

COVID-19 subject 437

2021-04-17

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.

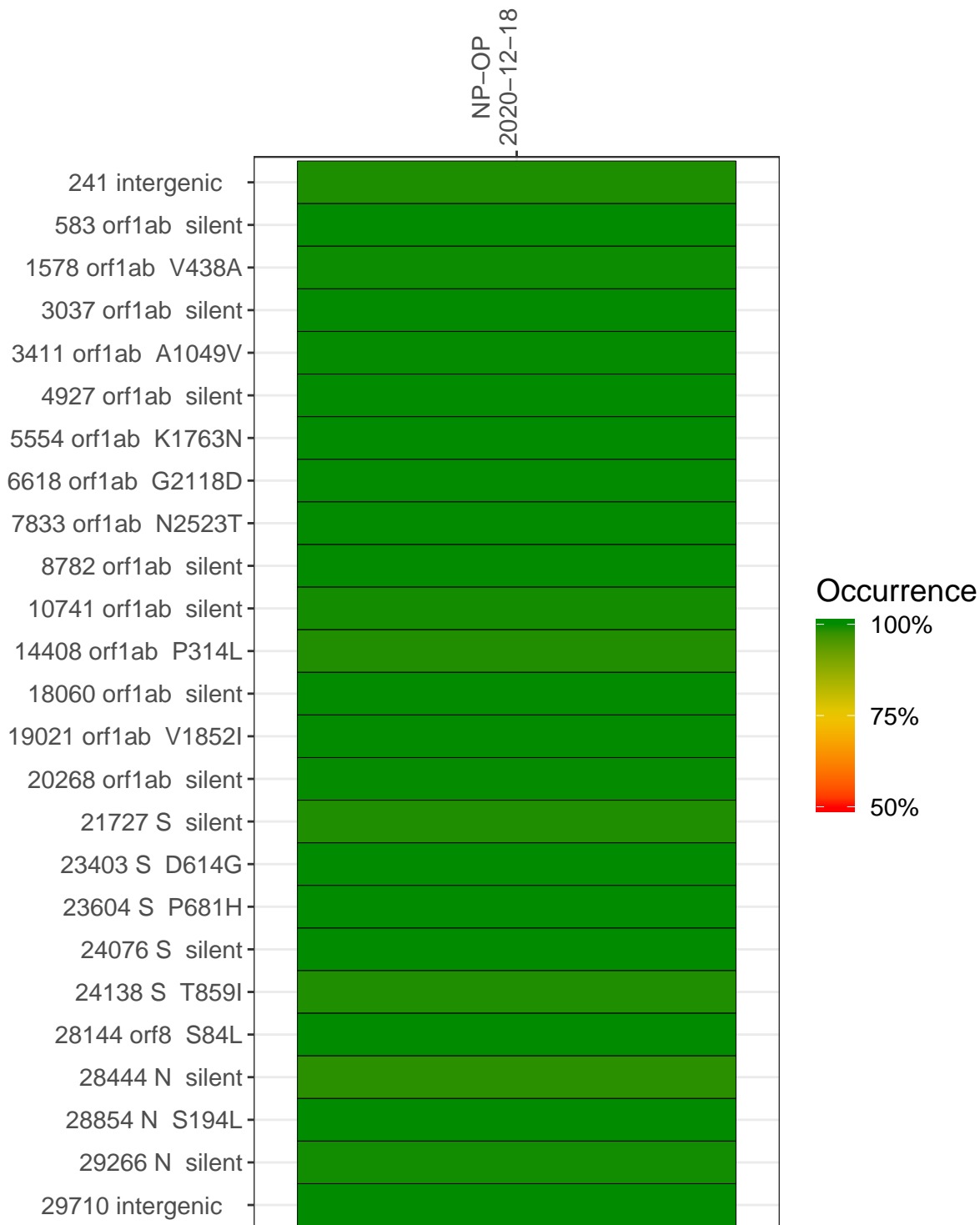
Experiment specific analyses are shown at the end of this report. Lineages are called with the Pangolin software tool (Rambaut et al 2020) for genomes with $> 90\%$ sequence coverage.

Table 1. Sample summary.

| Experiment | Type | Genomes | Sample type | Sample date | Largest contig (KD) | Lineage | Reference read coverage | Reference read coverage (≥ 5 reads) |
|------------|-------------------|---------|-------------|-------------|---------------------|---------|-------------------------|---|
| VSP0560 | composite | NA | NP-OP | 2020-12-18 | 28.05 | B.1.243 | 99.9% | 99.8% |
| VSP0560-1 | single experiment | NA | NP-OP | 2020-12-18 | 29.77 | B.1.243 | 99.9% | 99.8% |
| VSP0560-2 | single experiment | NA | NP-OP | 2020-12-18 | 15.09 | B.1.243 | 99.8% | 99.8% |

Variants shared across samples

The heat map below shows how variants (reference genome /home/everett/projects/SARS-CoV-2-Philadelphia/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 or 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
2020-12-18

| | | |
|---------------------|-------|------|
| 241 intergenic | 10010 | 4023 |
| 583 orf1ab silent | 6449 | 2322 |
| 1578 orf1ab V438A | 1523 | 517 |
| 3037 orf1ab silent | 6583 | 2438 |
| 3411 orf1ab A1049V | 9707 | 3422 |
| 4927 orf1ab silent | 12817 | 4833 |
| 5554 orf1ab K1763N | 6898 | 2330 |
| 6618 orf1ab G2118D | 8722 | 3589 |
| 7833 orf1ab N2523T | 12358 | 4416 |
| 8782 orf1ab silent | 7207 | 2354 |
| 10741 orf1ab silent | 6185 | 2127 |
| 14408 orf1ab P314L | 15332 | 5351 |
| 18060 orf1ab silent | 6253 | 2152 |
| 19021 orf1ab V1852I | 12614 | 4576 |
| 20268 orf1ab silent | 1885 | 530 |
| 21727 S silent | 4778 | 1676 |
| 23403 S D614G | 18984 | 6842 |
| 23604 S P681H | 15563 | 5779 |
| 24076 S silent | 3446 | 1165 |
| 24138 S T859I | 4906 | 1693 |
| 28144 orf8 S84L | 9336 | 3289 |
| 28444 N silent | 23936 | 7475 |
| 28854 N S194L | 2675 | 931 |
| 29266 N silent | 9815 | 3305 |
| 29710 intergenic | 1010 | 380 |

Base change

- Expected
- A
- T
- C
- G
- N
- Ins/Del
- No data

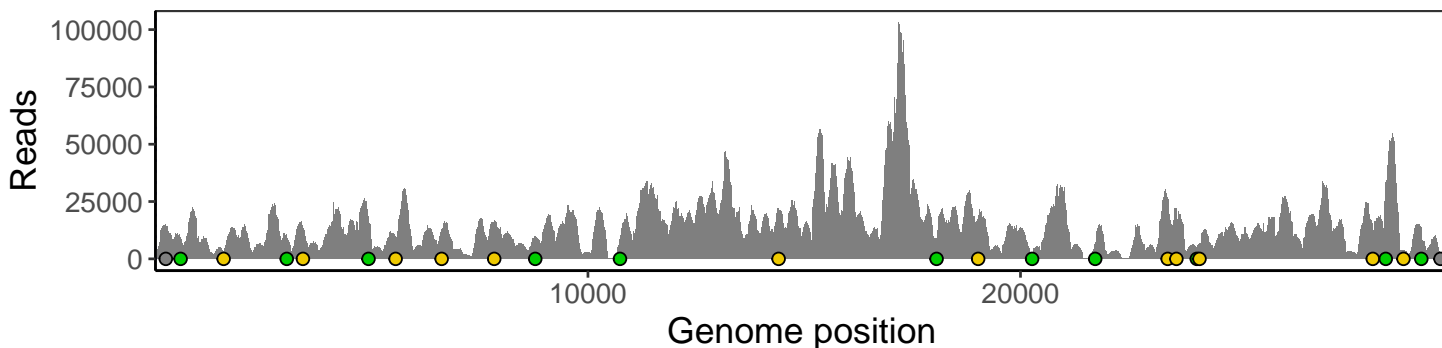
VSP0560-1

VSP0560-2

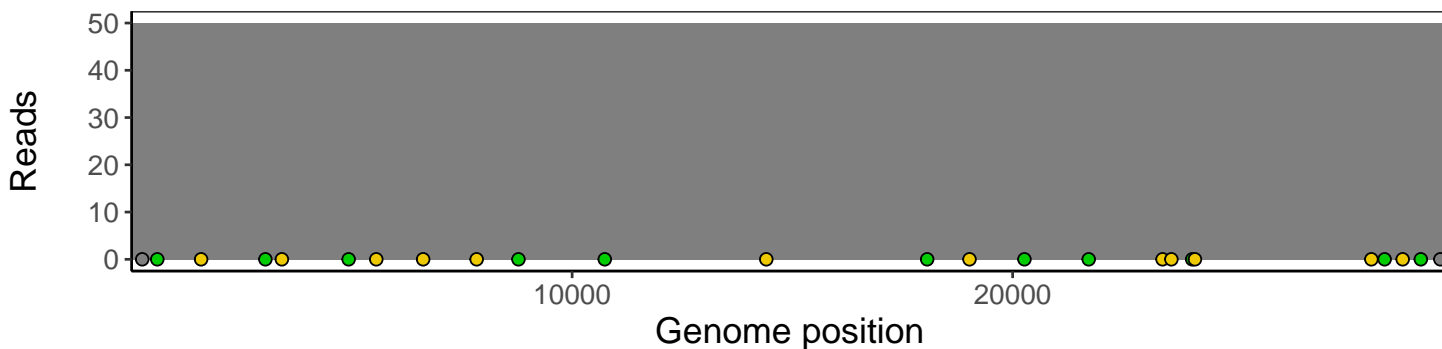
Analyses of individual experiments and composite results

VSP0560 | 2020-12-18 | NP-OP | 437no | composite result

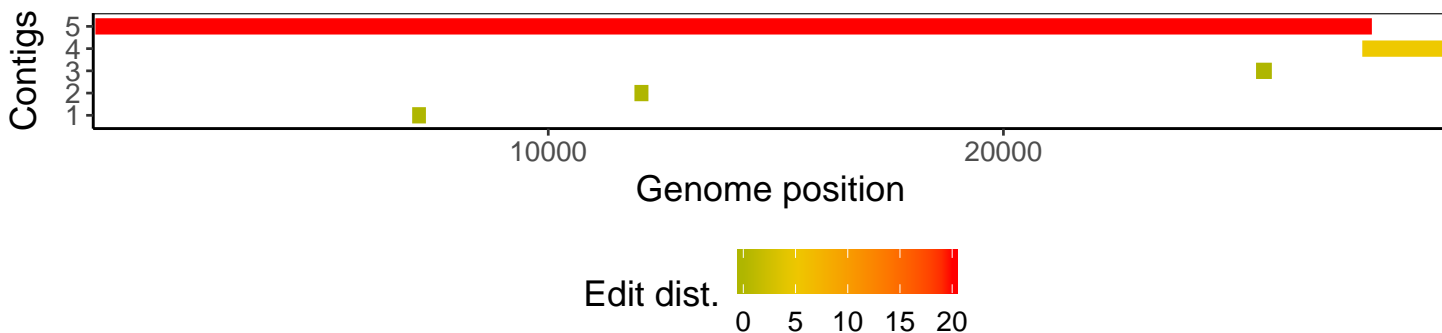
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according to variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



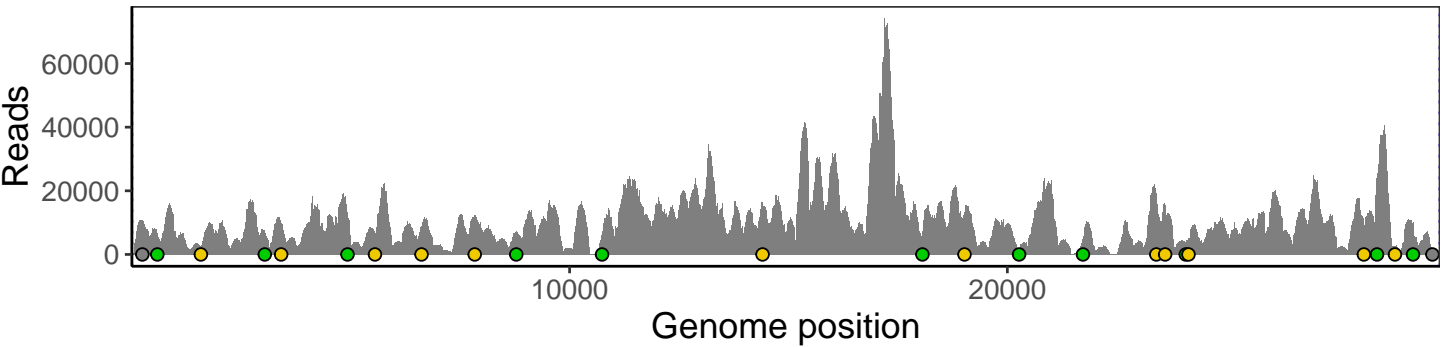
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



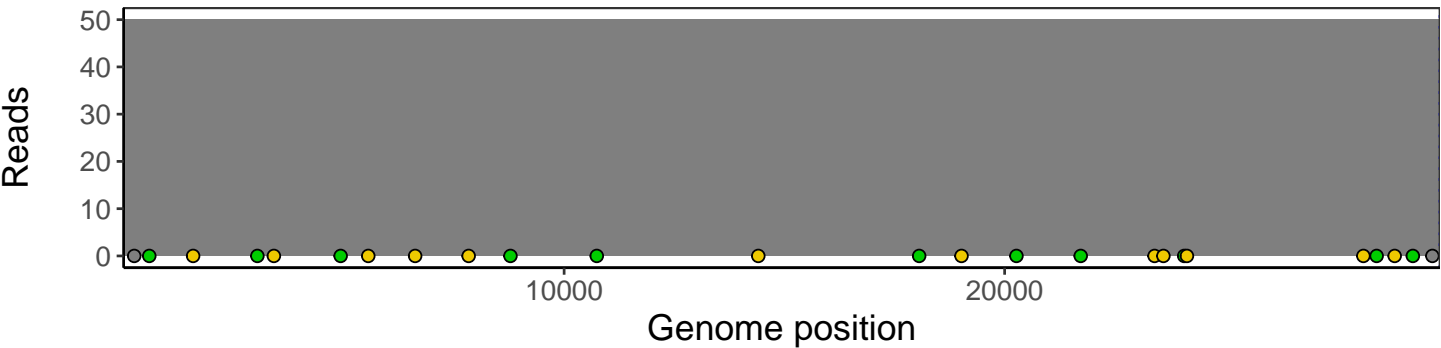
The longest five assembled contigs are shown below colored by their edit distance to the reference genome.



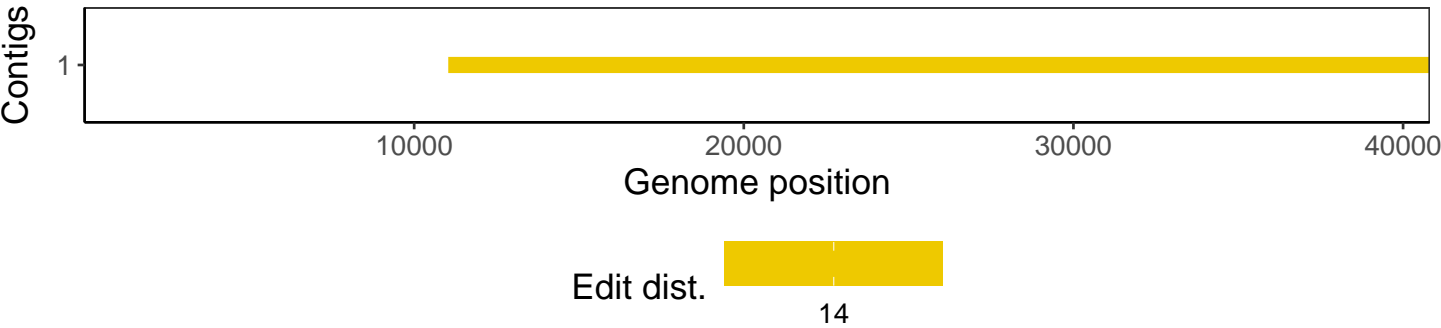
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



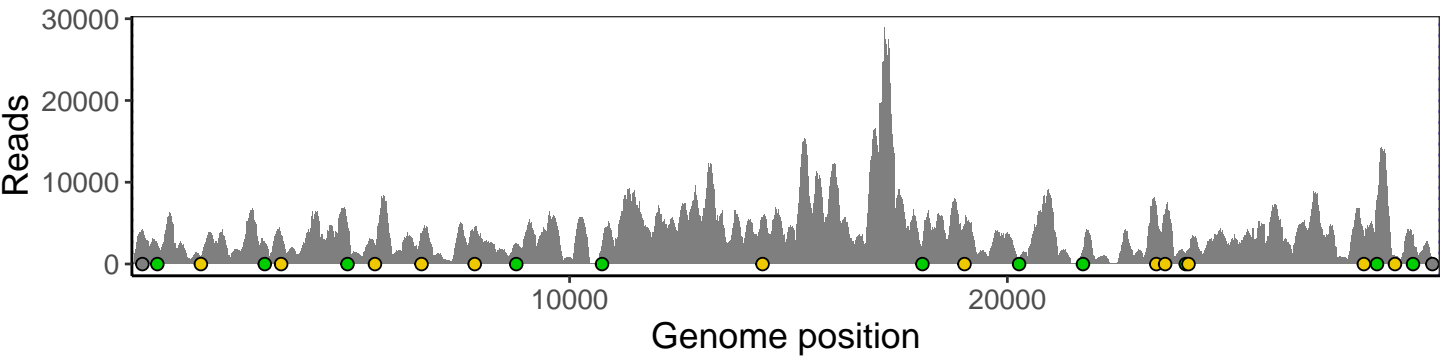
Excerpt from plot above focusing on reads coverage from 0 to 50 NT.



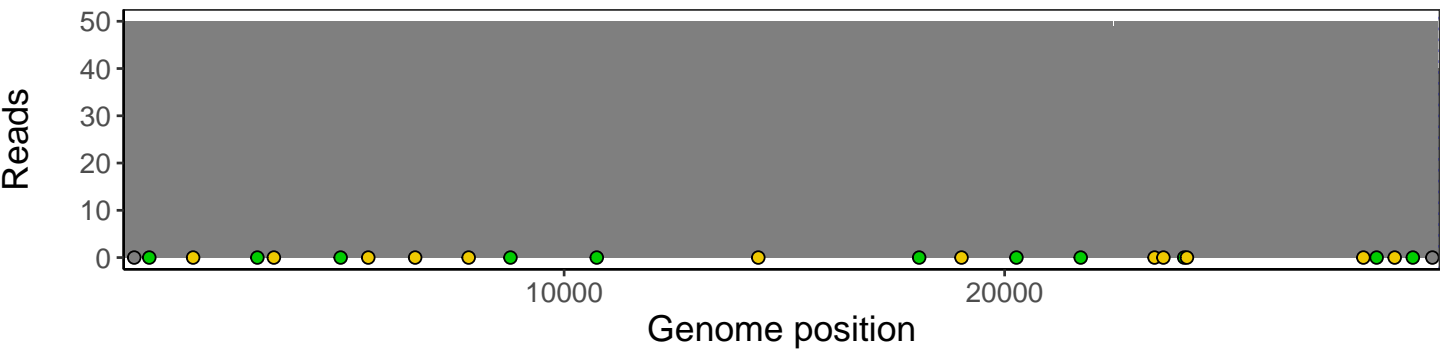
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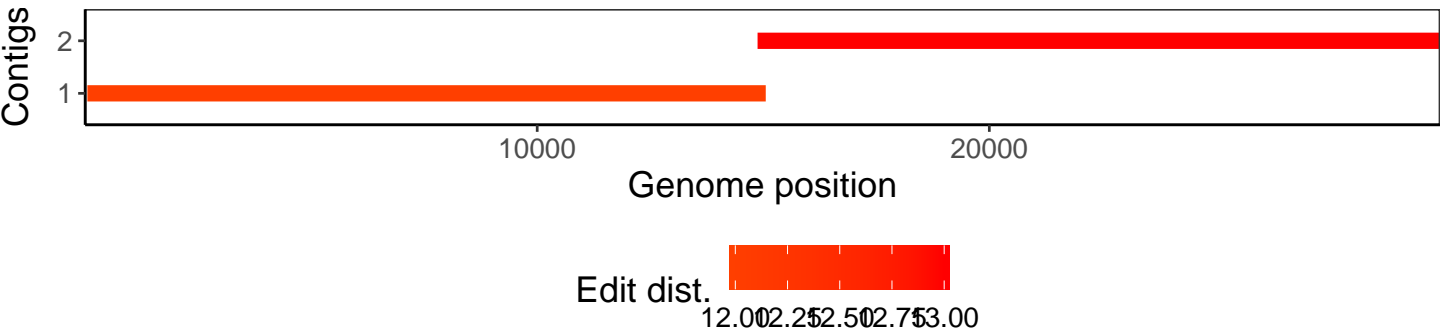
The plot below shows the number of reads covering each nucleotide position in the reference genome. Variants are shown as colored dots along the bottom of the plot and are color coded according by variant types: gray - transgenic, green - silent, gold - missense, red - nonsense, black - indel.



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Software environment

| Software/R package | Version |
|----------------------|---|
| R | 3.4.0 |
| bwa | 0.7.17-r1198-dirty |
| samtools | 1.10 Using htlib 1.10 |
| bcftools | 1.10.2-34-g1a12af0-dirty Using htlib 1.10.2-57-gf58a6f3 |
| pangolin | 2.3.8 |
| genbankr | 1.4.0 |
| optparse | 1.6.0 |
| forcats | 0.3.0 |
| stringr | 1.4.0 |
| dplyr | 0.8.1 |
| purrr | 0.2.5 |
| readr | 1.1.1 |
| tidyr | 0.8.1 |
| tibble | 2.1.2 |
| ggplot2 | 3.0.0 |
| tidyverse | 1.2.1 |
| ShortRead | 1.34.2 |
| GenomicAlignments | 1.12.2 |
| SummarizedExperiment | 1.6.5 |
| DelayedArray | 0.2.7 |
| matrixStats | 0.54.0 |
| Biobase | 2.36.2 |
| Rsamtools | 1.28.0 |
| GenomicRanges | 1.28.6 |
| GenomeInfoDb | 1.12.3 |
| Biostrings | 2.44.2 |
| XVector | 0.16.0 |
| IRanges | 2.10.5 |
| S4Vectors | 0.14.7 |
| BiocParallel | 1.10.1 |
| BiocGenerics | 0.22.1 |