

SARS-CoV-2 Sequencing Update 26 August 2022



Supported by the DSI and the SA MRC

Msomi N, Mlisana K, et al. Lancet Microbe 2020

The genomic data presented here are based on South African SARS-CoV-2 sequence data downloaded from GISAID (www.gisaid.org) on 26 August 2022 at 15h13



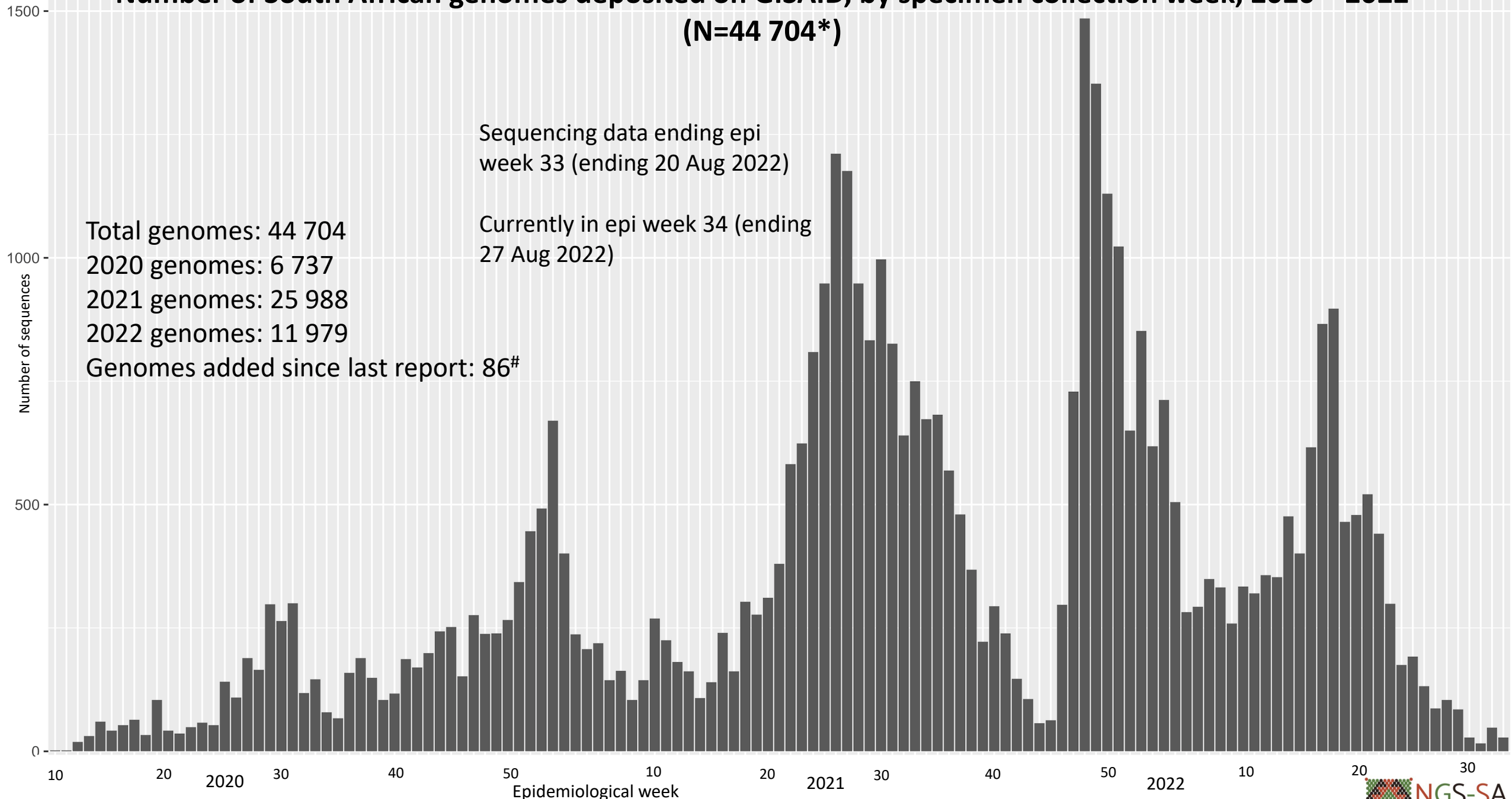
Data license: <https://www.gisaid.org/registration/terms-of-use/>

Elbe, S., and Buckland-Merrett, G. (2017) Data, disease and diplomacy: GISAID's innovative contribution to global health. *Global Challenges*, 1:33-46. DOI: 10.1002/gch2.1018 PMCID: 31565258

Shu, Y., McCauley, J. (2017) GISAID: Global initiative on sharing all influenza data – from vision to reality. *EuroSurveillance*, 22(13) DOI: 10.2807/1560-7917.ES.2017.22.13.30494 PMCID: PMC5388101

Number of South African genomes deposited on GISAID, by specimen collection week, 2020 – 2022

(N=44 704*)

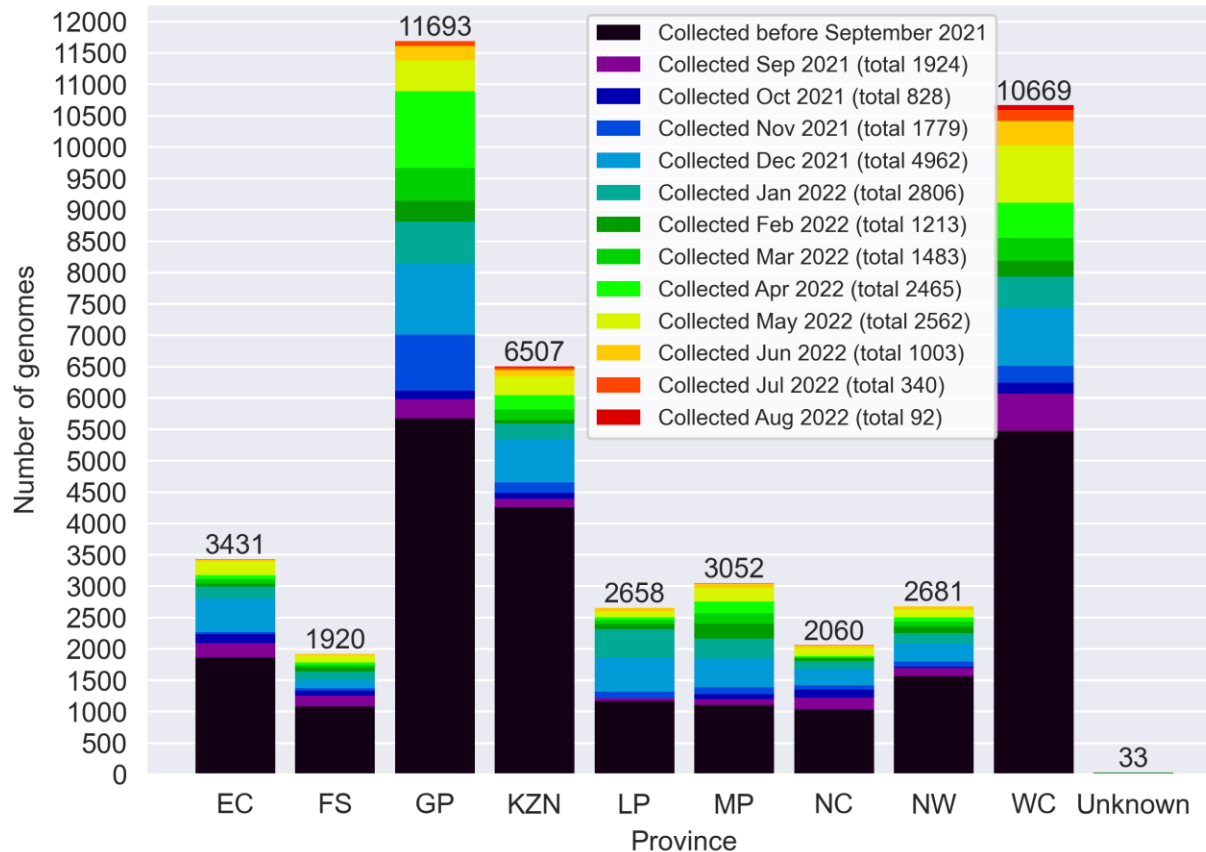


*This represents the cleaned, de-duplicated dataset of unique **National and Pneumonia Surveillance** sequences. This dataset will be used for all further figures.

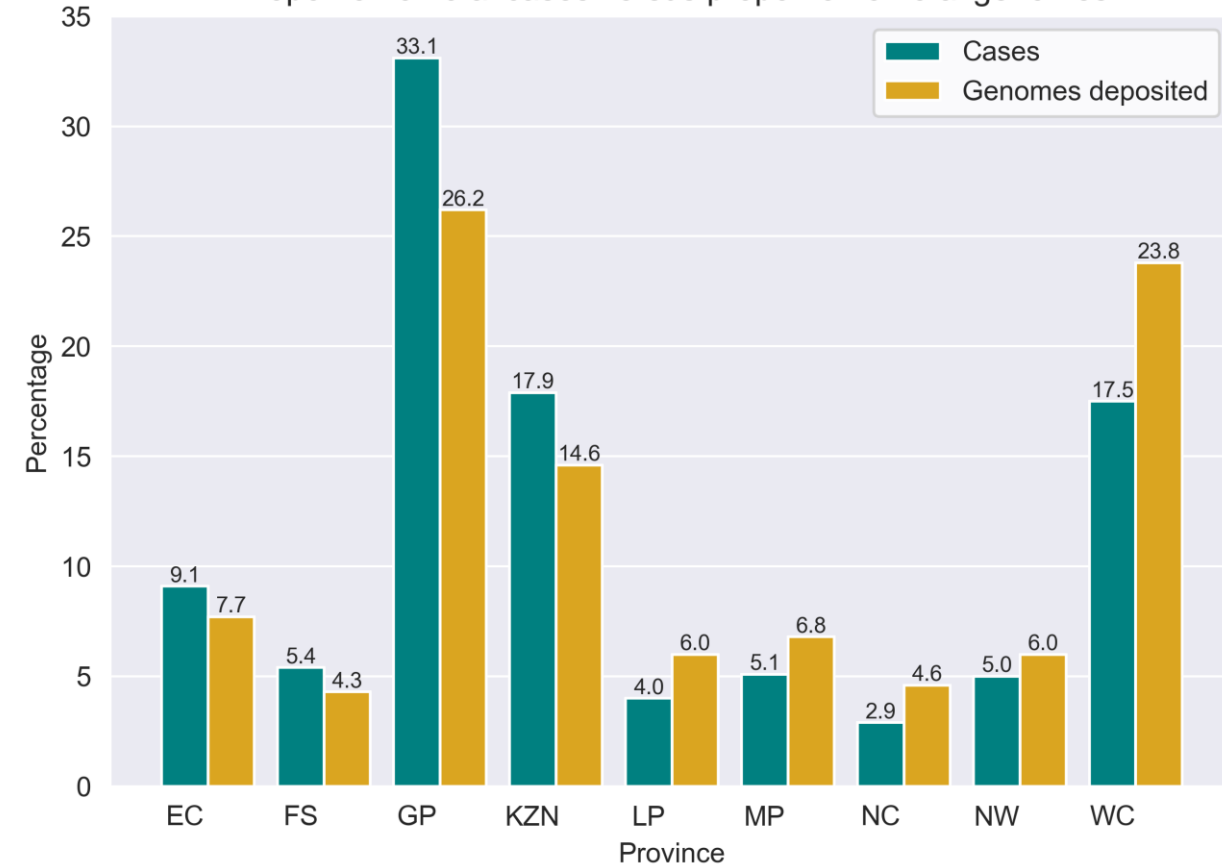
[#]3 longitudinal samples from the same individual removed since previous report

GISAID genomes vs total cases, 2020 – 2022 (N=44 704)

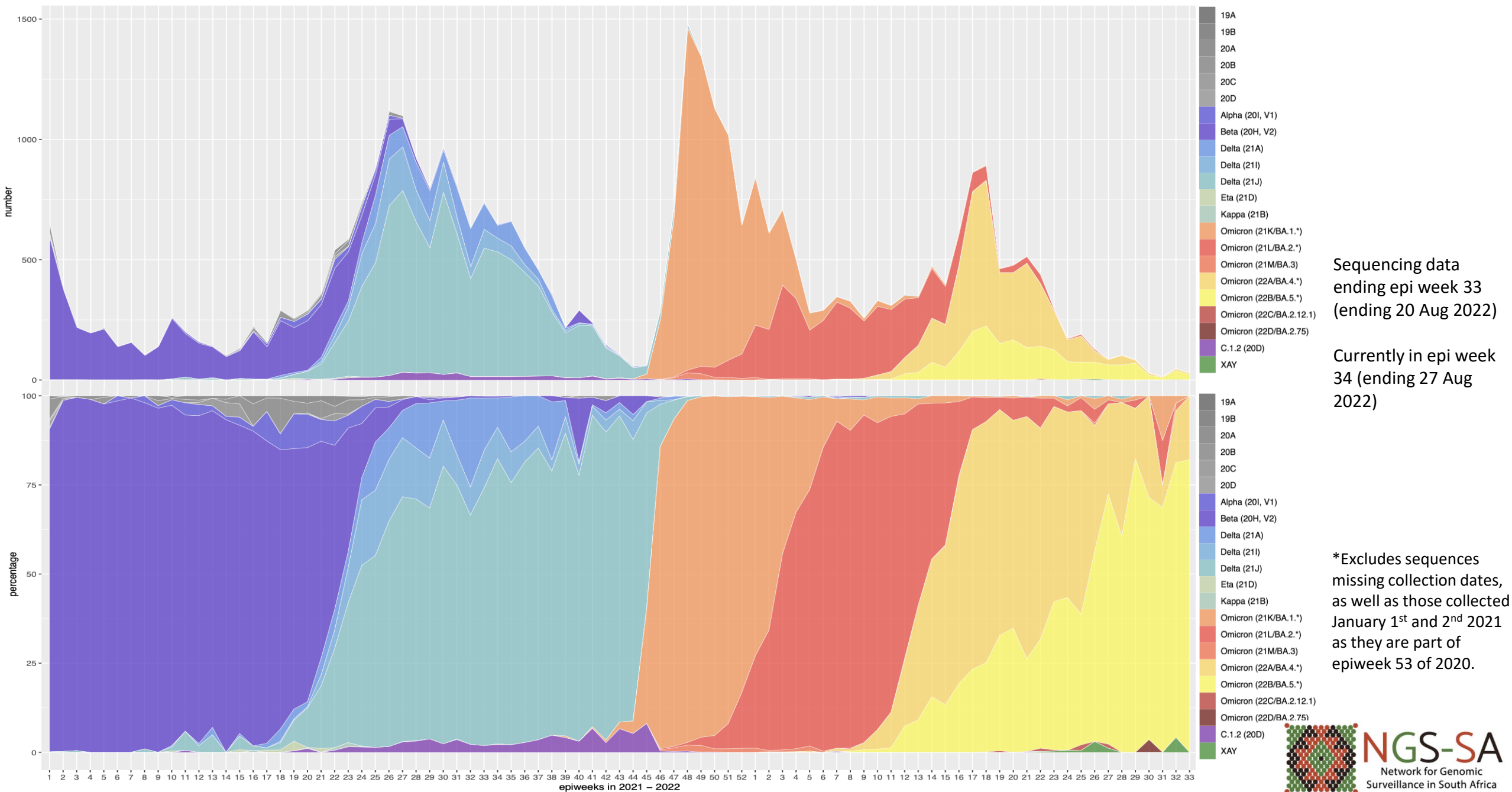
Provincial breakdown of genomes deposited into GISAID



Proportion of total cases versus proportion of total genomes

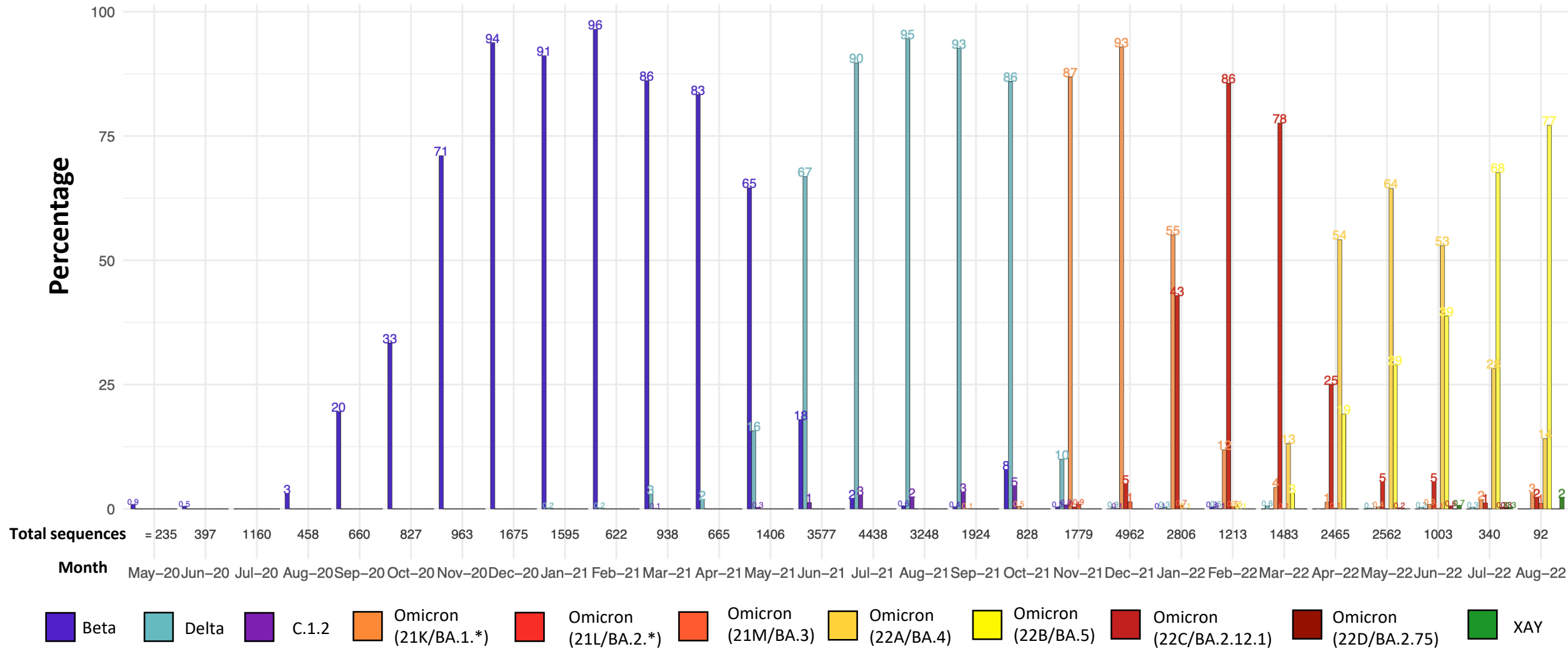


Number and percentage of clades by epiweek in South Africa, 2021 – 2022 (37 797*)



Detection Rates: Beta, Delta, C.1.2 and Omicron

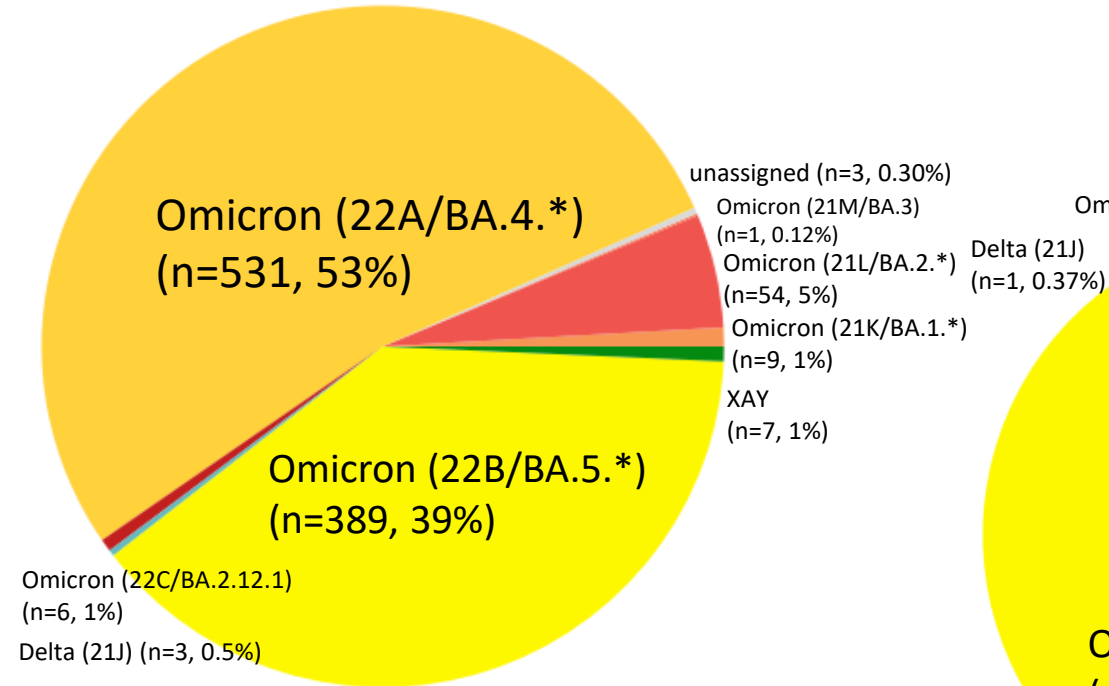
Detection rates of variants being monitored in South Africa



*Bars represent percentage prevalence of variant for the month; total number sequences collected for the month are given below the bar

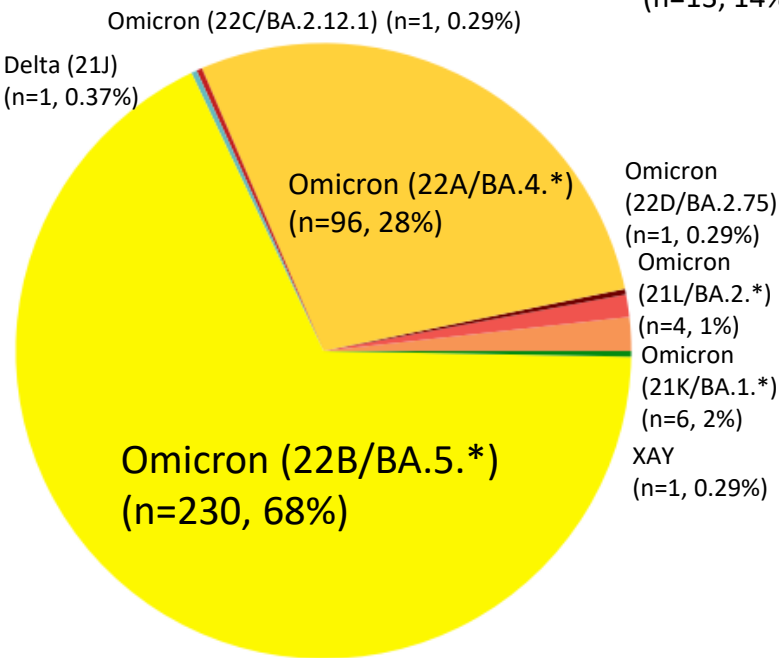
Prevalence of Variants of Concern (VOC) and Variants of Interest (VOI) in June – August 2022

June (N=1003)



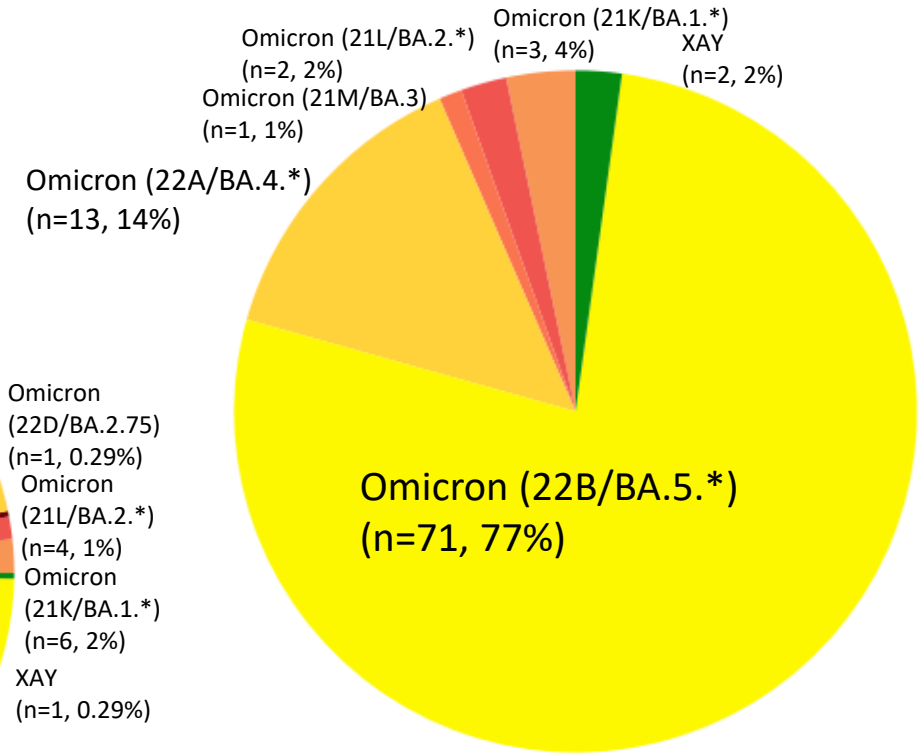
Total Omicron in June: 990 (98.7%)

July (N=340)



Total Omicron in July: 337 (99.1%)

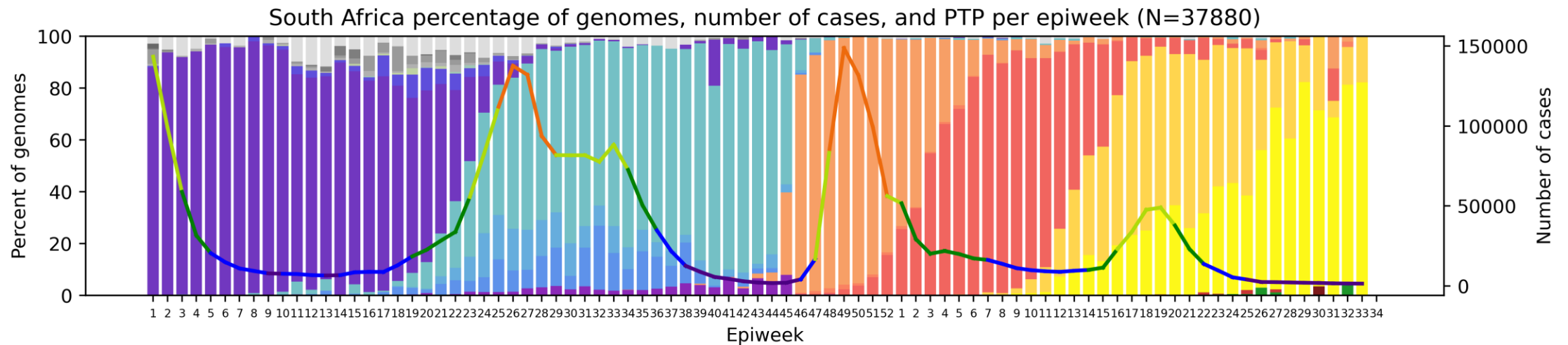
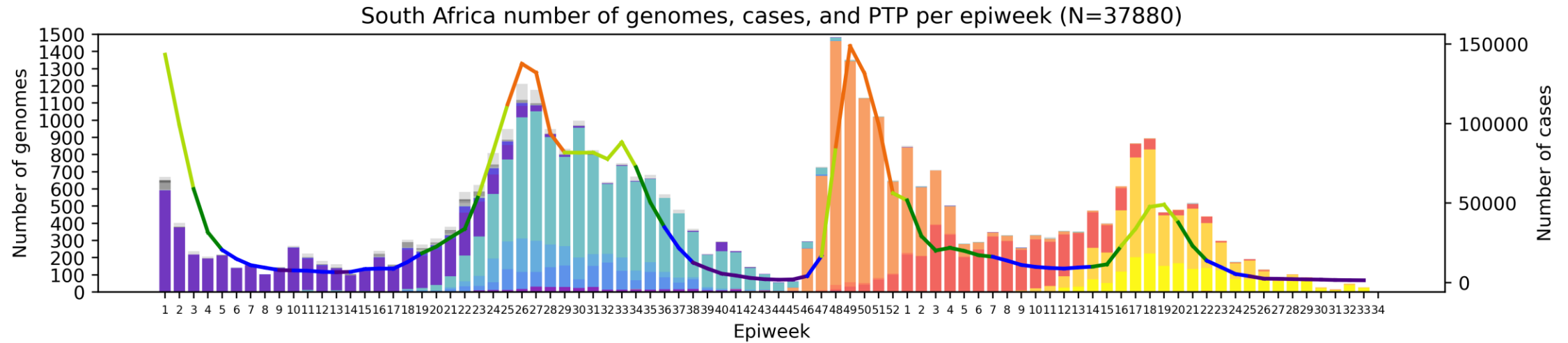
August (N=92)



Total Omicron in August: 90 (97.8%)



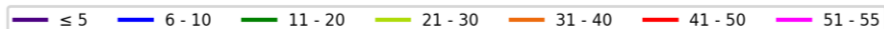
South Africa, 2021-2022, n = 37 880*



Clade key (bar graph)



Weekly percentage testing positive key (line graph)

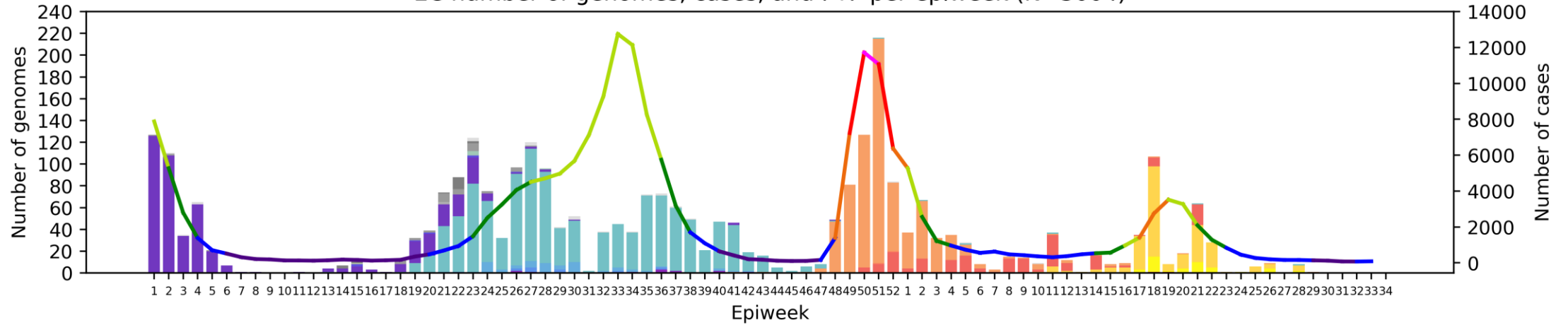


*Excludes sequences missing collection dates, as well as those collected January 1st and 2nd 2021 as they are part of epiweek 53 of 2020.

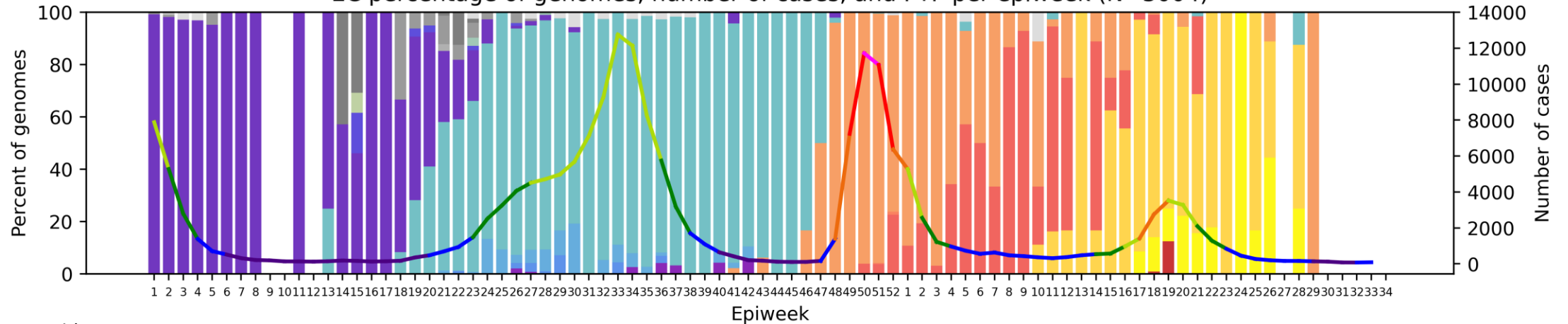
Eastern Cape Province, 2021-2022, n = 3004

Genomes added since last report: 0*

EC number of genomes, cases, and PTP per epiweek (N=3004)



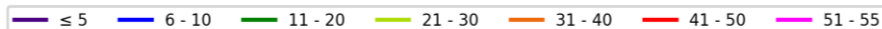
EC percentage of genomes, number of cases, and PTP per epiweek (N=3004)



Clade key (bar graph)

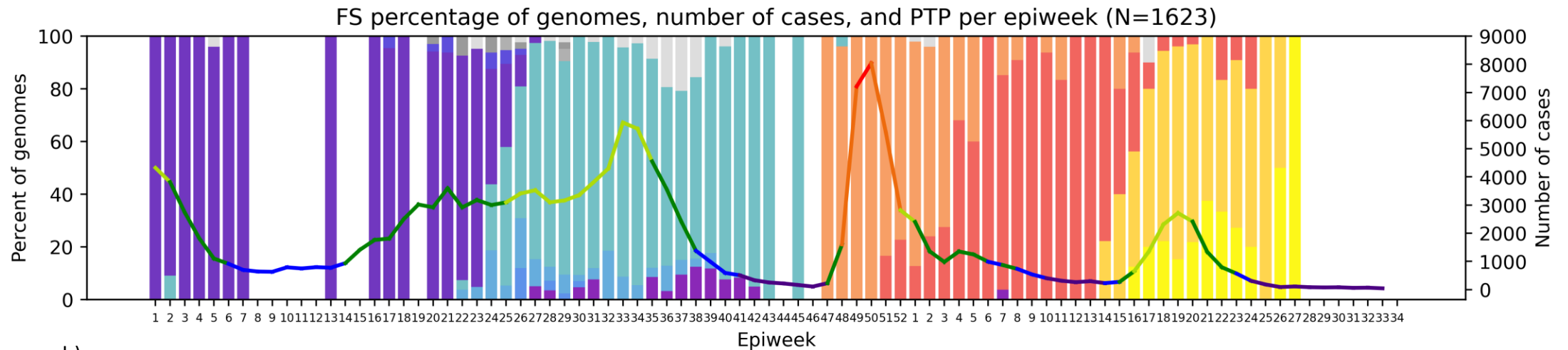
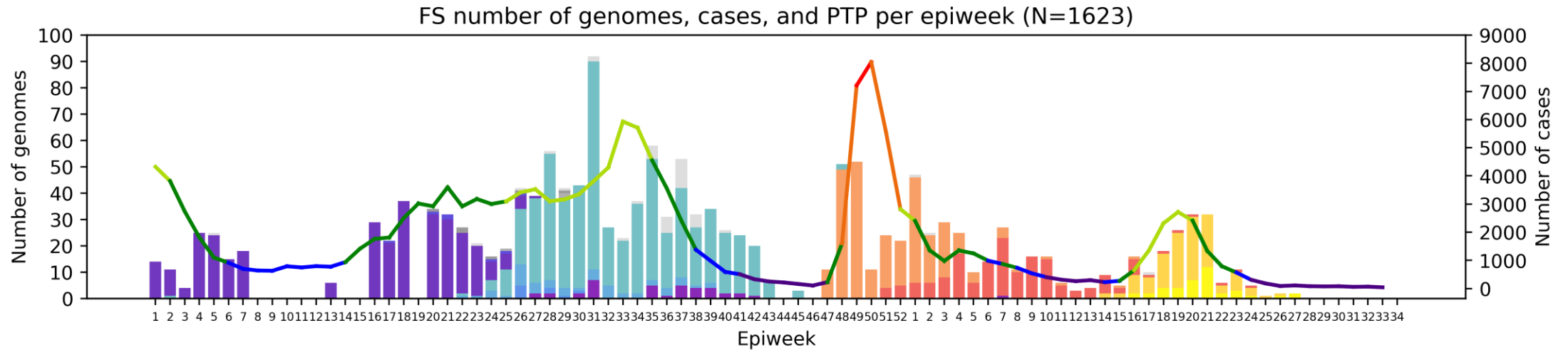


Weekly percentage testing positive key (line graph)



Free State Province, 2021-2022, n = 1623

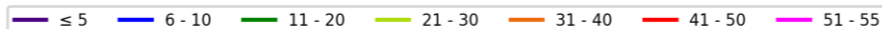
Genomes added since last report: 0*



Clade key (bar graph)



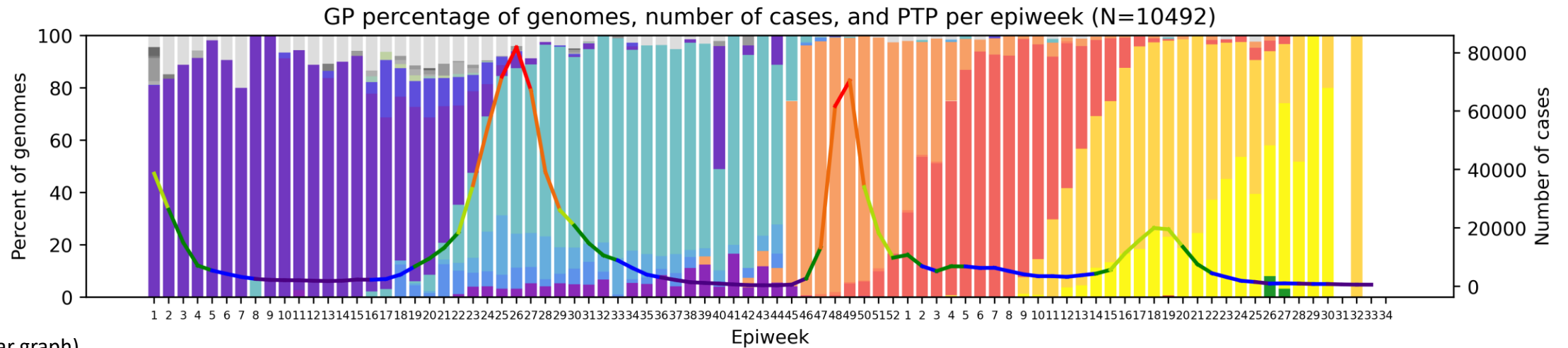
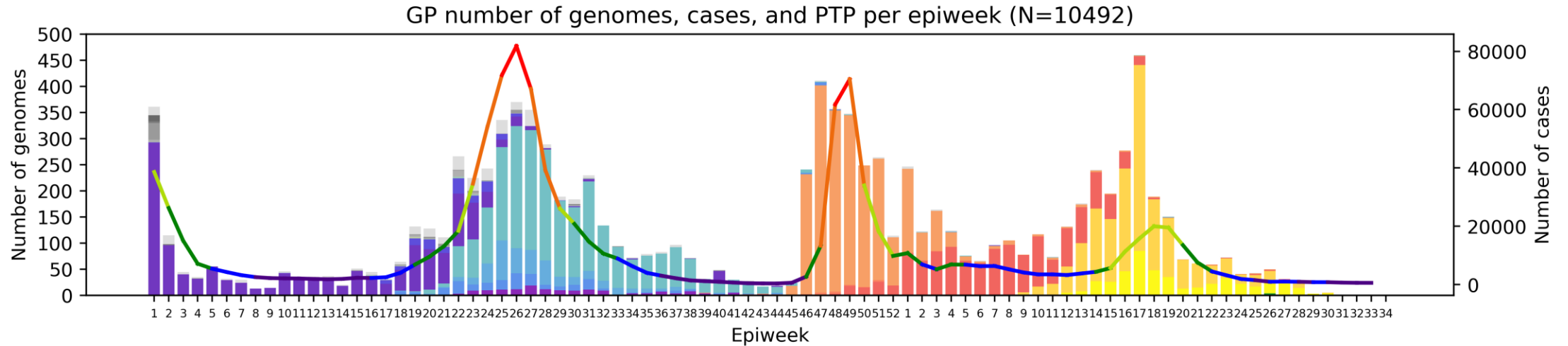
Weekly percentage testing positive key (line graph)



*May include genomes from 2020 which are not pictured here and are not included in the slide total.

Gauteng Province, 2021-2022, n = 10 492

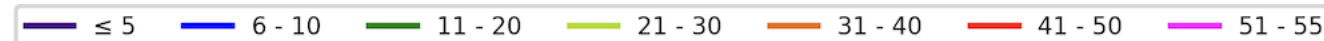
Genomes added since last report: 1*



Clade key (bar graph)



Weekly percentage testing positive key (line graph)

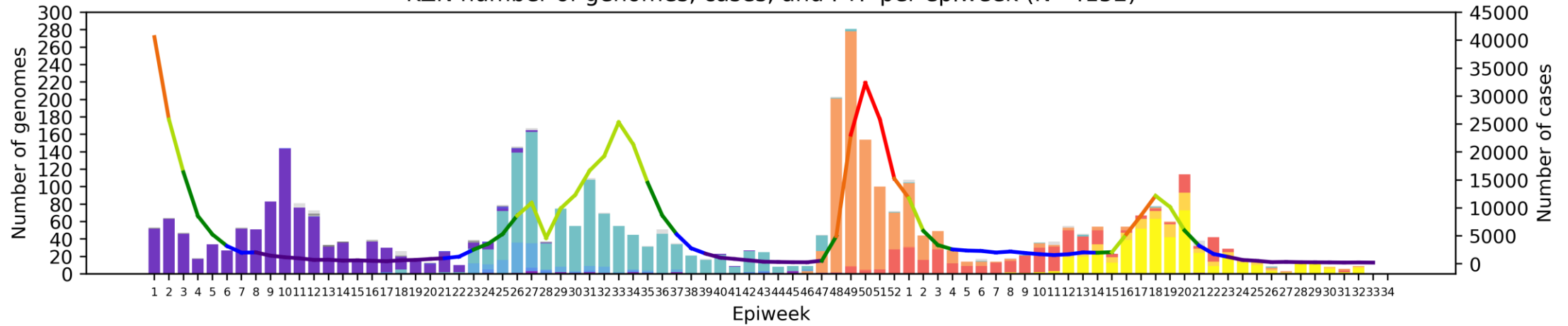


*May include genomes from 2020 which are not pictured here and are not included in the slide total.

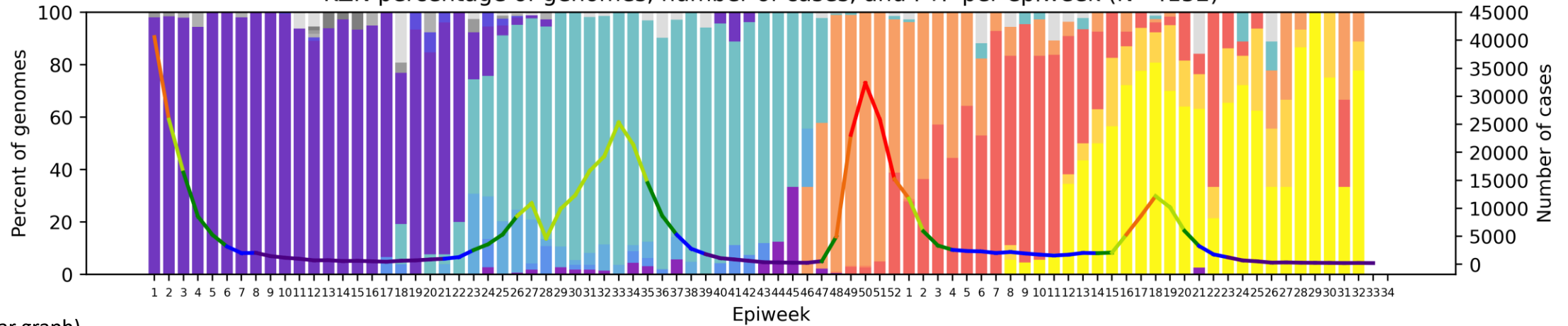
KwaZulu-Natal Province, 2021-2022, n = 4152

Genomes added since last report: 10*

KZN number of genomes, cases, and PTP per epiweek (N=4152)



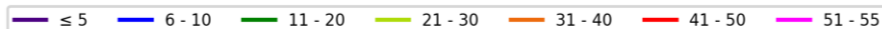
KZN percentage of genomes, number of cases, and PTP per epiweek (N=4152)



Clade key (bar graph)

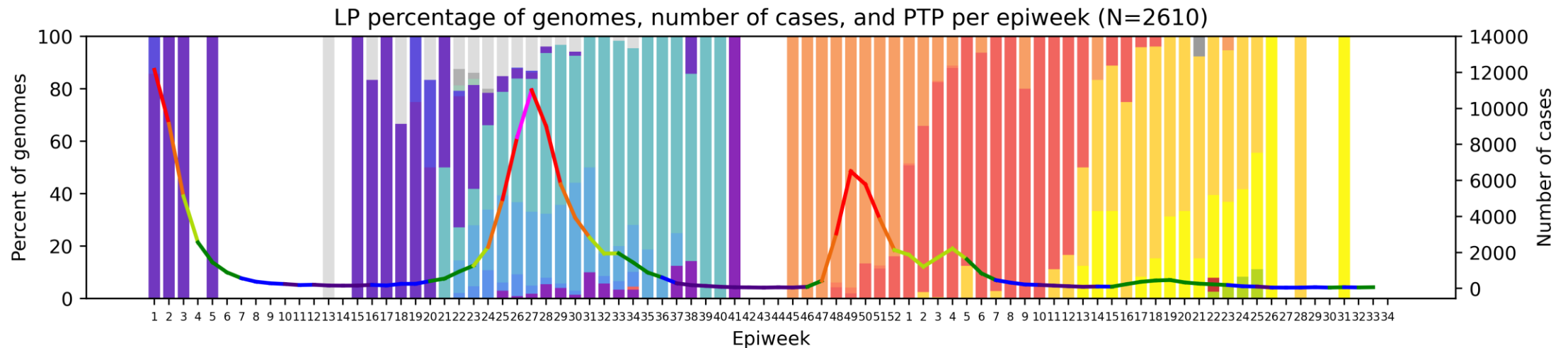
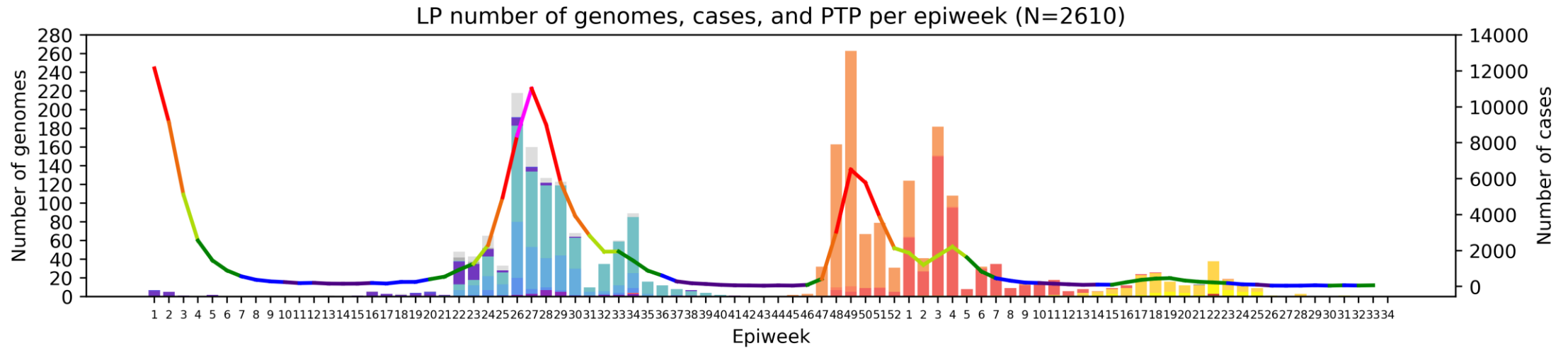


Weekly percentage testing positive key (line graph)



Limpopo Province, 2021-2022, n = 2610

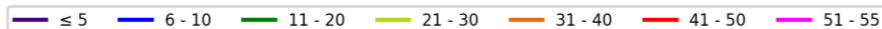
Genomes added since last report: 0*



Clade key (bar graph)



Weekly percentage testing positive key (line graph)

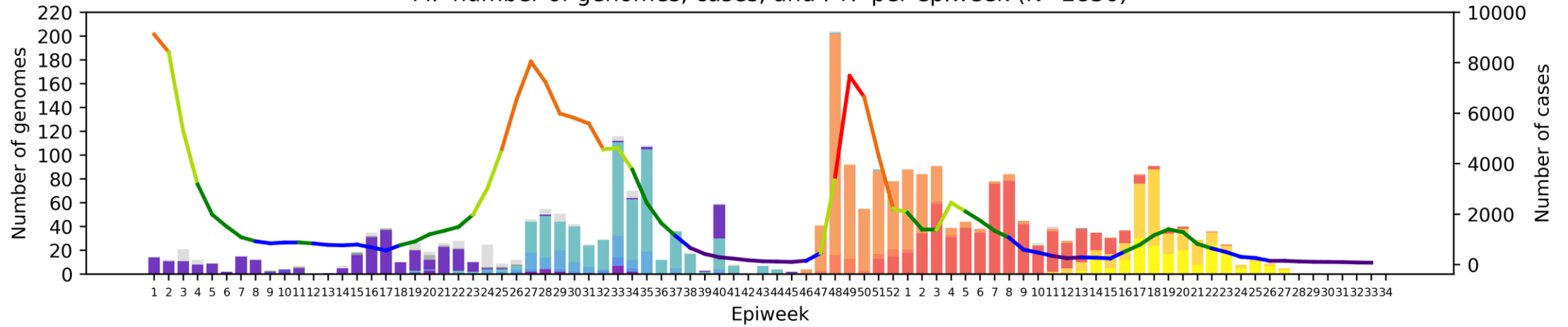


*May include genomes from 2020 which are not pictured here and are not included in the slide total.

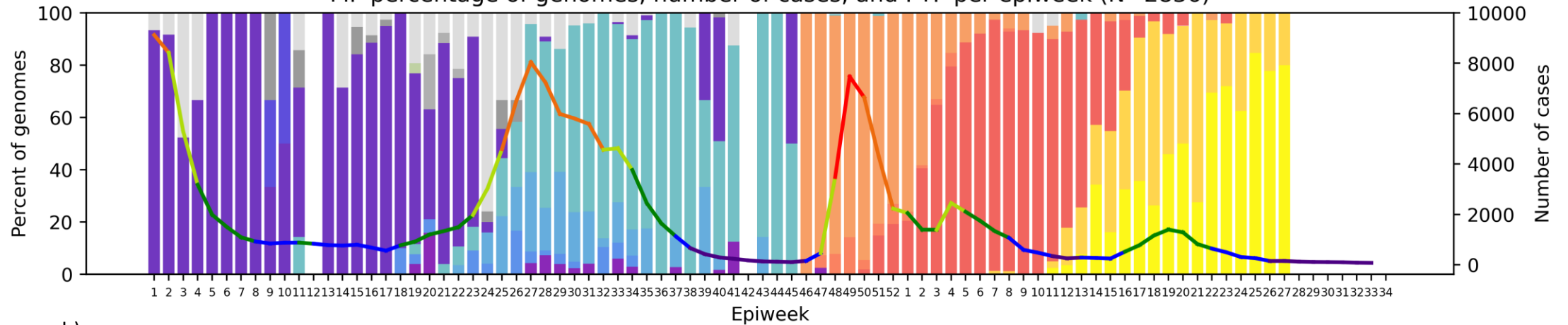
Mpumalanga Province, 2021-2022, n = 2836

Genomes added since last report: 0*

MP number of genomes, cases, and PTP per epiweek (N=2836)



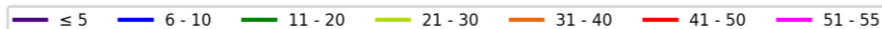
MP percentage of genomes, number of cases, and PTP per epiweek (N=2836)



Clade key (bar graph)

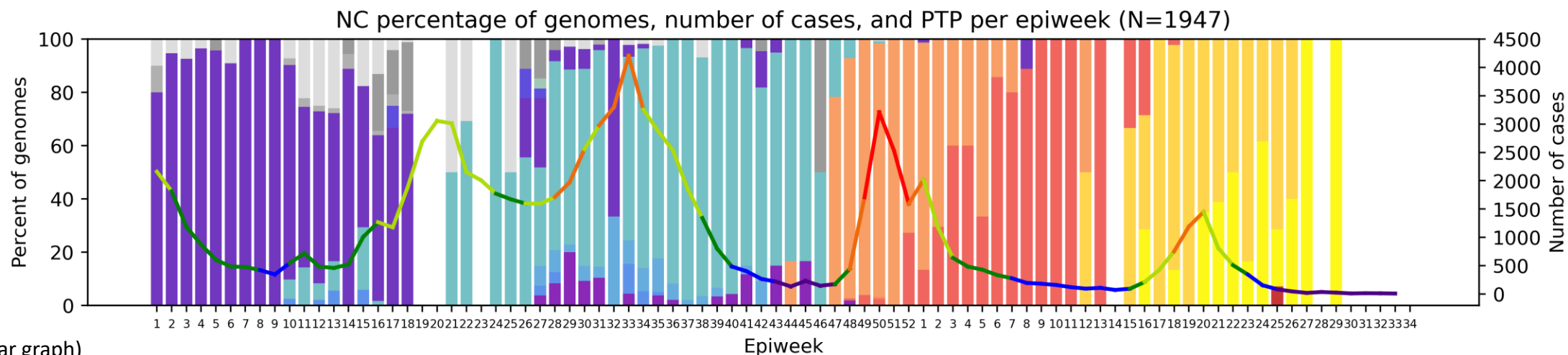
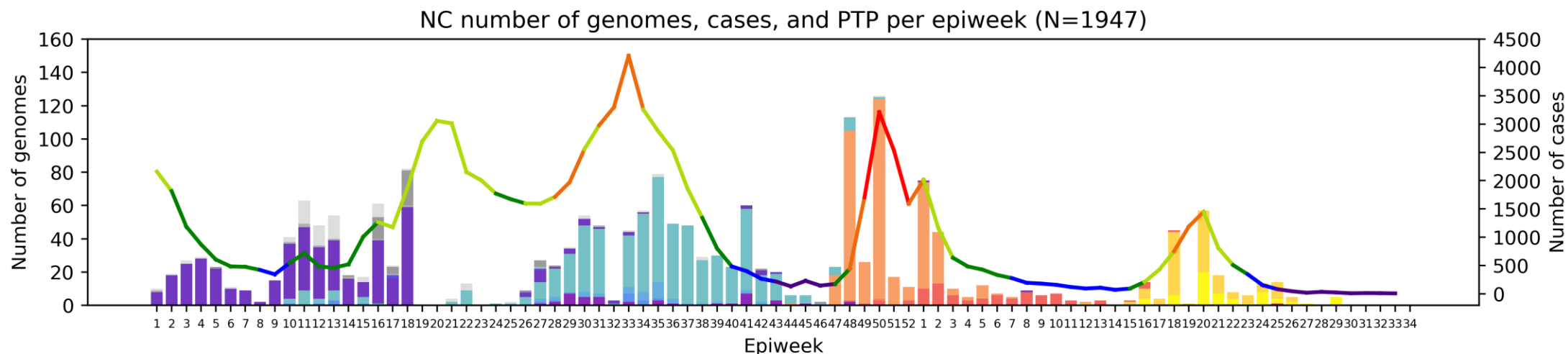


Weekly percentage testing positive key (line graph)



Northern Cape Province, 2021-2022, n = 1947

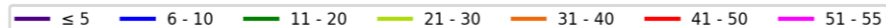
Genomes added since last report: 0*



Clade key (bar graph)



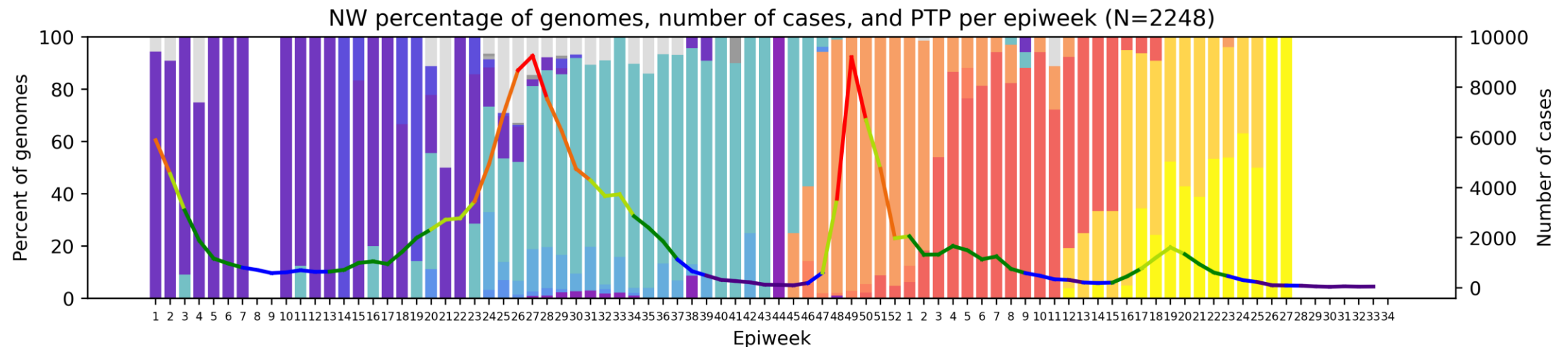
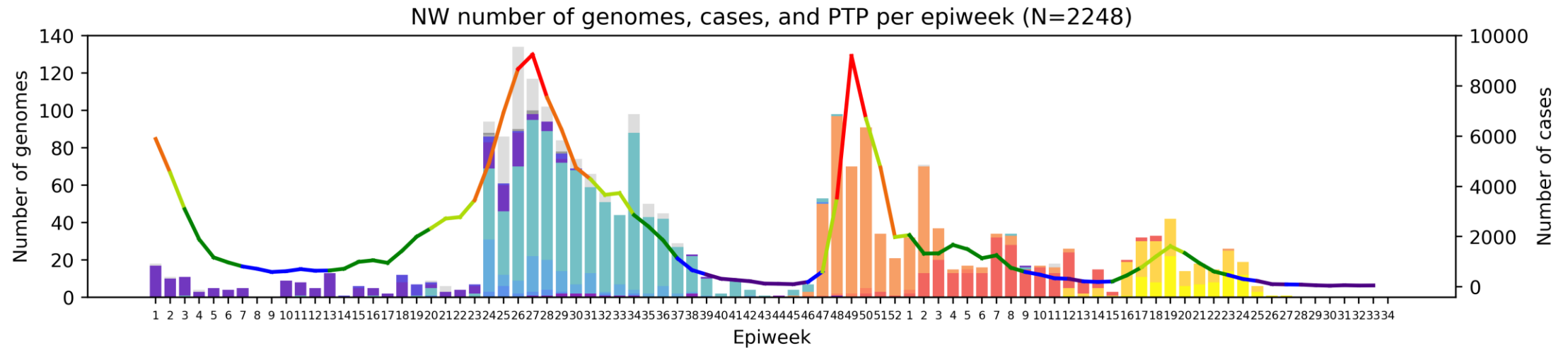
Weekly percentage testing positive key (line graph)



*May include genomes from 2020 which are not pictured here and are not included in the slide total.

North West Province, 2021-2022, n = 2248

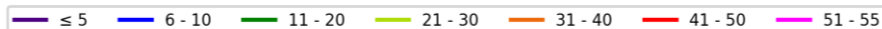
Genomes added since last report: 0*



Clade key (bar graph)



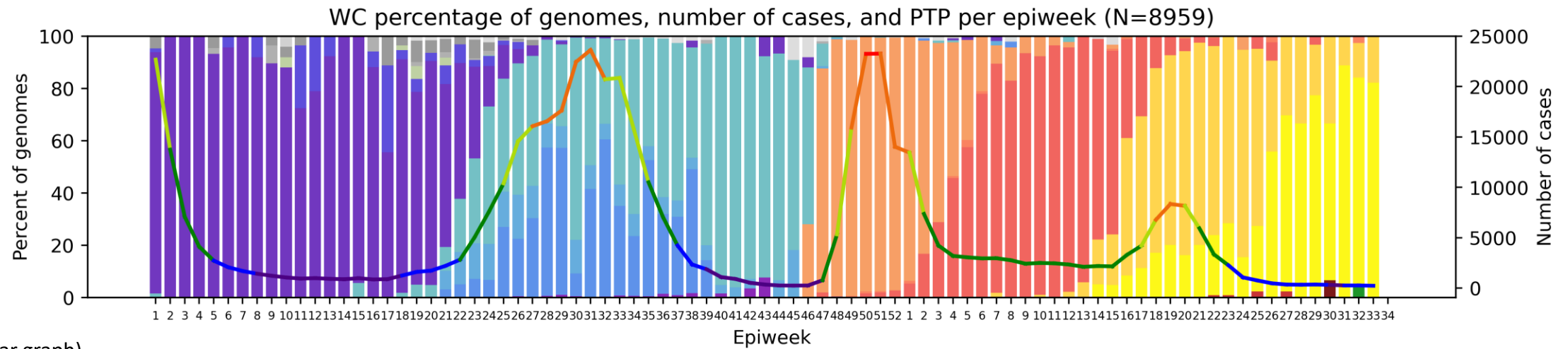
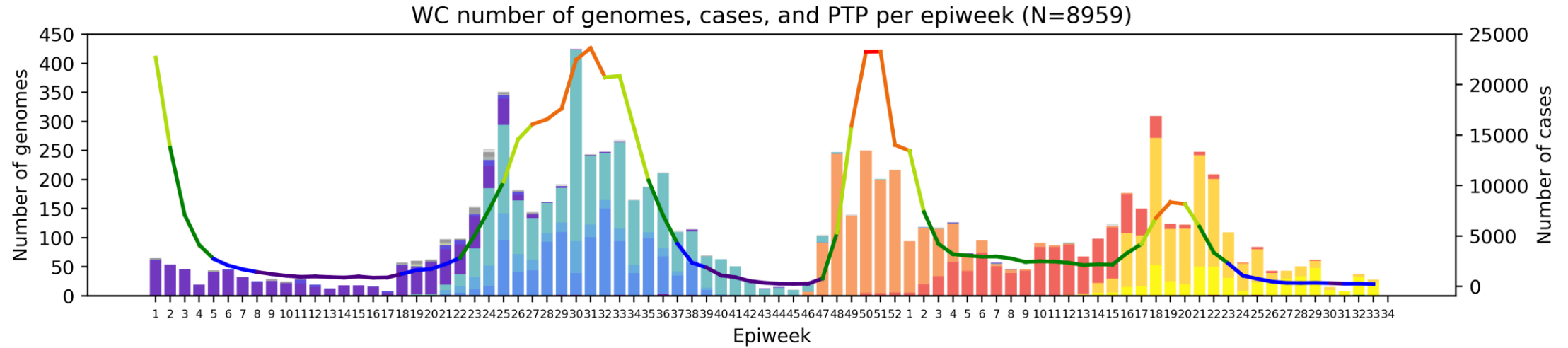
Weekly percentage testing positive key (line graph)



*May include genomes from 2020 which are not pictured here and are not included in the slide total.

Western Cape Province, 2021-2022, n = 8959

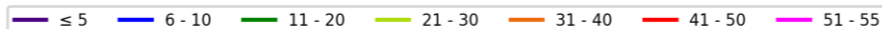
Genomes added since last report: 75*



Clade key (bar graph)



Weekly percentage testing positive key (line graph)



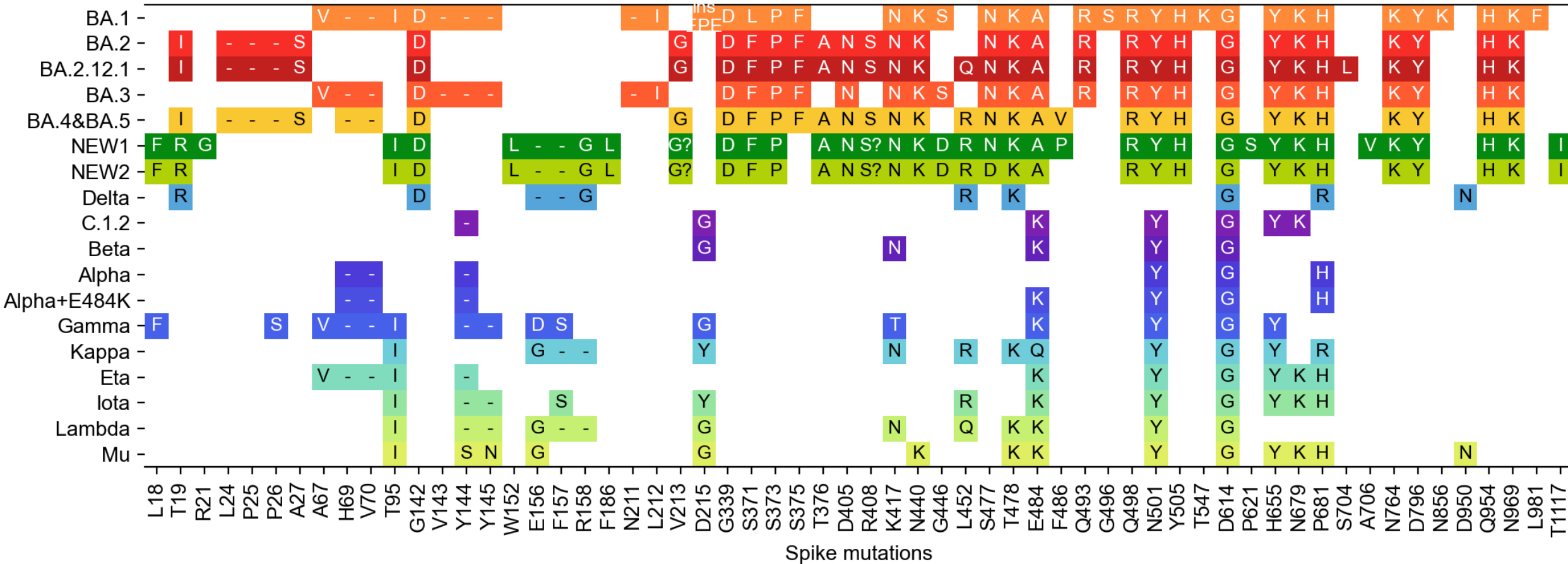
*May include genomes from 2020 which are not pictured here and are not included in the slide total.

Summary

- **Sequencing update**
 - All provinces have sequences for July.
 - August sequences are from KZN, GP, LP and WC.
 - Omicron dominated in June (98.7%), July (99.1%) and August (97.8%). BA.4 and BA.5 together were dominant in June, July and August.
- **N=11 sequences with novel mutational profile**
 - N=2 new sequences have been detected, collected 8 Aug 2022 in the Western Cape (first detection in the Western Cape)
 - Sequences are currently designated XAY but this is likely to change if more sequences are found¹.
- **Variant of Concern Omicron in South Africa**
 - Dominates 2022 sequencing data at >98% of genomes.
 - While BA.1 was the predominant lineage in January (55%), BA.2 dominated in February (86%) and March (78%).
 - Omicron lineages BA.4 and BA.5 increased in prevalence in March (16%), and together are dominant in April (73%), May (93%), June (92%), July (96%), and August (91%).
 - BA.2.12.1 was detected in South Africa at low prevalence in May, June and July (<1%)
 - BA.2.75 was detected in South Africa at low prevalence in July (<1%)
- Low frequency of previously circulating variants such as Delta still detected in recent data.

¹ <https://github.com/cov-lineages/pango-designation/issues/844#issuecomment-1227532073>

Spike protein mutation* profile of Variants of Interest and Concern

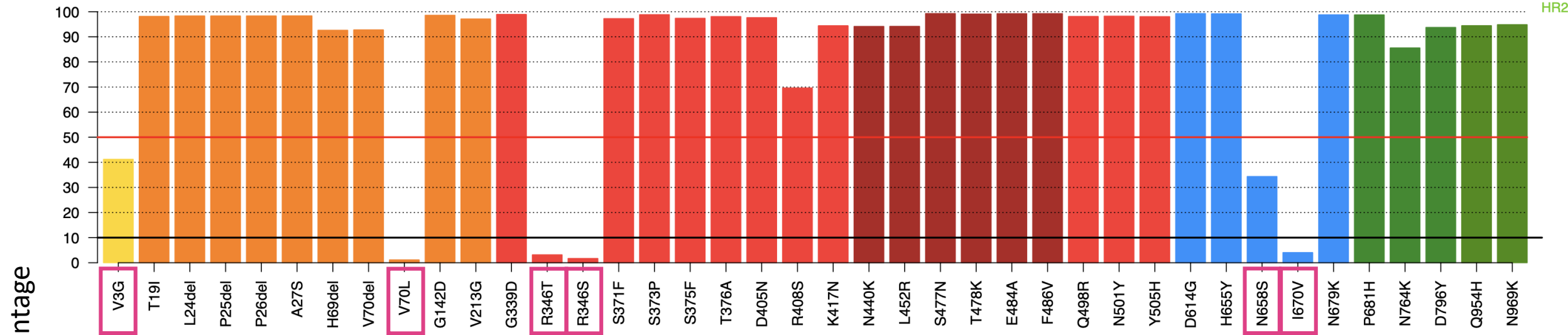


- Multiple changes within the two immunogenic regions in S1 (NTD and RBD)
 - Including a three amino acid insertion
- Accumulation of mutations surrounding the furin cleavage site
 - Including combination of N679K and P681H
- Effect of most spike S2 subunit changes have not been defined, but may be linked to immune escape

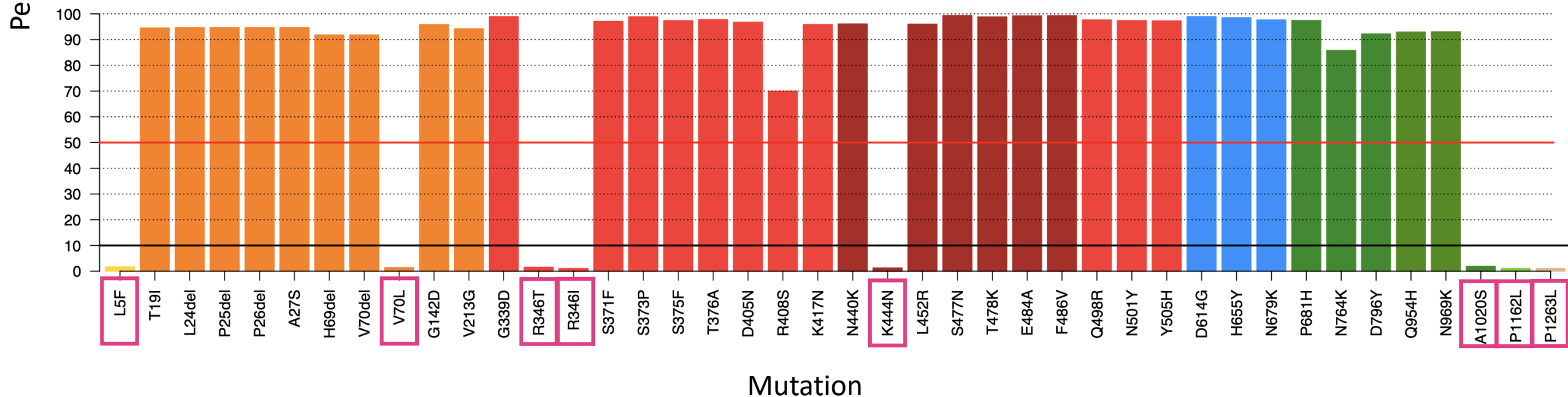
*Only mutations present in Omicron, Delta, or the new SGTP sequences are picture

BA.4 and BA.5 spike mutations

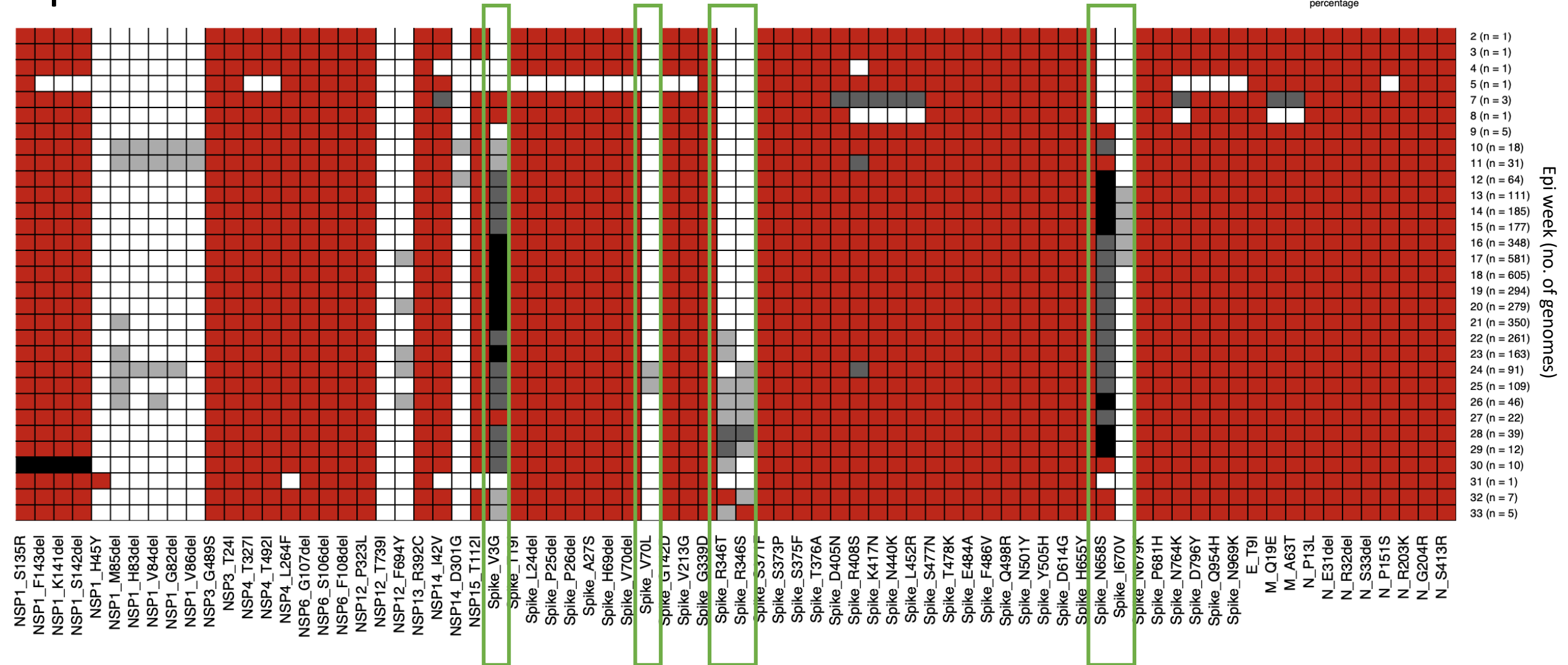
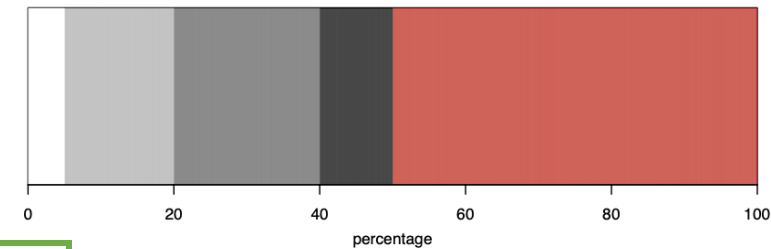
SP
NTD
RBD
RBM
S1
S2
HR1
HR2



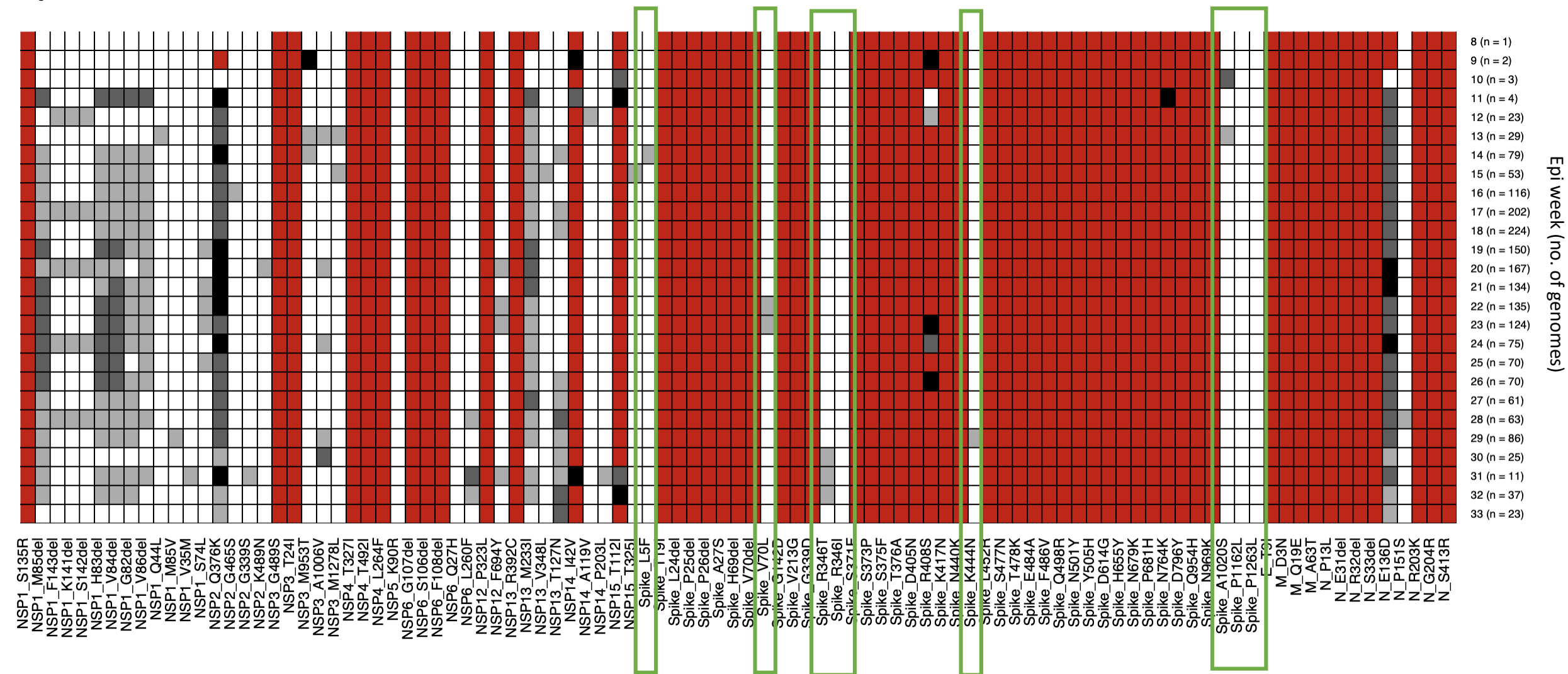
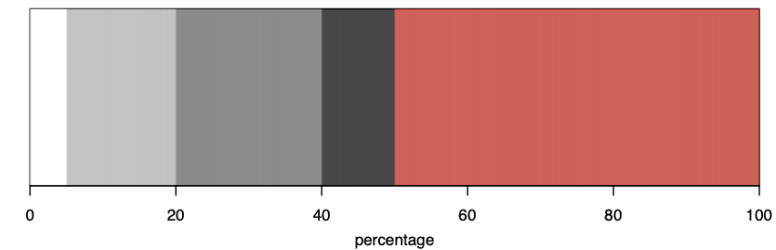
Frequency of spike SNVs for Omicron (22B/BA.5.*) (n = 1967)



BA.4 whole genome mutation prevalence over time



BA.5 whole genome mutation prevalence over time





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INYUVESI YAKWAZULU-NATALI



EDCTP

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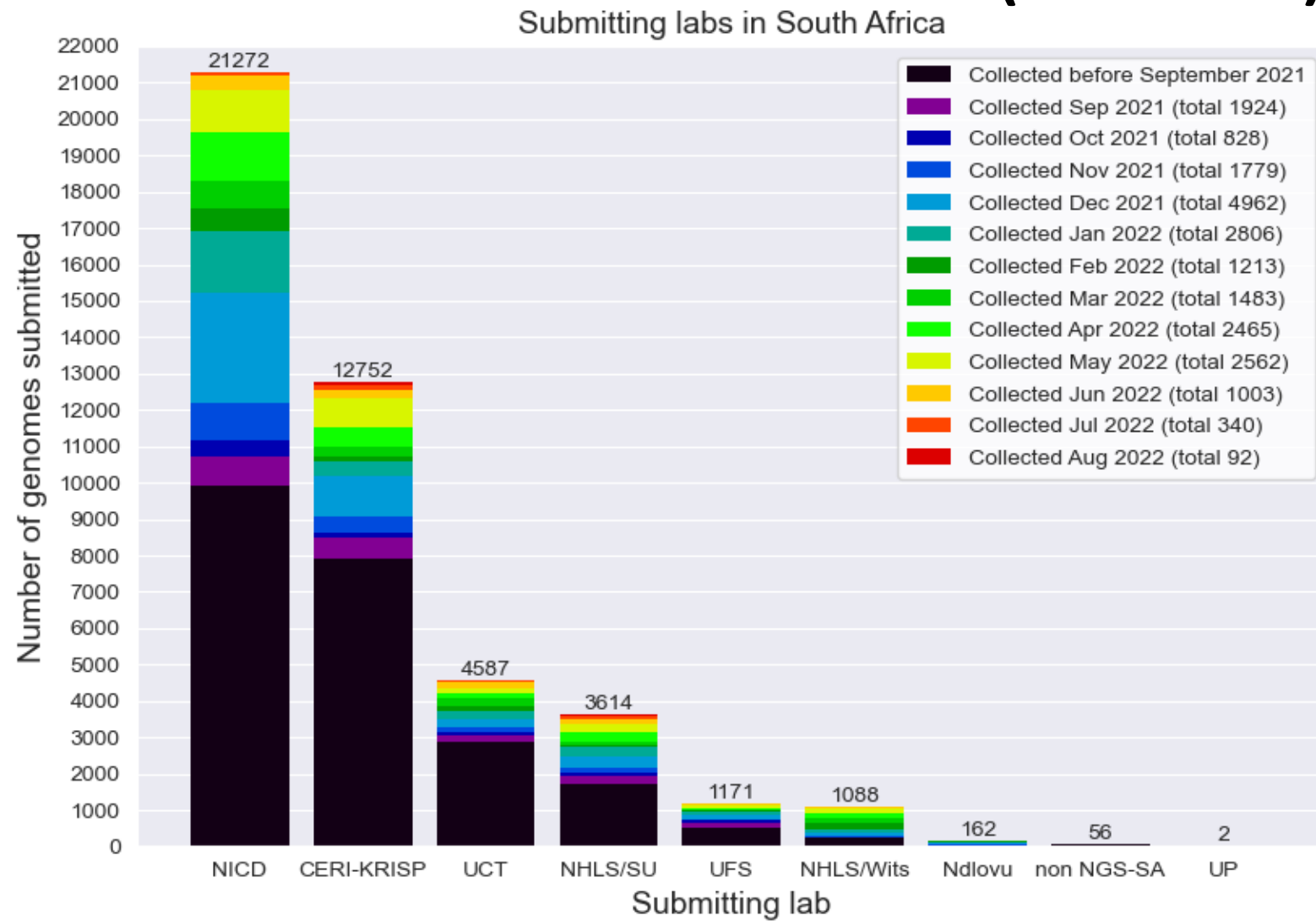
Glaudina Loots

SA MRC

Glenda Gray



South African genomes submitted per submitting lab, 2020 - 2022 (N=44 704)



NGS-SA Labs

CERI: Centre for Epidemic Response and Innovation

KRISP: KZN Research Innovation and Sequencing Platform

NDLOVU: Ndlovu Research Laboratories

NICD: National Institute for Communicable Diseases

NHLS: National Health Laboratory Service

SU: Stellenbosch University

UCT: University of Cape Town

UFS: University of the Free State

UP: University of Pretoria

Multiple labs from NGS-SA and collaborating public and private laboratories are contributing to sequencing, both as originating and as submitting (pictured here) laboratories.

Currently circulating Variants of Concern (VOC)

WHO label	Pango lineage•	GISAID clade	Nextstrain clade	Additional amino acid changes monitored°	Earliest documented samples	Date of designation
Omicron*	B.1.1.529	GR/484A	21K, 21L, 21M, 22A, 22B, 22C, 22D	+S:R346K +S:L452X +S:F486V	Multiple countries, Nov-2021	VUM: 24-Nov-2021 VOC: 26-Nov-2021

* Includes BA.1, BA.2, BA.3, BA.4, BA.5 and descendent lineages. It also includes BA.1/BA.2 circulating recombinant forms such as XE. WHO emphasizes that these descendant lineages should be monitored as distinct lineages by public health authorities and comparative assessments of their virus characteristics should be undertaken.

• Only found in a subset of sequences

Previously circulating Variants of Concern

WHO label	Pango lineage•	GISAID clade	Nextstrain clade	Earliest documented samples	Date of designation
Alpha	B.1.1.7	GRY	20I (V1)	United Kingdom, Sep-2020	VOC: 18-Dec-2020 Previous VOC: 09-Mar-2022
Beta	B.1.351	GH/501Y.V2	20H (V2)	South Africa, May-2020	VOC: 18-Dec-2020 Previous VOC: 09-Mar-2022
Gamma	P.1	GR/501Y.V3	20J (V3)	Brazil, Nov-2020	VOC: 11-Jan-2021 Previous VOC: 09-Mar-2022
Delta	B.1.617.2	G/478K.V1	21A, 21I, 21J	India, Oct-2020	VOI: 4-Apr-2021 VOC: 11-May-2021 Previous VOC: 7-Jun-2022

• Includes all descendant lineages. See the cov-lineages.org and the Pango network websites for further details.

<https://www.who.int/en/activities/tracking-SARS-CoV-2-variants/> accessed 12 August 2022

Submission of routine specimens for sequencing

- representative of multiple geographic regions (provinces/districts/health facilities) from individuals of
 - all ages
 - over as many time periods during the SARS-CoV-2 epidemic in South Africa
- requested that testing laboratories in both the private and public sectors, submit respiratory samples to their closest NGS-SA sequencing laboratory on a routine basis (ideally every week) as follows, depending on the capacity of the testing laboratory:
 - All positives samples should be sent every week (NGS-SA laboratory will perform random sampling as described below) **OR**
 - A weekly selection of approximately 10%-20% of randomly selected positive samples should be sent every week. Number of selected samples will depend on the size of laboratory and how many other laboratories are drained by the submitting laboratory.

Submission of special interest specimens for sequencing

In addition to routine samples mentioned above, please send specimens separately to above and clearly marked if:

- Suspected vaccine breakthrough (≥ 14 days after vaccine), especially if hospitalised and clinically severe
- Suspected re-infection (≥ 90 days after previous episode), especially if hospitalised and clinically severe
- Prolonged shedding with high SARS-CoV-2 viral loads (i.e. Ct values less than 30 for more than 1 month post-primary diagnosis) in immunocompromised individuals
- Possible animal-to-human transmission
- Suspected cases of importation from another country, especially countries known to harbour SARS-CoV-2 variants of concern or countries with little available information
- Clusters of “unusual” cases (e.g., in terms of disease presentation, patient groups affected, etc.)