Viral Beacon Statistics part 2

1 Stats on variants

Statistics are calculated using Query by region -by names/aliases and Filters.

I would keep separate this by sequencing technology (platform): Illumina vs Nanopore (for now only Illumina) and of course fastq vs consensus (for now only fastq)

- 1. Number of genomic positions with variants: 28754/Number of genomic positions: 29903 (96.15758%)
- 2. Frequency of variants per position: fig 1
- 3. Number of variants per position: (3: 5769; 2: 11021; 1:11964) fig 1
- 4. Number of variants in database: 51313
- 5. Graph: Number of variants, split by options
- option Split by variant frequency (groups) fig 2
- option Split by variant type field: SNP, indels (only SNPs) Mau to check pipeline? fig 3
- option Split by genomic region: coding: all with genomic region=CODING, non-coding: the rest fig (further UTRs, intergenic?) fig 3
- option Split by molecular consequence (grouped: SYN: SILENT, NON-SYN: MISSENSE+NONSENSE, NONCODING: the rest) and further in all classes as in fig 3
- option See Per region Statistics: Distribution in genomic regions. Upon click on genomic region graph, expand to see per number of variants distribution in genomic regions: select $\underline{\text{NON-CODING}}$, and within $\underline{\text{CODING}}$ show options: $\underline{\text{gene}}$, $\underline{\text{cds/mature peptide}}$. This will show individual components of each class fig 4 and 5.
- 6. Number of positions with aminoacid substitutions in database/ Number of coding positions
- 7. Number of variants producing unique aminoacid substitutions in database: 18308
- option Filter upon cds/mature peptide. Number of variants producing aminoacid substitutions (aminoacid change) (eg. "G507C") are aggregated in mature protein region

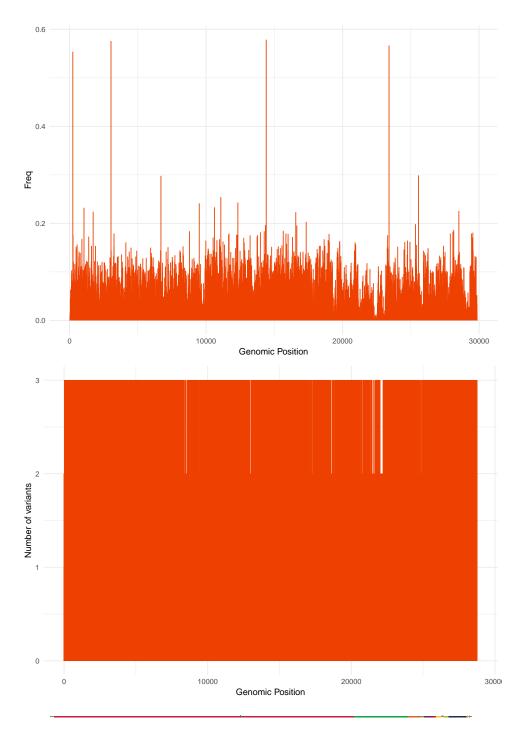


Figure 1: Needle plot: Top: Frequency of variants per position. Bottom: Number of alternates per position

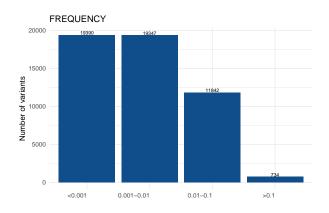


Figure 2: Number of variants per frequency group

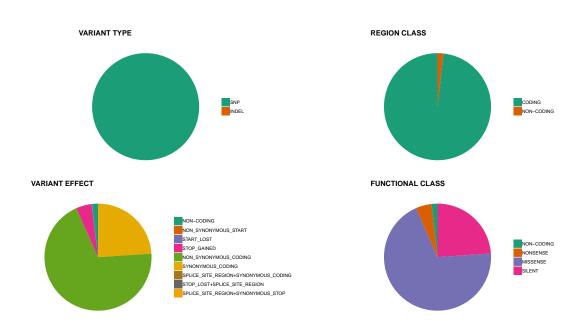


Figure 3: Number of variants, split by variant type, region class, variant effect and mol consequence

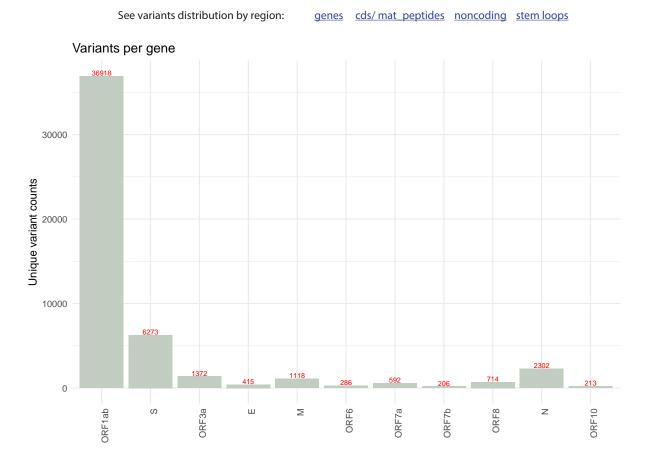


Figure 4: Number of variants in coding regions of SARS-CoV2, shown per genes, shown directly on clicking CODING area of genomic region graph or in option "See variants distribution by region" genes.



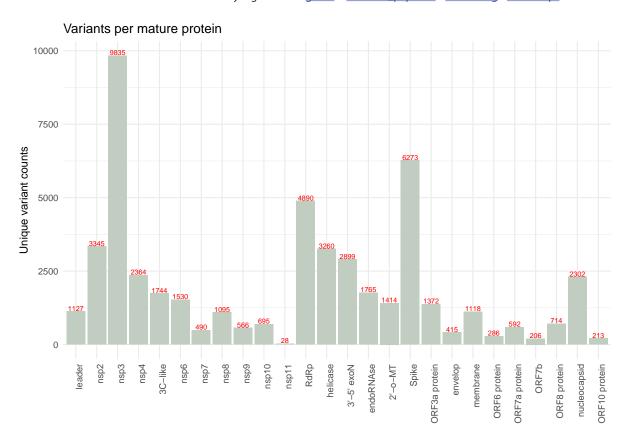


Figure 5: Number of variants in coding regions of SARS-CoV2, shown per mature proteins, shown on clicking cds/mature peptide option within CODING area of genomic region graph (which will show directly upon clicking the variants per gene graph 4 or in option "See variants distribution by region" cds/mat_peptides. Additional options within this graph: split by syn/non syn (or default in stack bar)

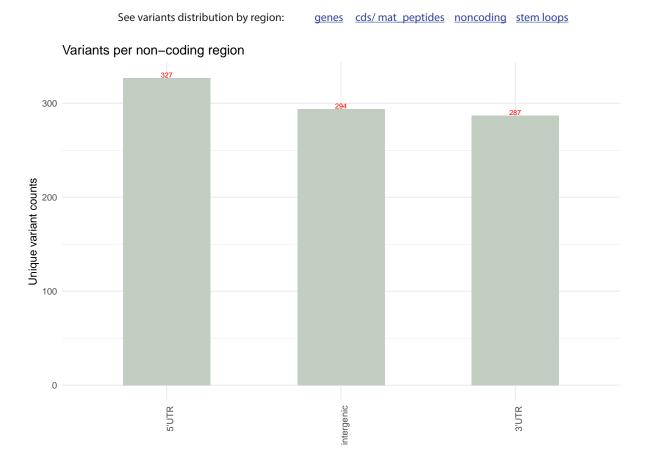


Figure 6: Number of variants in non-coding regions of SARS-CoV2, shown per region, shown on clicking $\underline{\text{NON-CODING}}$ area of genomic region graph or in option "See variants distribution by region" $\underline{\text{noncoding}}$.

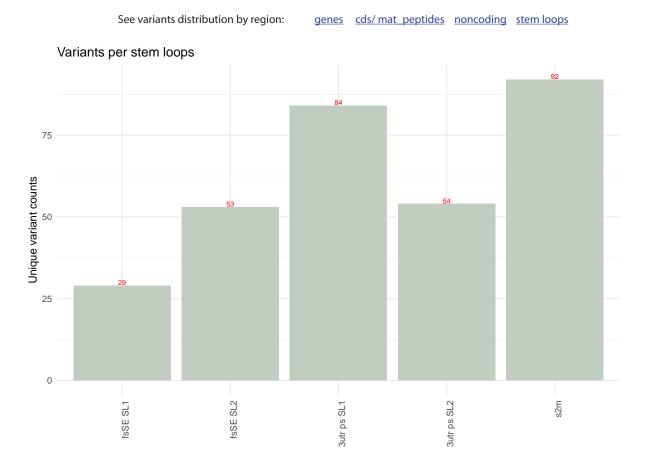


Figure 7: Number of variants in stem loops regions of SARS-CoV2, shown per region, shown on clicking in option "See variants distribution by region" stem loops.