# COVID-19 subject 237

2020-08-13

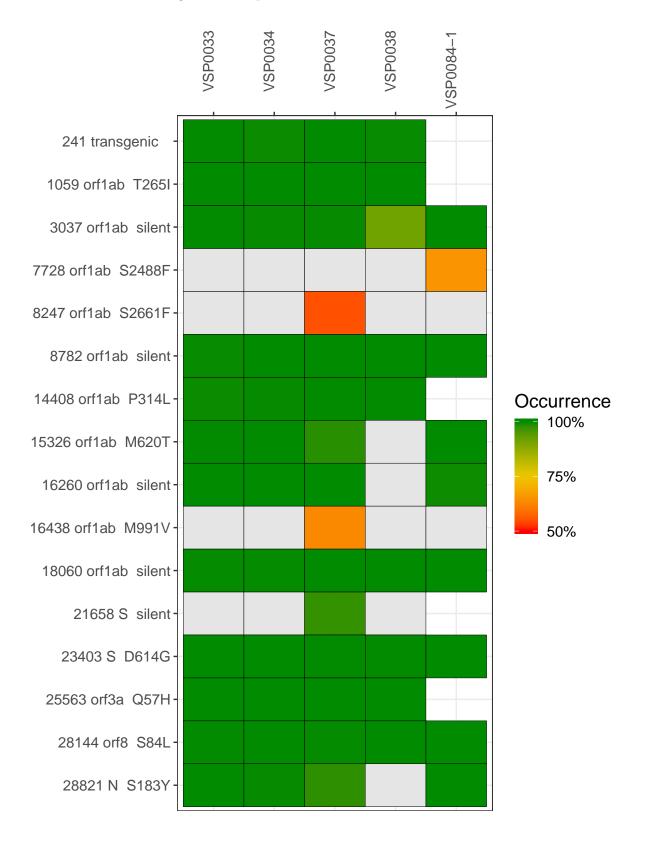
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report.

Table 1. Sample summary.

Experiment	Type	Input genomes	Sample type	Sample date	Largest contig (KD)	Reference read coverage	Reference read coverage $(>= 5 \text{ reads})$
VSP0033	composite	NA	NP-OP	4/27/2020	29.97	99.9%	99.8%
VSP0034	composite	NA	NP-OP	4/27/2020	29.84	99.8%	99.8%
VSP0037	composite	NA	NP	4/29/2020	24.92	99.3%	98.9%
VSP0038	composite	NA	OP	4/29/2020	29.92	99.9%	99.5%
VSP0033-1a	single experiment	597	NP-OP	4/27/2020	28.58	99.9%	99.8%
VSP0033-1b	single experiment	597	NP-OP	4/27/2020	0.20	12.0%	0.0%
VSP0033-2	single experiment	2985	NP-OP	4/27/2020	29.89	99.9%	99.5%
VSP0034-1a	single experiment	379000	NP-OP	4/27/2020	29.81	99.8%	99.8%
VSP0034-1b	single experiment	379000	NP-OP	4/27/2020	NA	NA	NA
VSP0034-2	single experiment	1895000	NP-OP	4/27/2020	22.45	98.8%	98.5%
VSP0037-1a	single experiment	154	NP	4/29/2020	6.66	89.3%	86.2%
VSP0037-1b	single experiment	154	NP	4/29/2020	NA	NA	NA
VSP0037-2	single experiment	770	NP	4/29/2020	11.47	95.2%	94.4%
VSP0038-1a	single experiment	412	OP	4/29/2020	0.80	64.3%	61.0%
VSP0038-1b	single experiment	412	OP	4/29/2020	0.56	64.2%	62.1%
VSP0038-2	single experiment	2060	OP	4/29/2020	22.32	99.7%	98.9%
VSP0084-1	single experiment	NA	Saliva	05/06/2020	4.70	78.4%	77.0%

#### Variants shared across samples

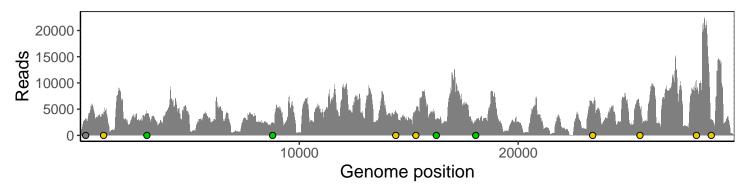
The heat map below shows how variants are shared across subject samples. The quality scores are PHRED scaled values  $[Q = -10\log 10(error\ rate)]$  where a score of 30 represents a probabilty of 99.9% that a variant is called correctly and a score of 50 represents a probabilty of 99.999% Gray tiles denote that 10 or more reads covered the variant position and the reference base was observed. Tiles are ommitted if there are less than 10 reads covering a variant position.



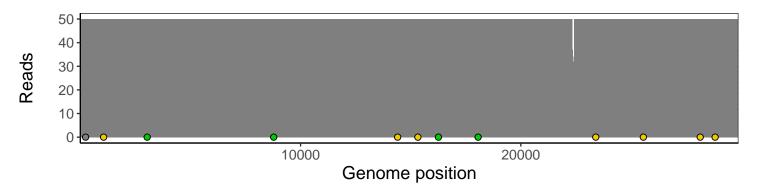
# Analyses of individual experiments and composite results.

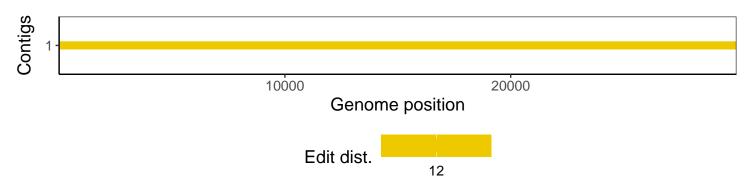
# VSP0033 | 4/27/2020 | NP-OP | 237-tri | composite result

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



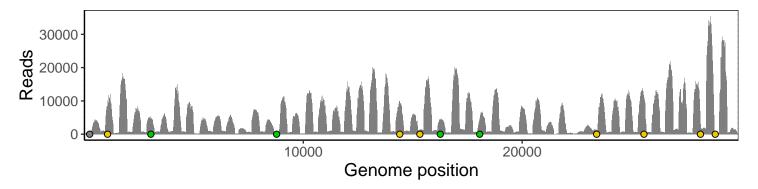
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



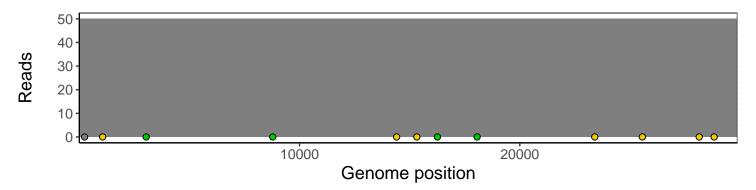


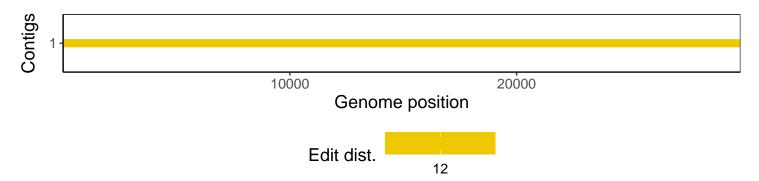
## VSP0034 | 4/27/2020 | NP-OP | 237-qia | composite result

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



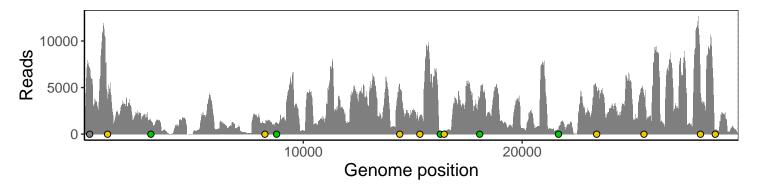
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



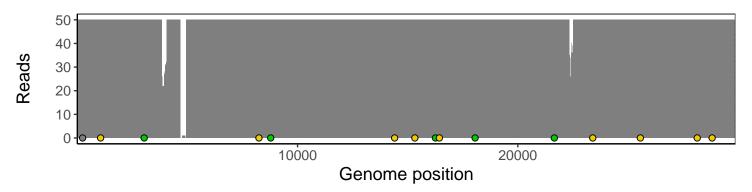


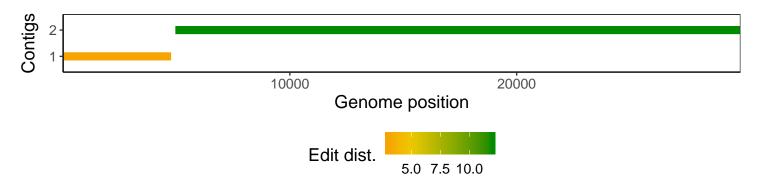
# VSP0037 | 4/29/2020 | NP | 237n-tri | composite result

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



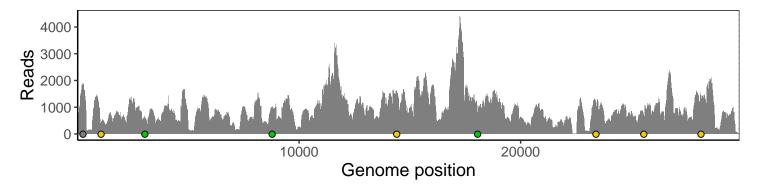
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



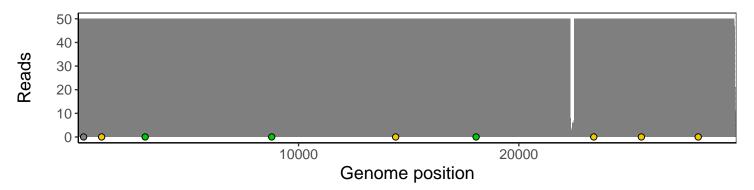


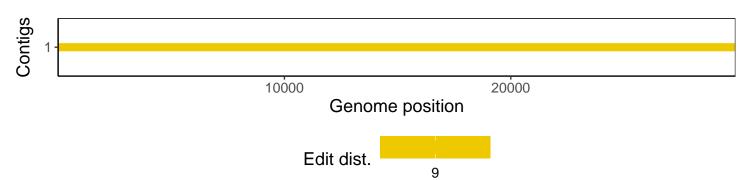
# VSP0038 | 4/29/2020 | OP | 2370-tri | composite result

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



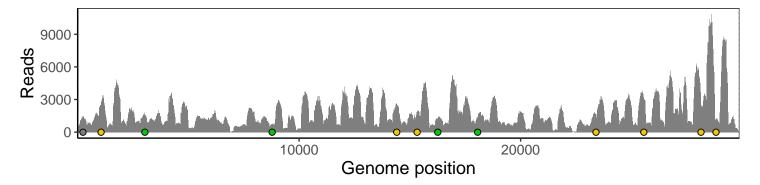
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



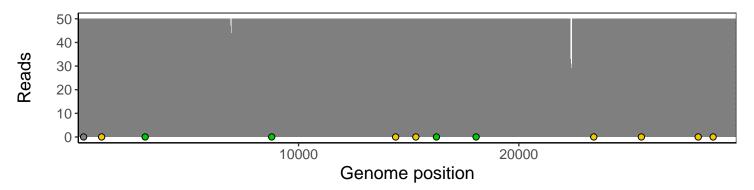


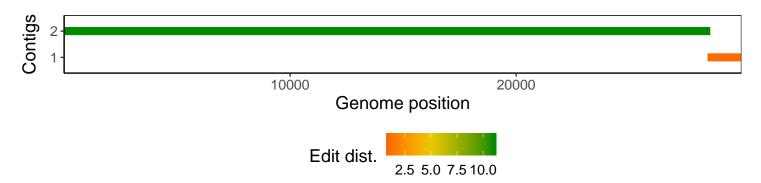
## VSP0033-1a | 4/27/2020 | NP-OP | 237-tri | 597 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



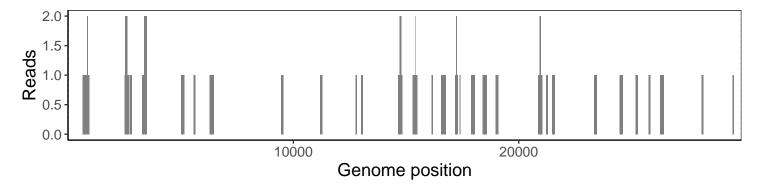
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



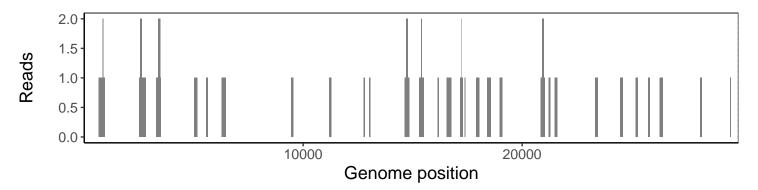


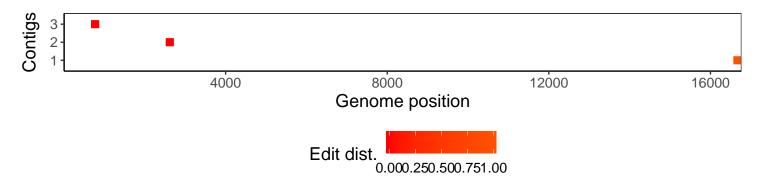
### VSP0033-1b | 4/27/2020 | NP-OP | 237-tri | 597 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



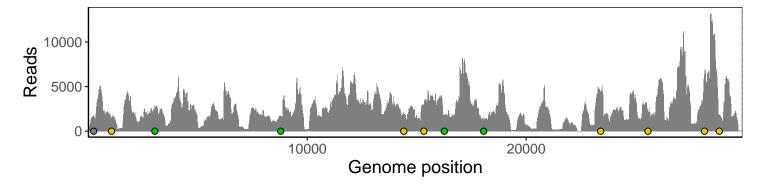
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



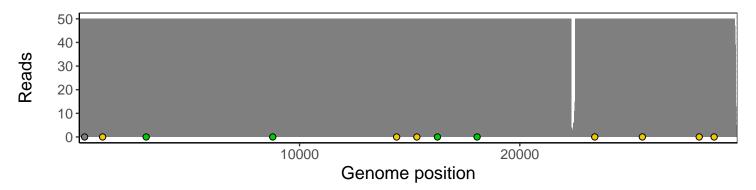


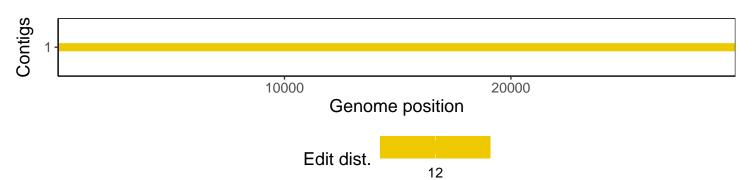
## VSP0033-2 | 4/27/2020 | NP-OP | 237-tri | 2985 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



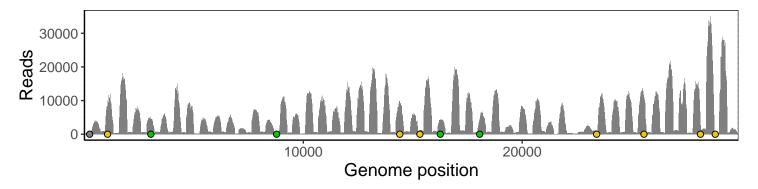
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



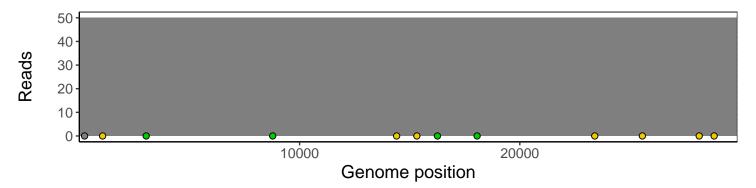


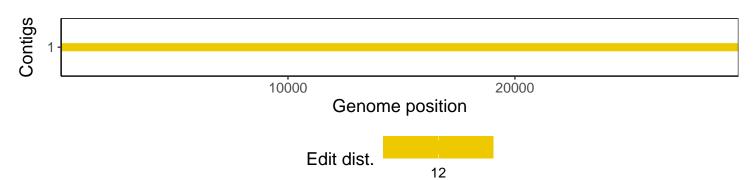
## VSP0034-1a | 4/27/2020 | NP-OP | 237-qia | 379000 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.





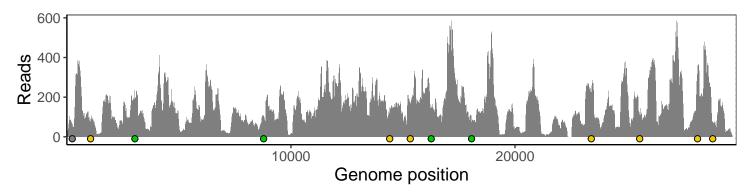
VSP0034-1<br/>b | 4/27/2020 | NP-OP | 237-qia | 379000 genomes | single experiment

No pileup data available.

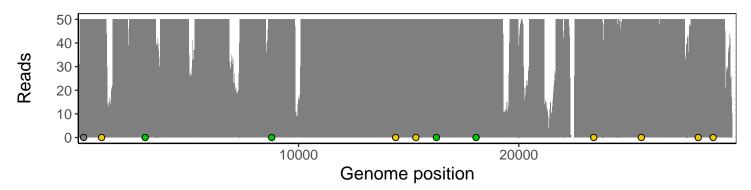
No contig data available.

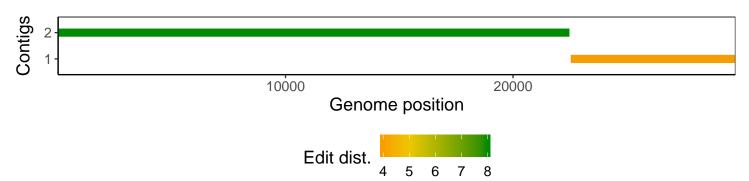
## VSP0034-2 | 4/27/2020 | NP-OP | 237-qia | 1895000 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



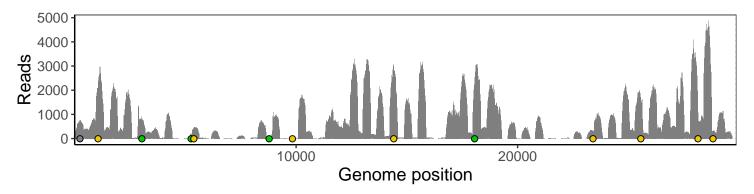
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



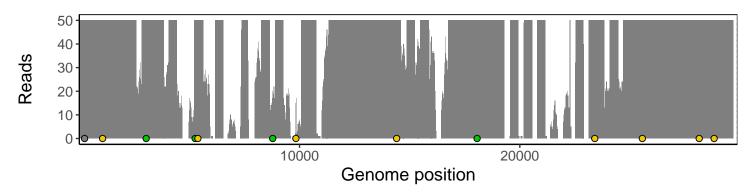


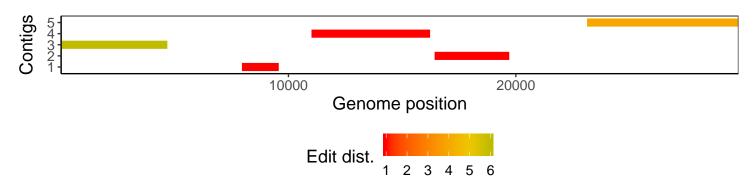
### VSP0037-1a | 4/29/2020 | NP | 237n-tri | 154 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.





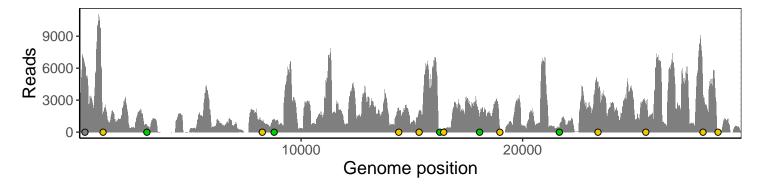
VSP0037-1b | 4/29/2020 | NP | 237<br/>n-tri | 154 genomes | single experiment

No pileup data available.

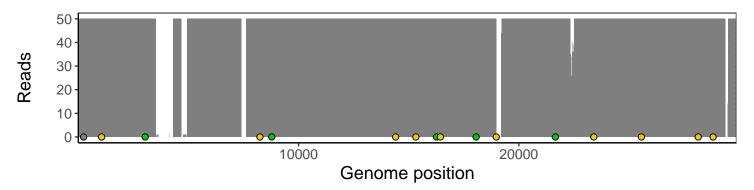
No contig data available.

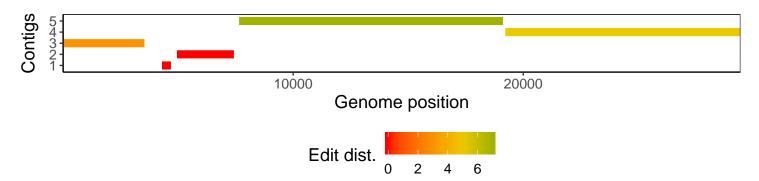
# VSP0037-2 | 4/29/2020 | NP | 237n-tri | 770 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



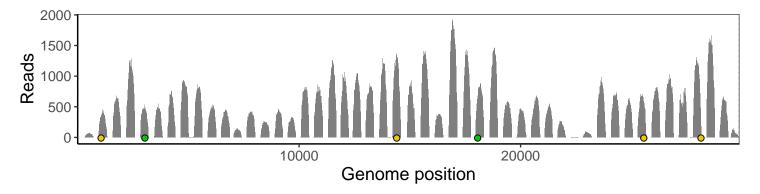
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



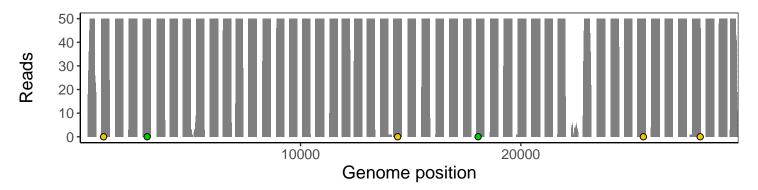


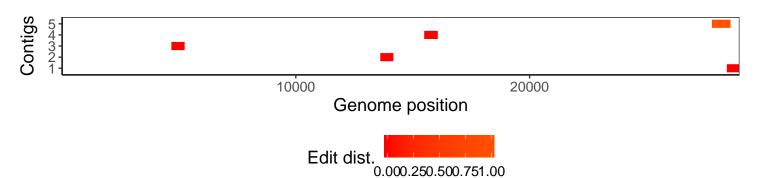
### VSP0038-1a | 4/29/2020 | OP | 2370-tri | 412 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



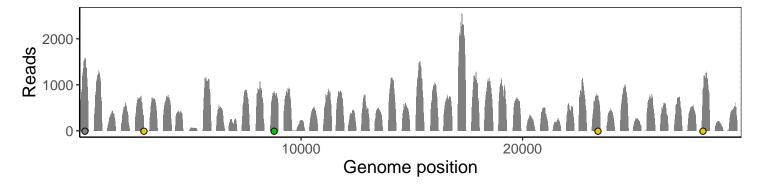
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



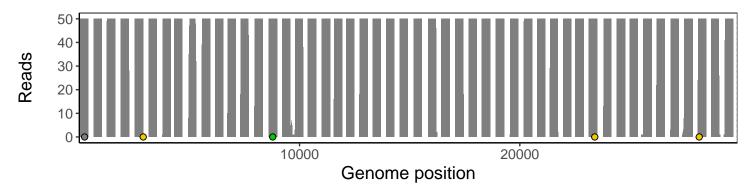


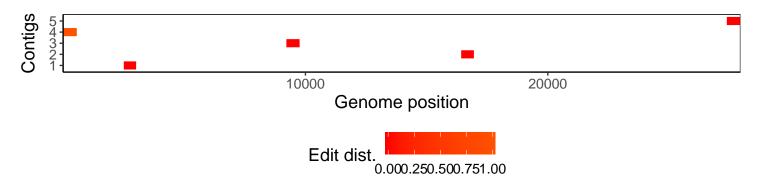
### VSP0038-1b | 4/29/2020 | OP | 2370-tri | 412 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



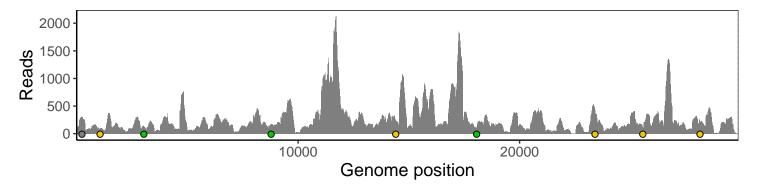
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



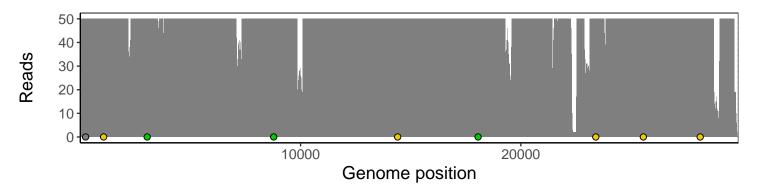


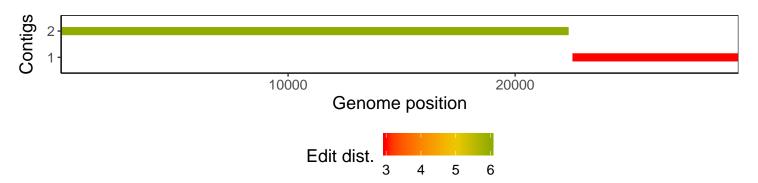
# VSP0038-2 | 4/29/2020 | OP | 2370-tri | 2060 genomes | single experiment

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.



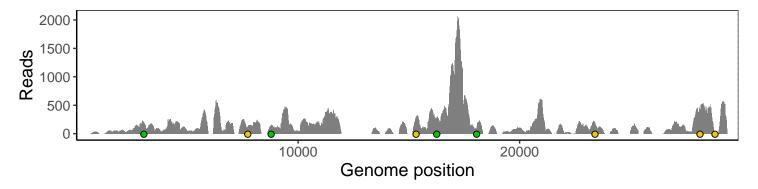
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.





### $VSP0084-1 \mid 05/06/2020 \mid Saliva \mid 237s-q \mid genomes \mid single experiment$

The plot below shows the number of reads covering each nucleotide position in the reference genome (USA-WA1-2020). 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.

