

# COVID-19 subject 235

*2021-01-10*

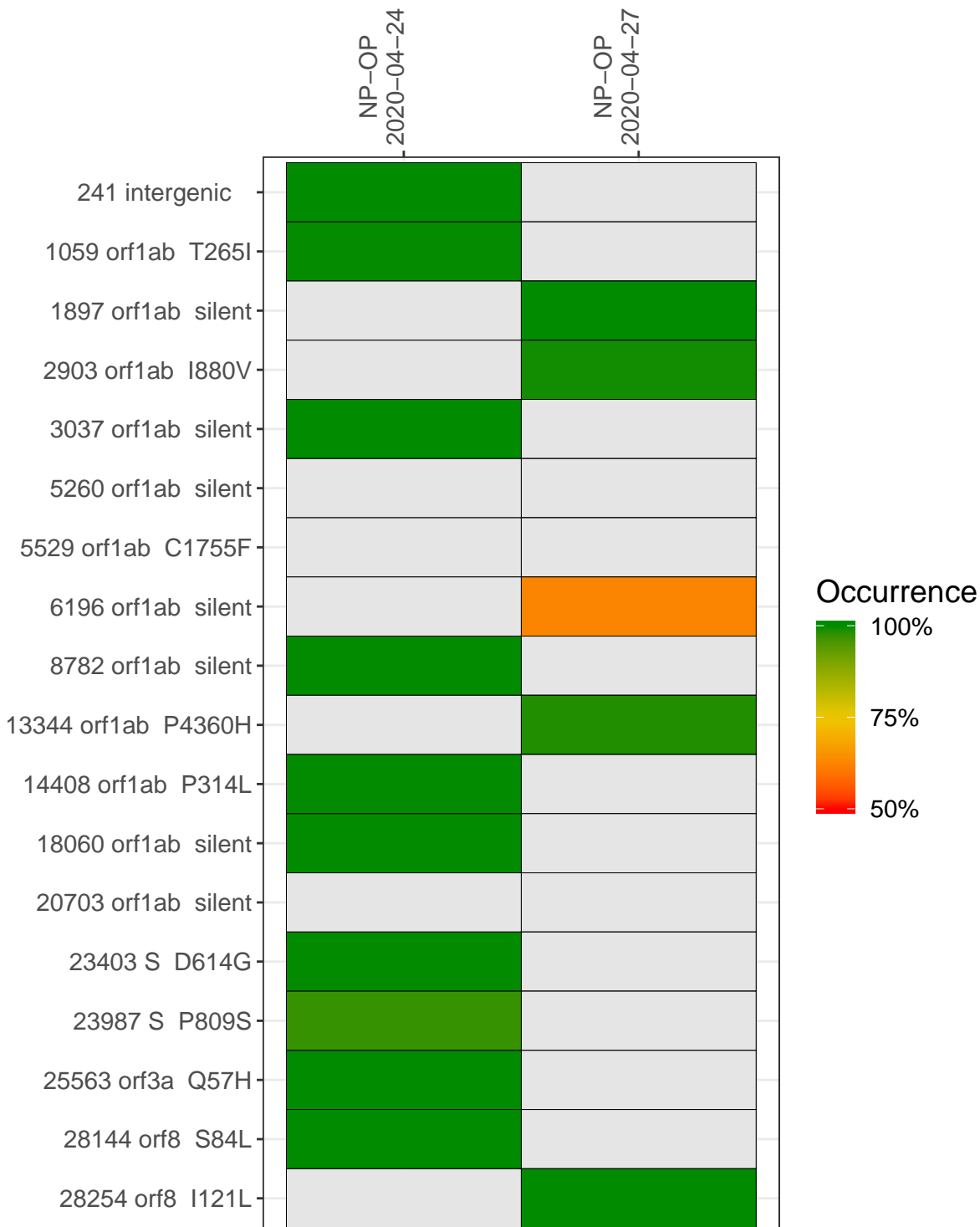
The table below provides a summary of subject samples for which sequencing data is available. The experiments column shows the number of sequencing experiments performed for each specimen. Experiment specific analyses are shown at the end of this report. The code base for this analysis can be found ([here](#)).

Table 1. Sample summary.

Experiment	Type	Input genomes	Sample type	Sample date	Largest contig (KD)	Reference read coverage	Reference read coverage (>= 5 reads)
VSP0030	composite	NA	NP-OP	2020-04-27	2.47	76.7%	54.3%
VSP0029-1m	single experiment	NA	NP-OP	2020-04-27	0.50	17.5%	12.1%
VSP0030-1m	single experiment	NA	NP-OP	2020-04-27	1.05	51.1%	20.1%
VSP0030-2	single experiment	NA	NP-OP	2020-04-27	1.96	55.0%	52.1%
VSP0080-1	single experiment	176000	NP-OP	2020-04-24	29.87	99.9%	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.



	NP-OP 2020-04-24		NP-OP 2020-04-27	
241 intergenic	3231		1	
1059 orf1ab T265I	1007			
1897 orf1ab silent	2028	15	6	1
2903 orf1ab I880V	1851	1274	83	206
3037 orf1ab silent	1844	1		244
5260 orf1ab silent	1160	2	6	2
5529 orf1ab C1755F	3107			417
6196 orf1ab silent	2455	8	20	723
8782 orf1ab silent	2504		2	
13344 orf1ab P4360H	4586	87	180	367
14408 orf1ab P314L	3499		261	596
18060 orf1ab silent	2225		343	395
20703 orf1ab silent	3696	185	258	620
23403 S D614G	7457		1	
23987 S P809S	826			
25563 orf3a Q57H	1987			494
28144 orf8 S84L	3607		1	
28254 orf8 I121L	4227	116	1	
	VSP0080-1	VSP0029-1m	VSP0030-1m	VSP0030-2

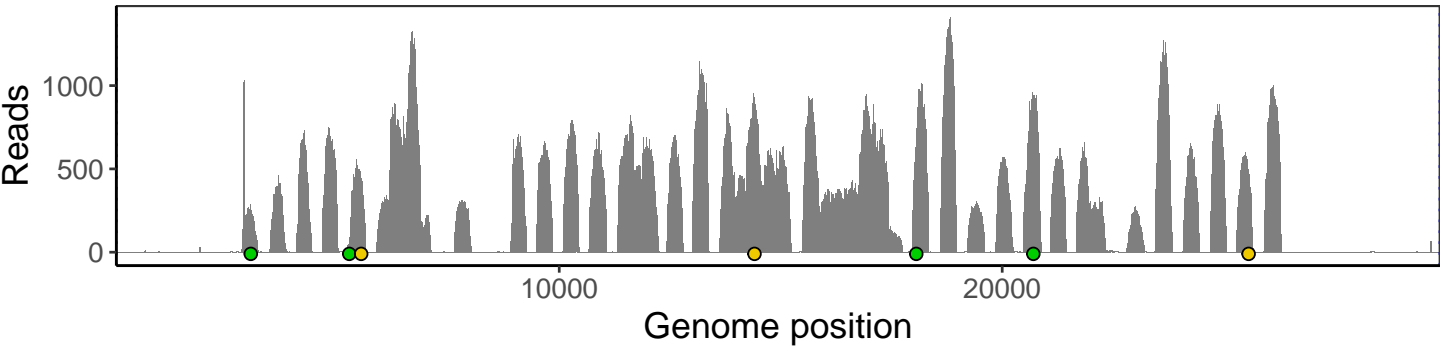
Base change



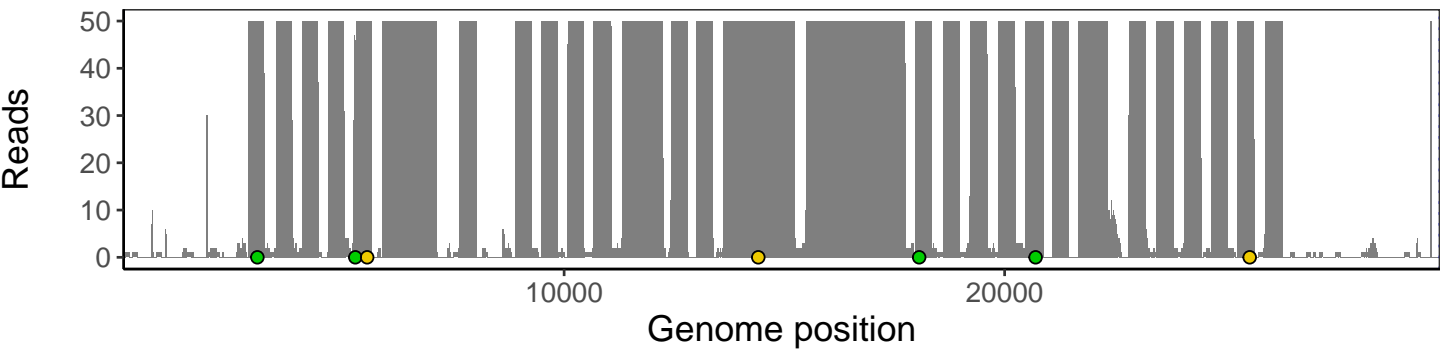
Analyses of individual experiments and composite results.

VSP0030 | 2020-04-27 | NP-OP | 235-qia | composite result

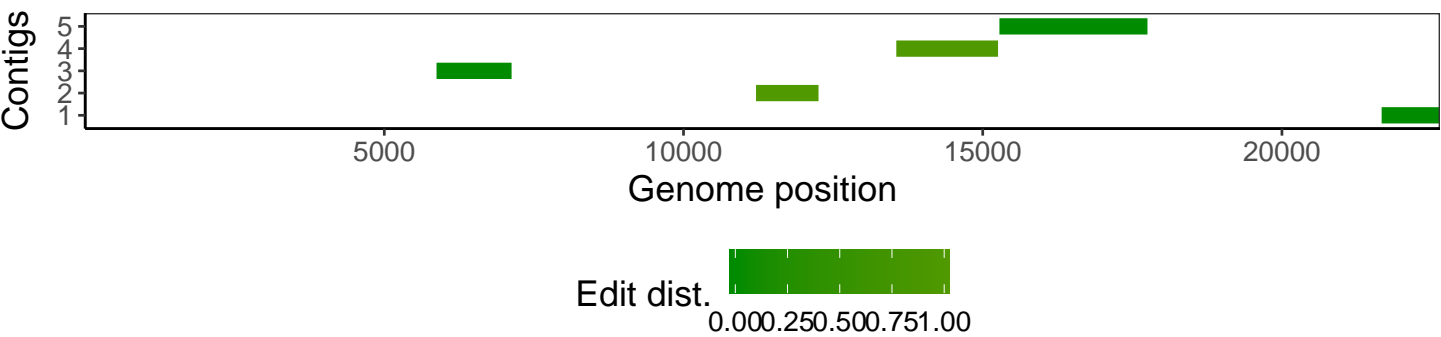
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



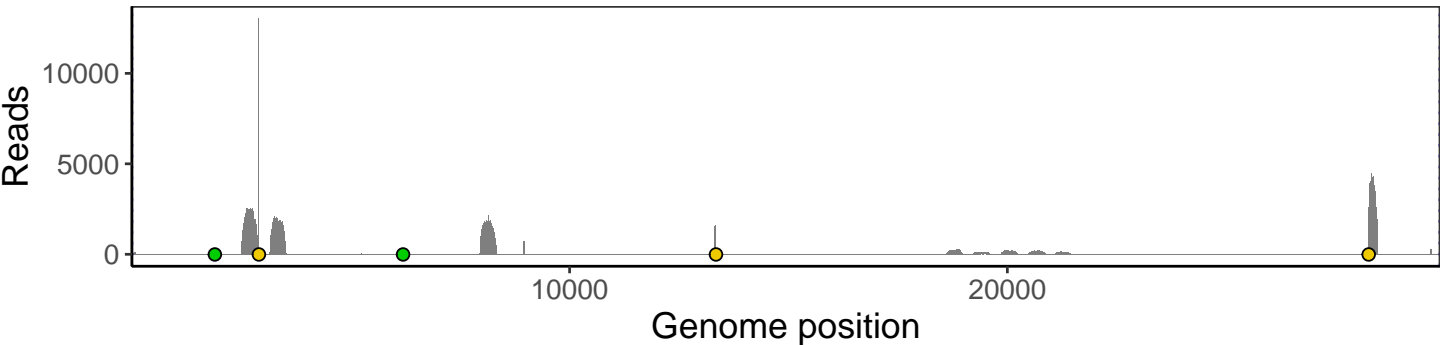
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



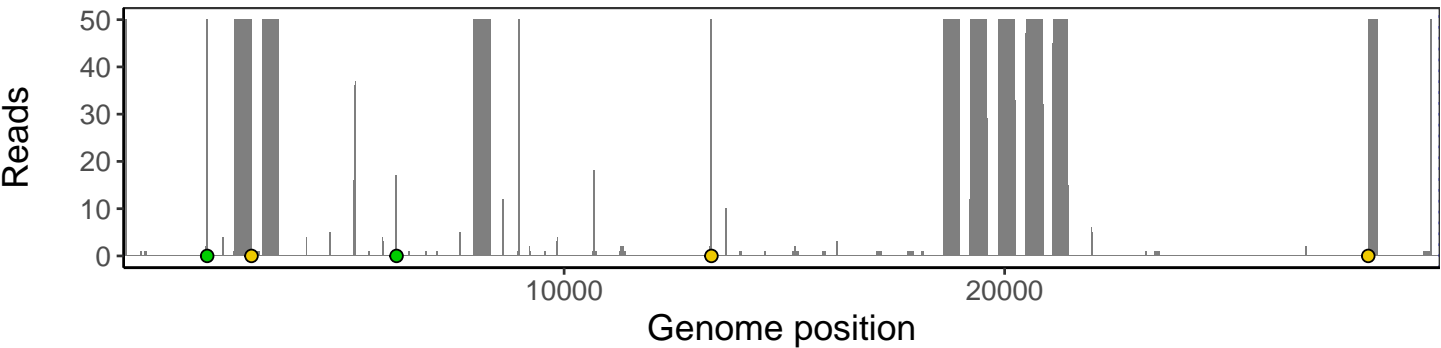
The longest five assembled contigs are shown below colored by their edit distance to the reference genome.



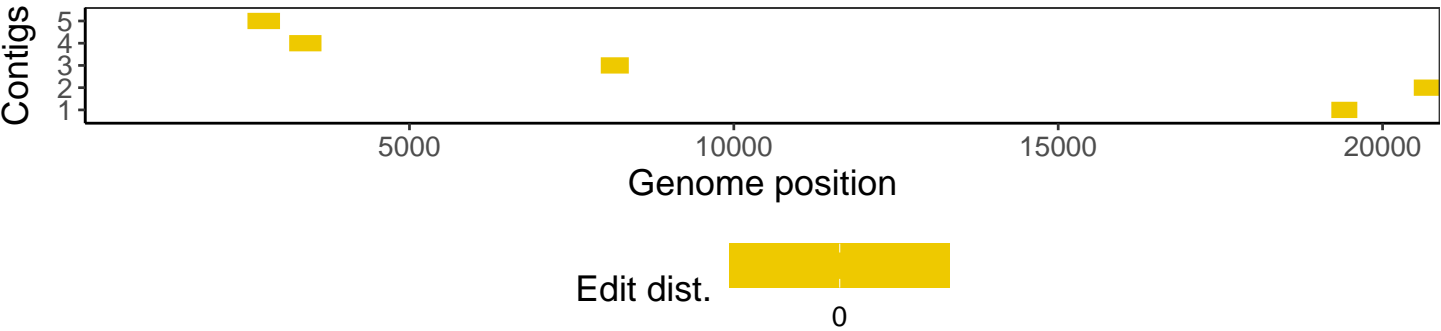
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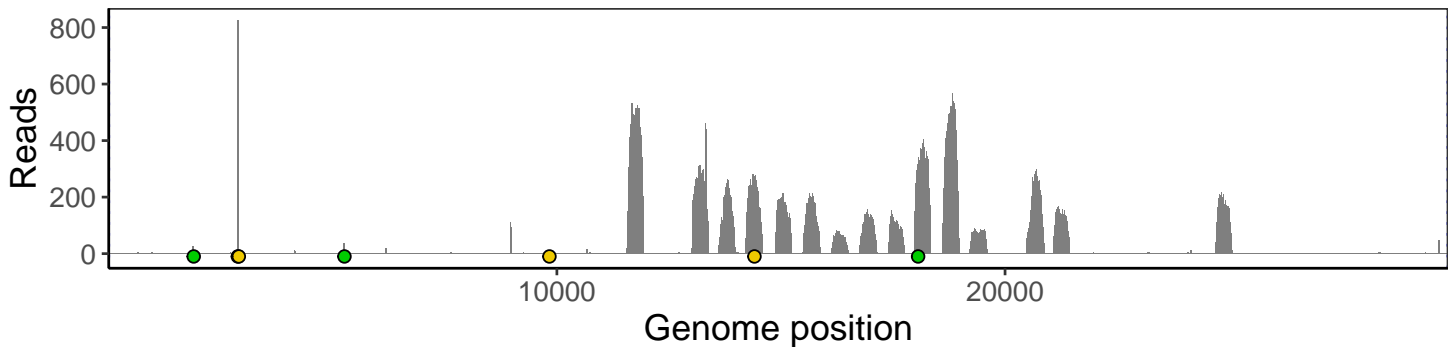
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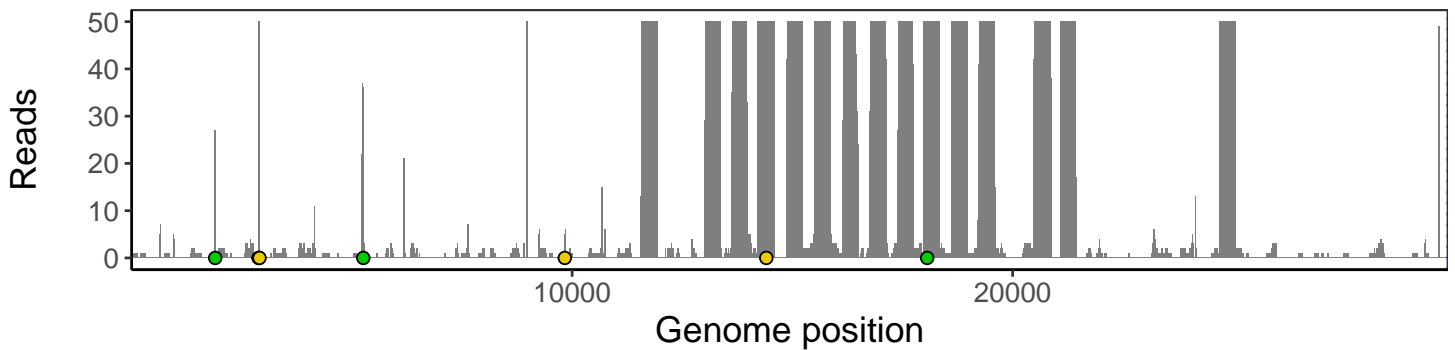
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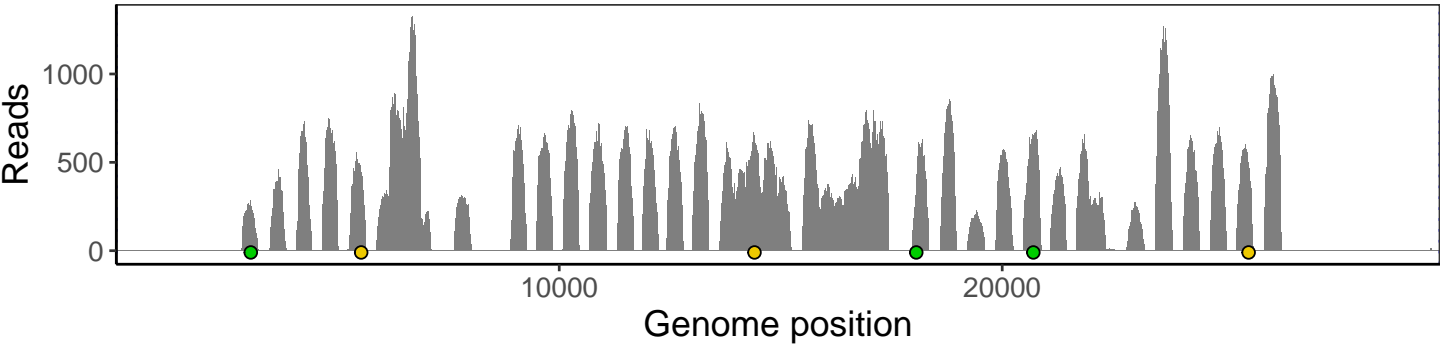
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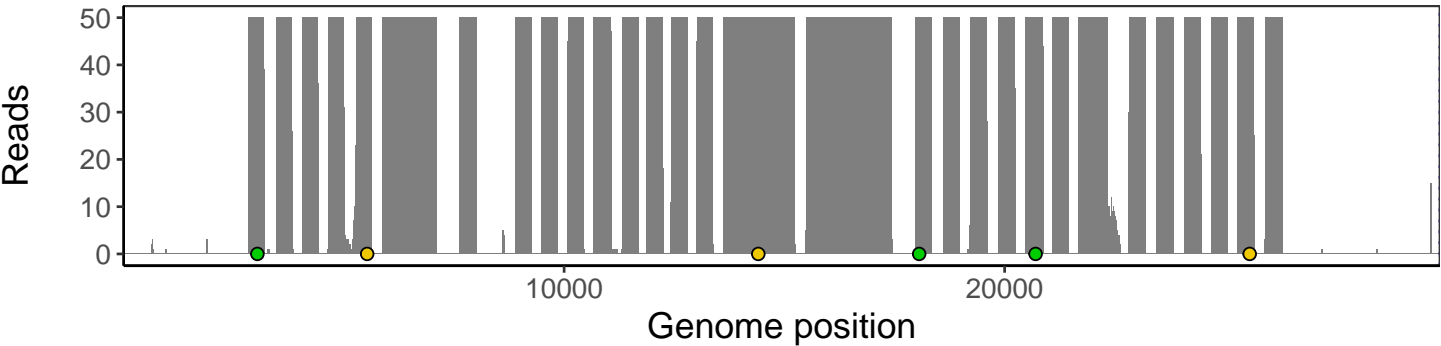
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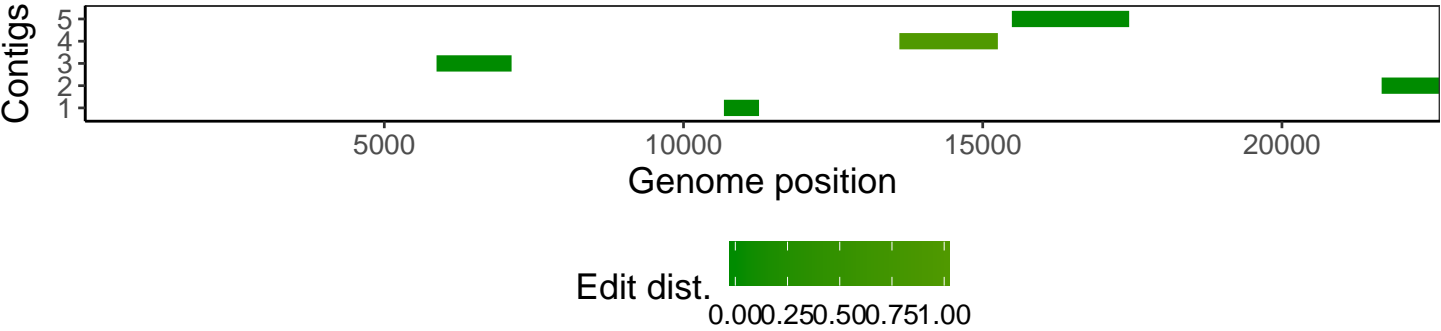
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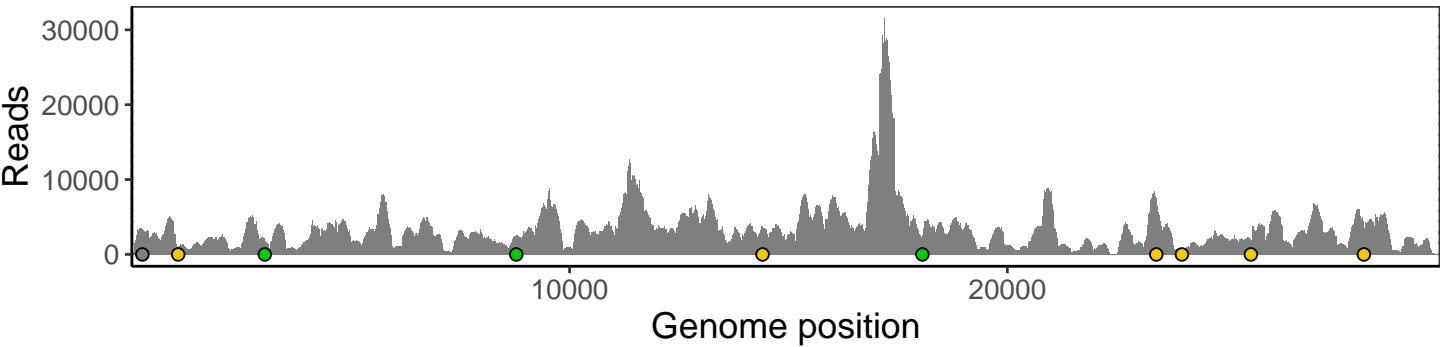
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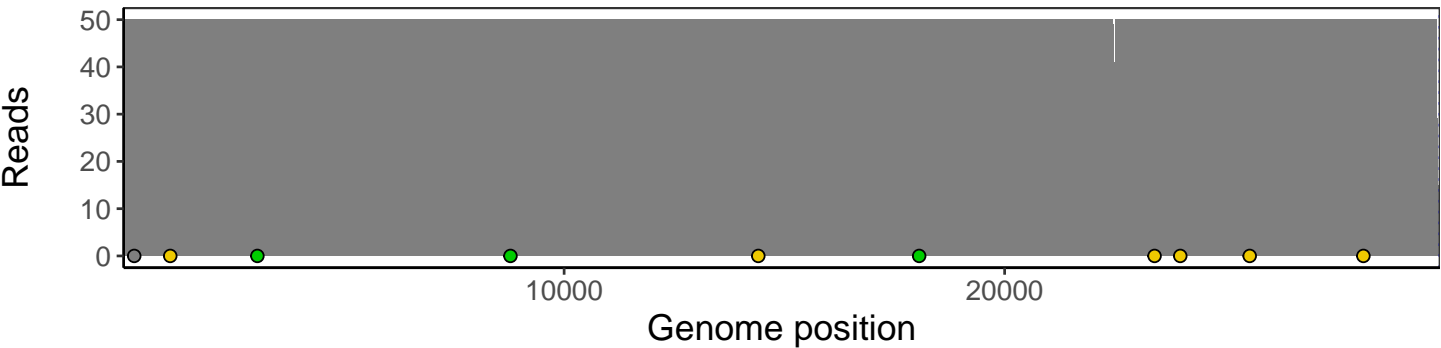
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