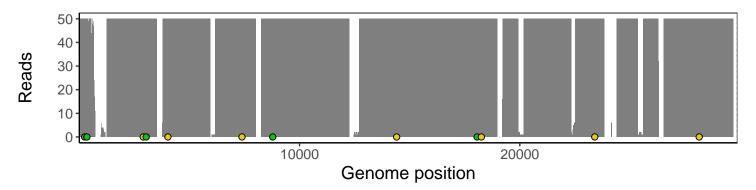


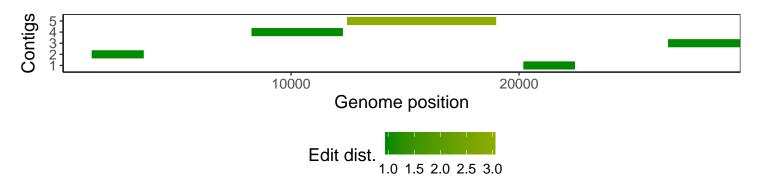
## VSP0069-1 | 04/24/2020 | NP-OP | 228no-t | 97.5 genomes | single experiment

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.



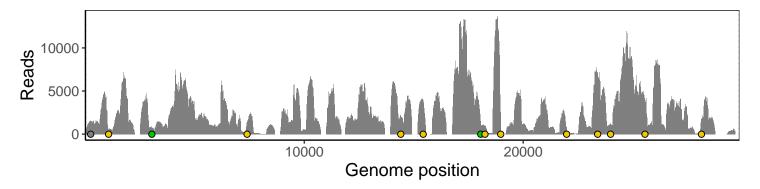
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



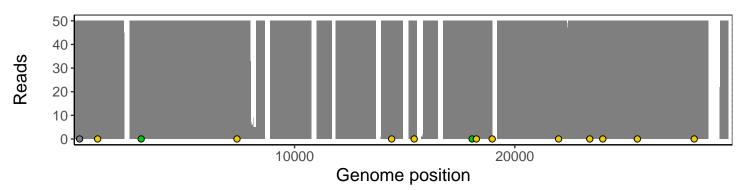


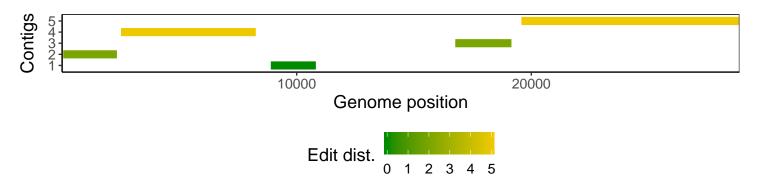
## $VSP0069-2\mid 04/24/2020\mid NP-OP\mid 228no-t\mid 97.5\ genomes\mid single\ experiment$

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.



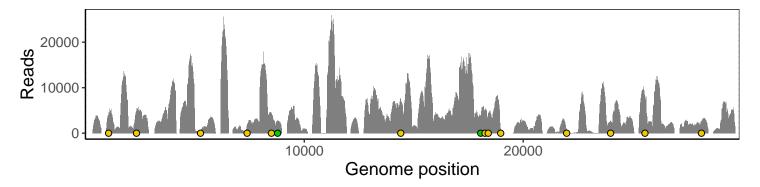
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



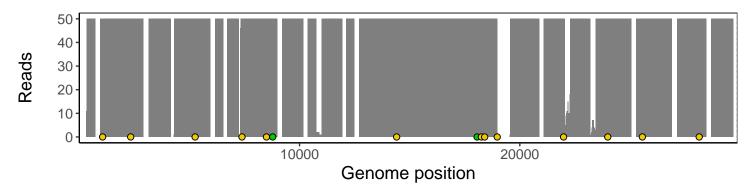


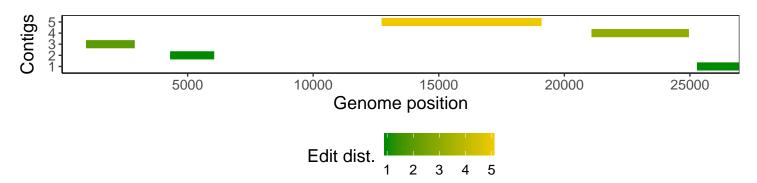
## VSP0069-3 | 04/24/2020 | NP-OP | 228<br/>no-t | 97.5 genomes | single experiment

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.





VSP0187-1 | 05/29/2020 | ETA | 228e-q | 60.2 genomes | single experiment

No pileup data available.

No contig data available.

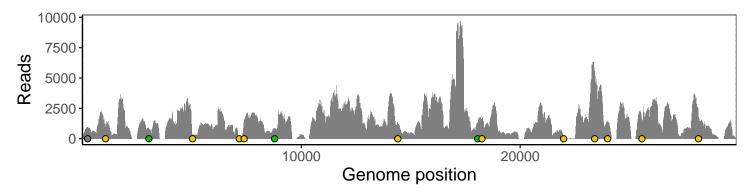
VSP0187-2 | 05/29/2020 | ETA | 228e-q | 301 genomes | single experiment

No pileup data available.

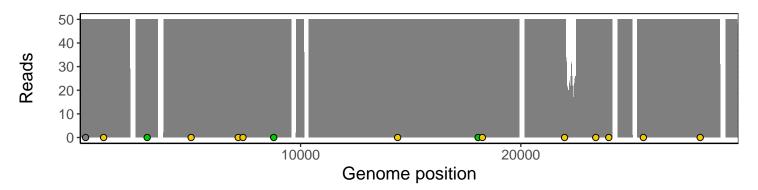
No contig data available.

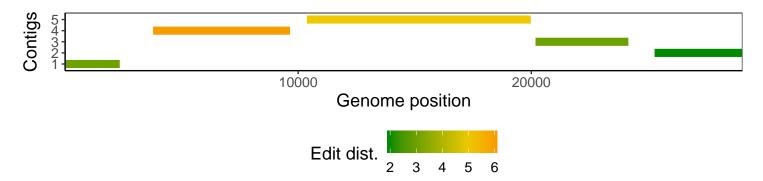
#### VSP0188-1 | 05/29/2020 | NP-OP | 228no-q | 2040 genomes | single experiment

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.



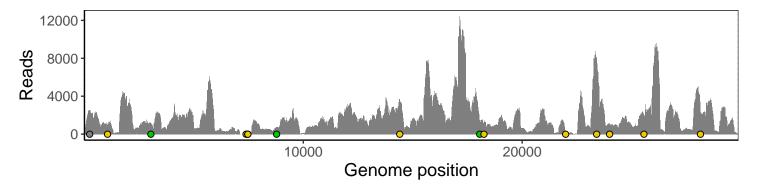
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



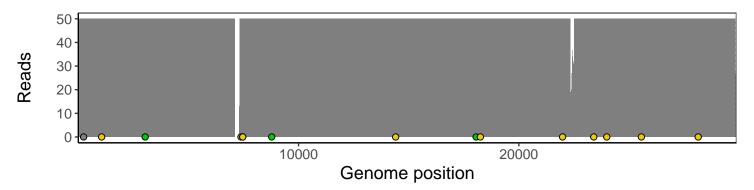


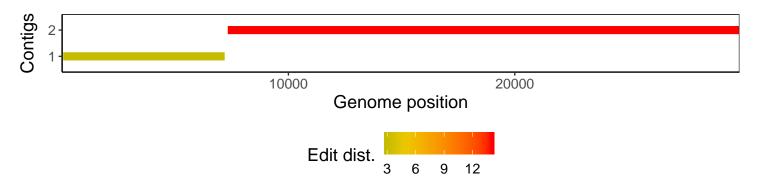
## VSP0188-2 | 05/29/2020 | NP-OP | 228<br/>no-q | 10200 genomes | single experiment

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.



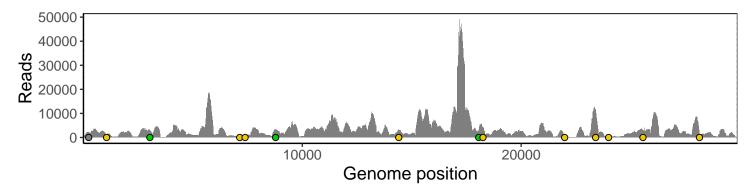
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



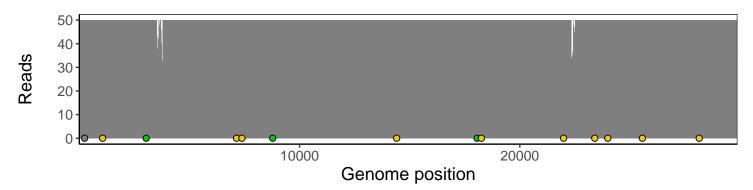


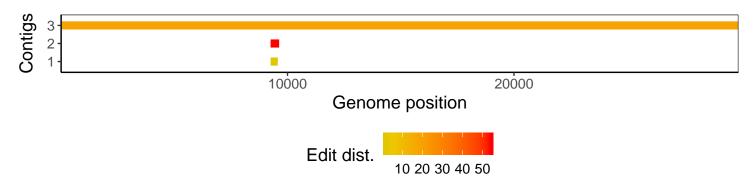
# VSP0188-3 | 05/29/2020 | NP-OP | 228<br/>no-q | 10200 genomes | single experiment

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.



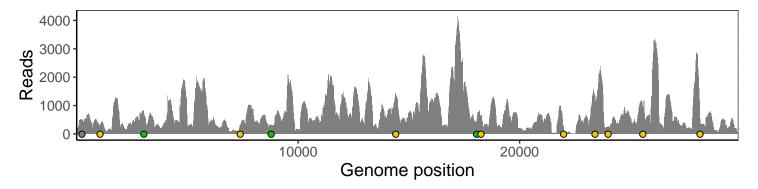
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.





## VSP0189-1 | 05/29/2020 | Saliva | 228s-q | 85100 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.

