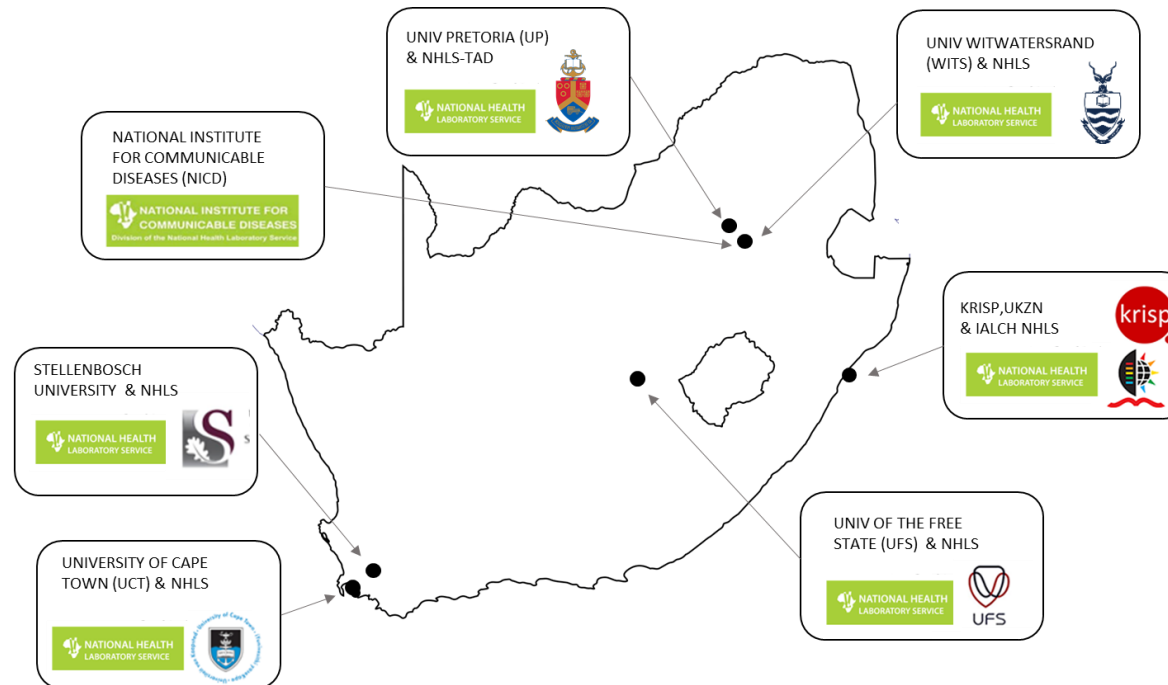


## SARS-CoV-2 Sequencing Update 17 March 2023



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](http://www.gisaid.org)) on 17 March 2023 at 15h00



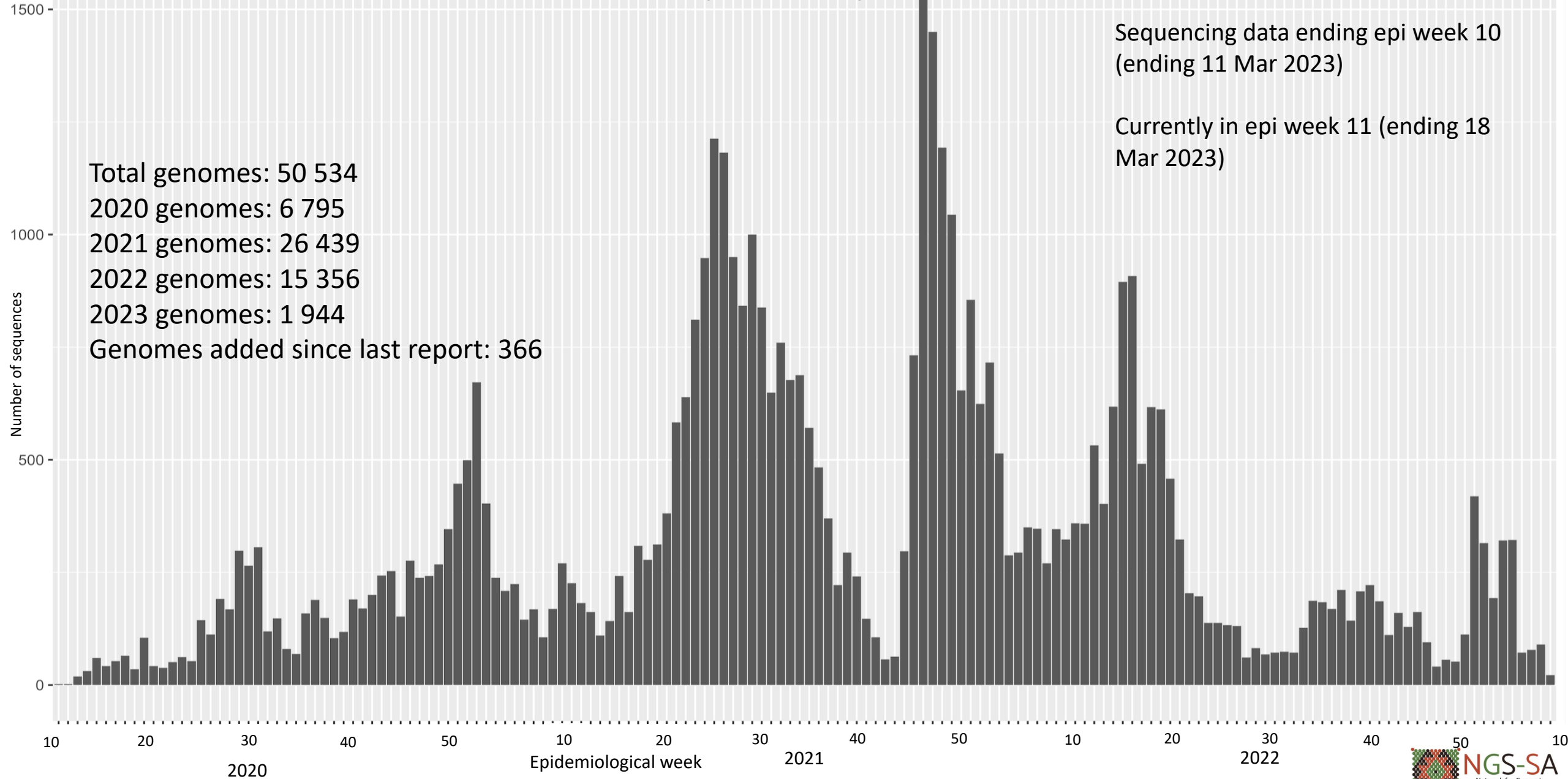
Data license: <https://www.gisaid.org/registration/terms-of-use/>

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

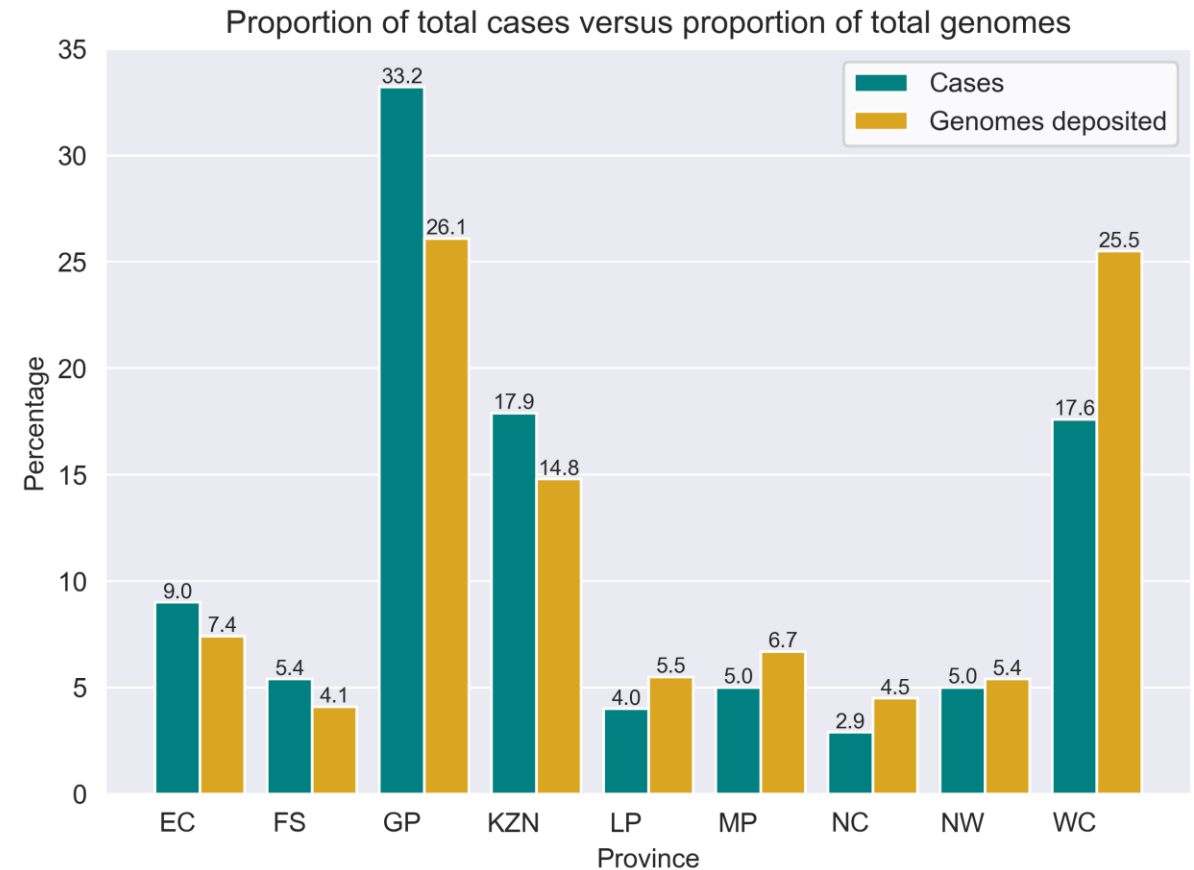
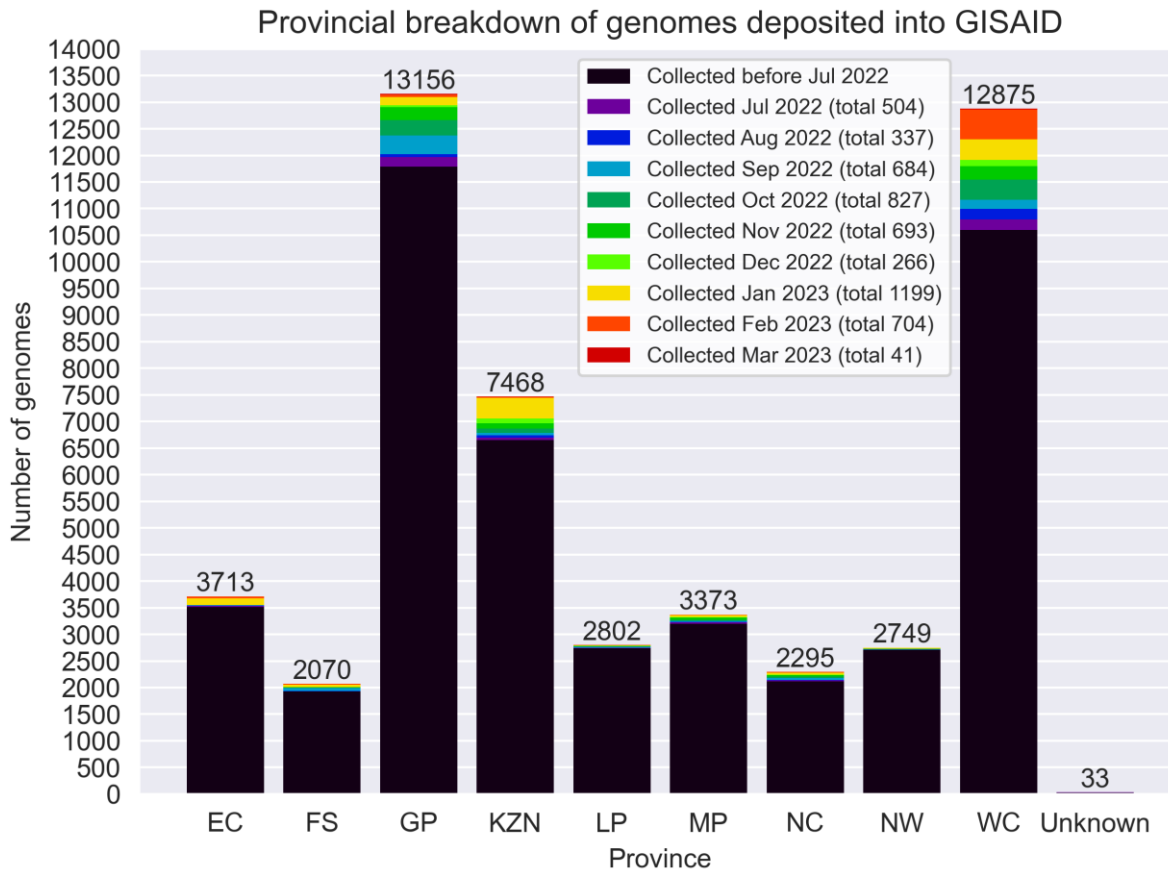
# Number of South African genomes deposited on GISAID, by specimen collection week, 2020 – 2023

(N=50 534\*)

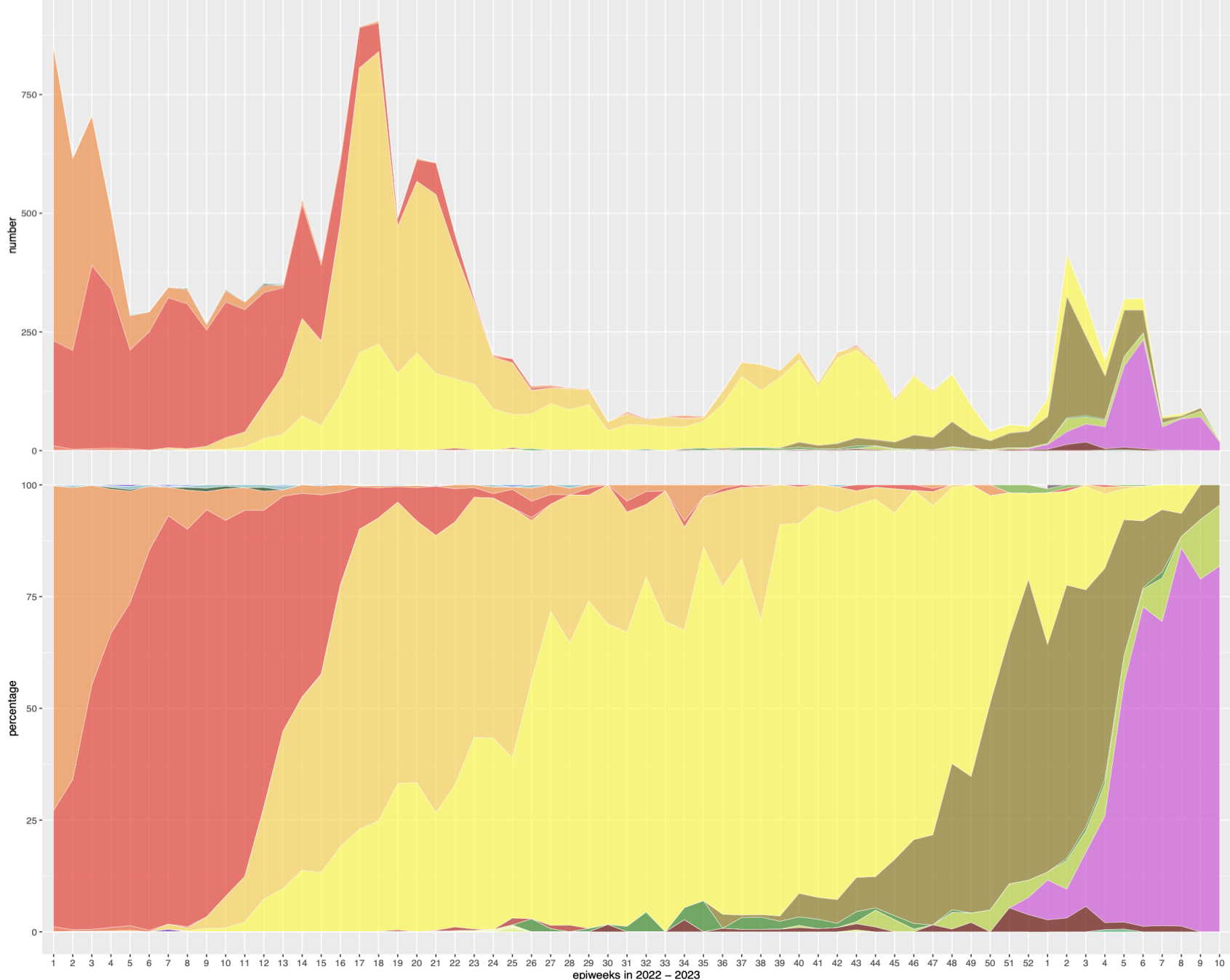


\*This represents the cleaned, de-duplicated dataset of unique **National and Pneumonia Surveillance** sequences. This dataset will be used for all further figures.

# GISAID genomes vs total cases, 2020 – 2023 (N=50 534)



# Number and percentage of clades by epiweek in South Africa, 2022-2023 (17 257\*)

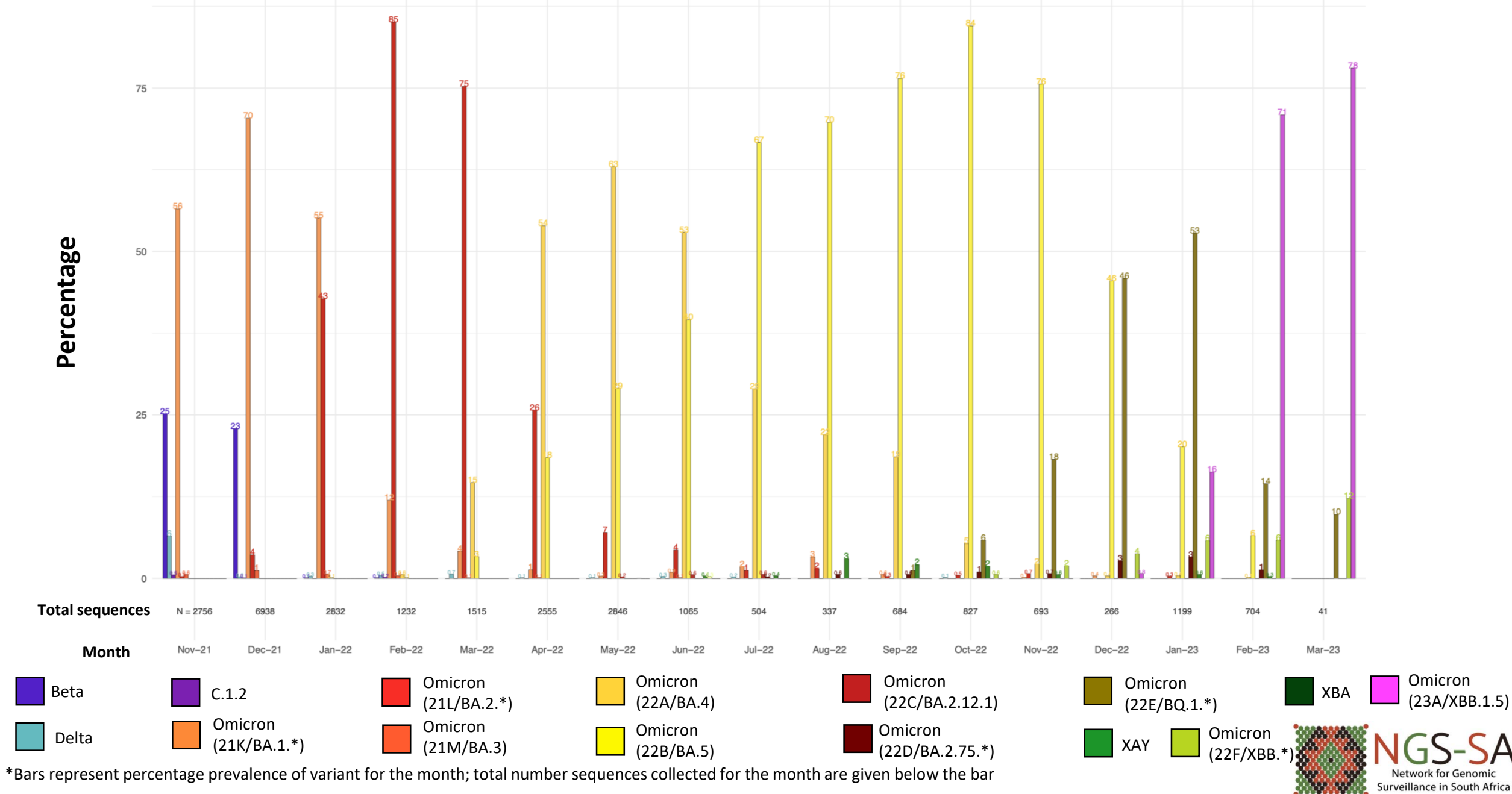


Sequencing data  
ending epi week 10  
(ending 11 Mar 2023)

Currently in epi week  
11 (ending 18 Mar  
2023)

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

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

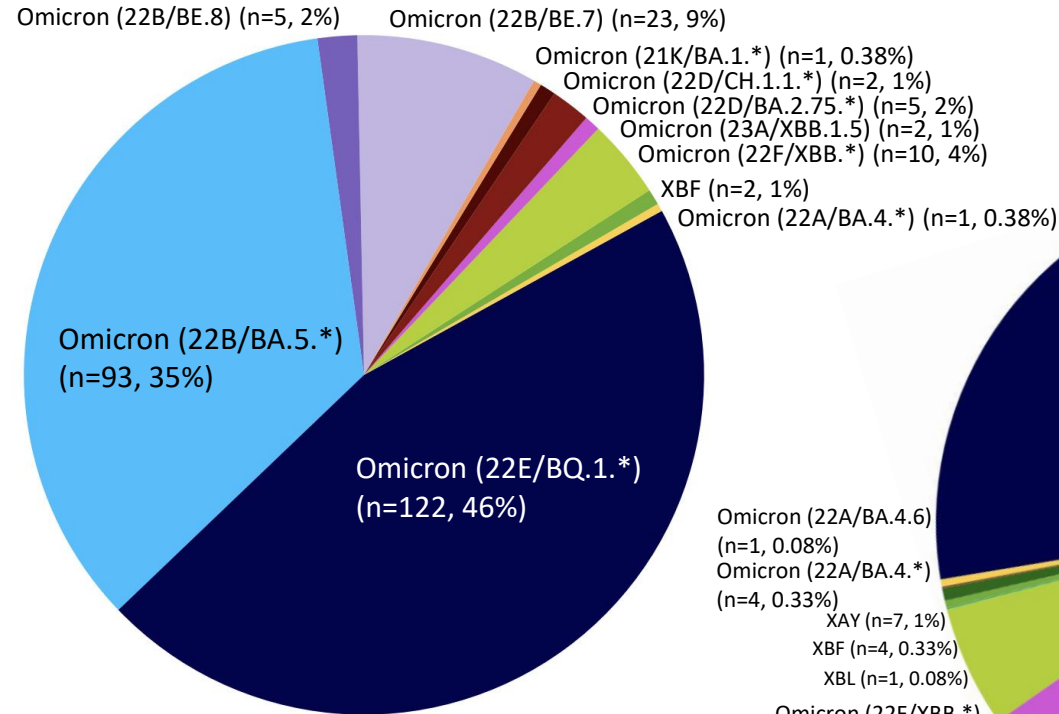


# Prevalence of Variants of Concern (VOC) and Variants of Interest (VOI) in

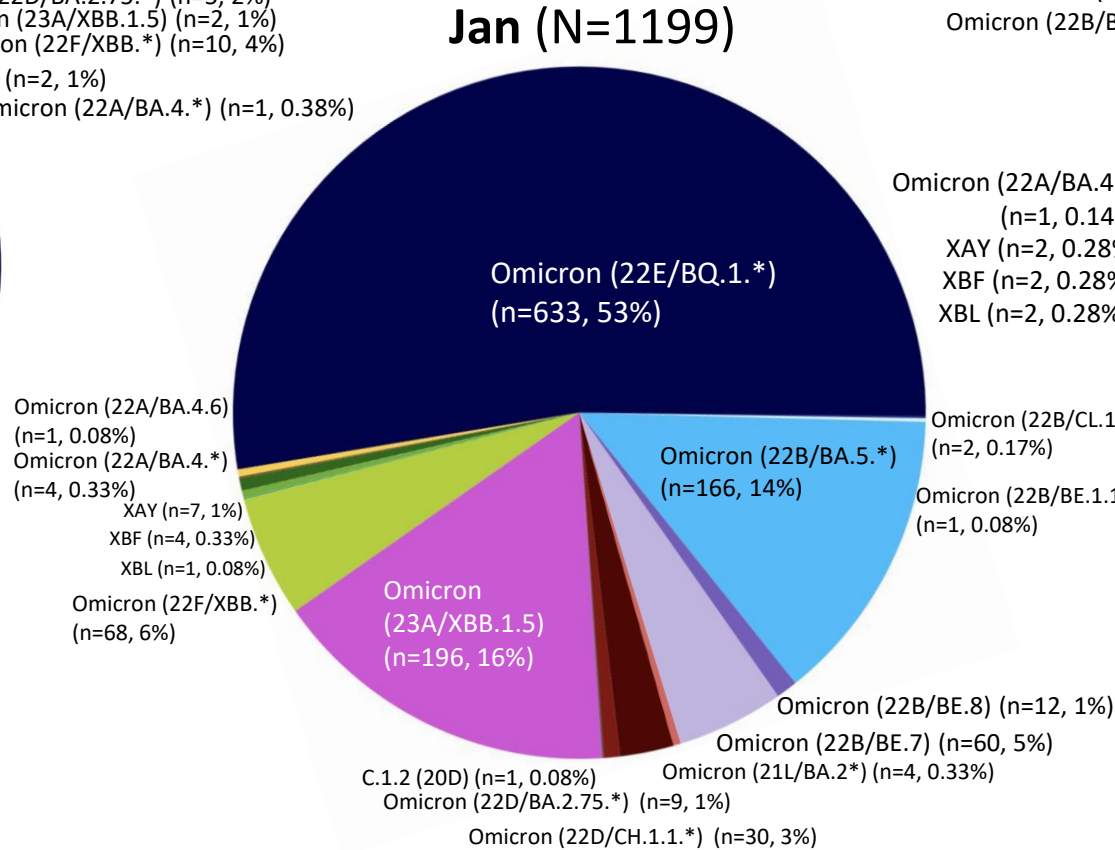
Dec (N=266)

Dec 2022– Feb 2023

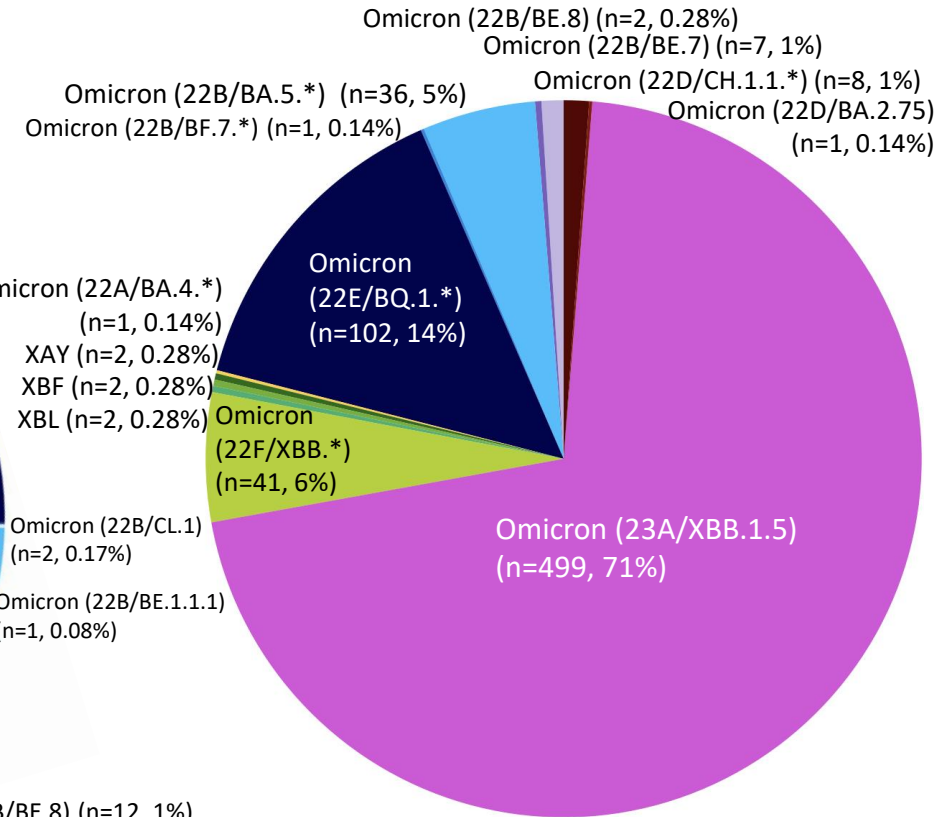
Feb (N=704)



Total Omicron in December: 266 (100%)



Total Omicron in January: 1191 (99.3%)

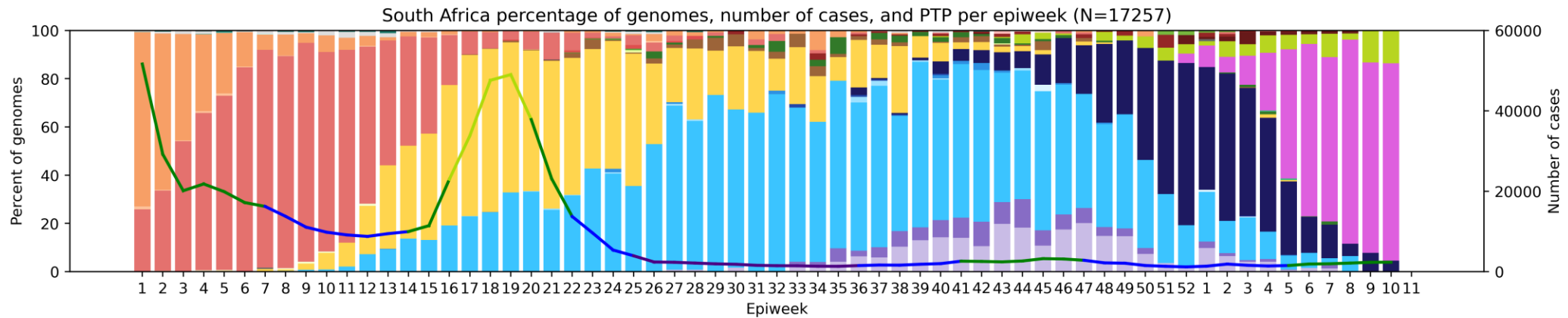
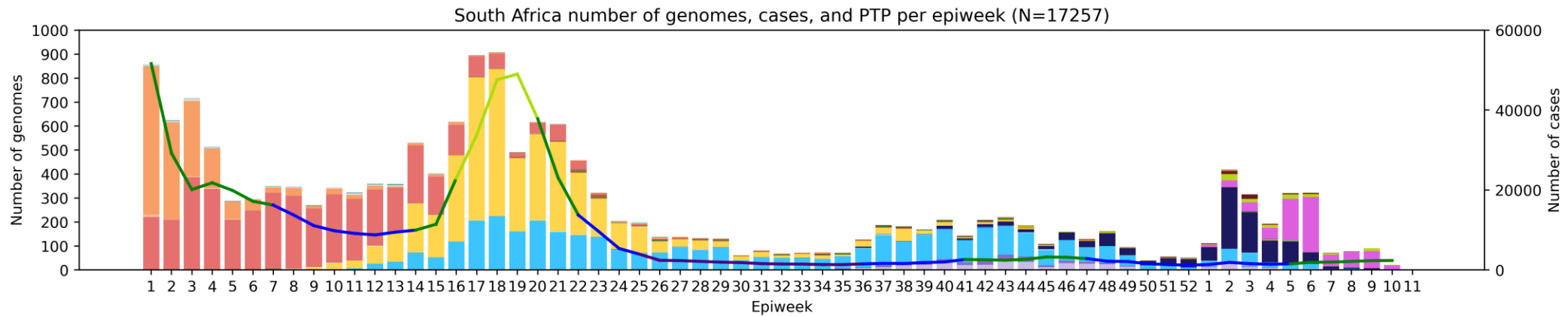


Total Omicron in February: 702 (99.7%)

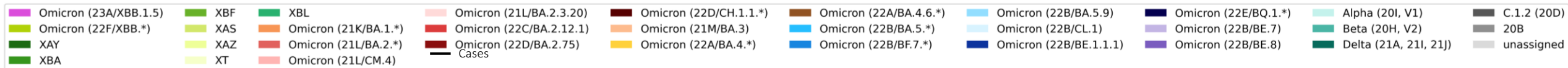
Omicron (23A/XBB.1.5)	XBF	XBL	Omicron (21L/BA.2.3.20)	Omicron (22D/CH.1.1.*)	Omicron (22A/BA.4.6.*)	Omicron (22B/BA.5.9)	Omicron (22E/BQ.1.*)	Alpha (20I, V1)	C.1.2 (20D)
Omicron (22F/XBB.*)	XAS	XBL	Omicron (22C/BA.2.12.1)	Omicron (22D/CH.1.1.*)	Omicron (21M/BA.3)	Omicron (22B/CL.1)	Omicron (22B/BE.7)	Beta (20H, V2)	20B
XAY	XAZ	XBL	Omicron (21L/BA.2.*)	Omicron (22D/BA.2.75)	Omicron (22A/BA.4.*)	Omicron (22B/BE.7.*)	Omicron (22B/BE.1.1.1)	Delta (21A, 21I, 21J)	unassigned
XBA	XT	XBL	Omicron (21L/CM.4)						

Note: XBF and XBL are Omicron-Omicron recombinants and so are counted in the total number of Omicrons.

# South Africa, 2022-2023, n = 17 257\*



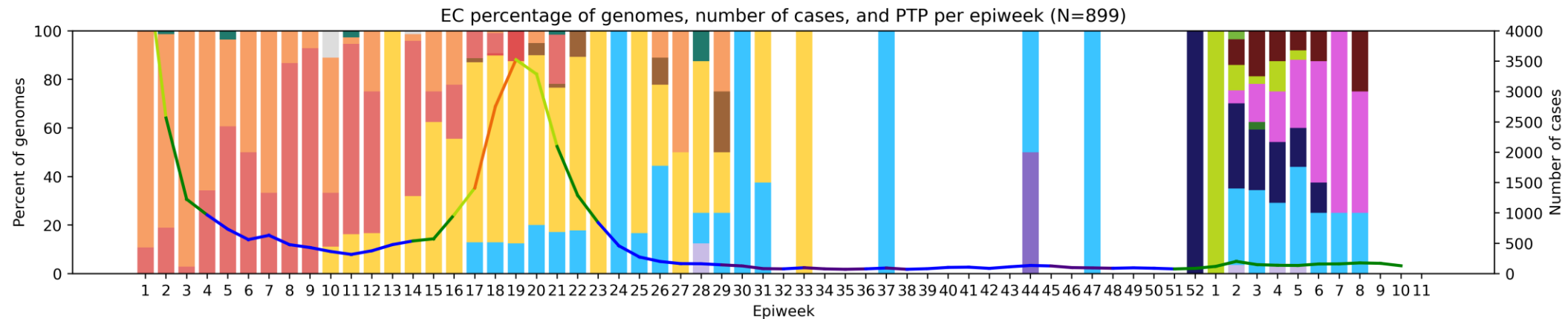
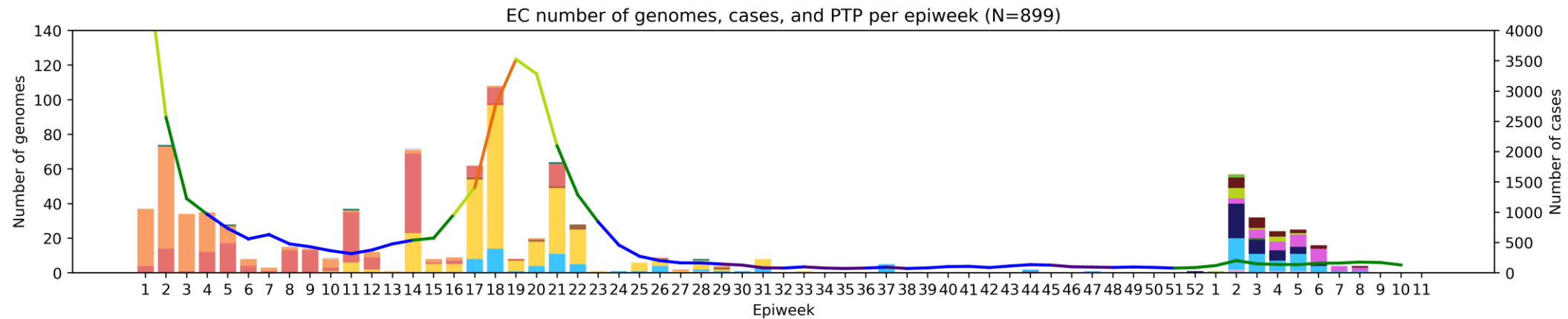
## Clade key (bar graph)



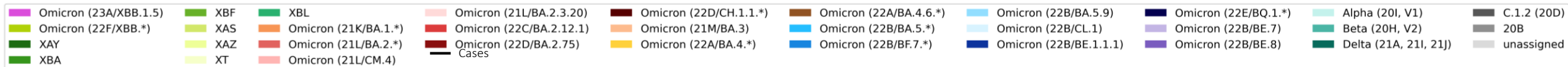
## Weekly percentage testing positive key (line graph)

# Eastern Cape Province, 2022-2023, n = 899

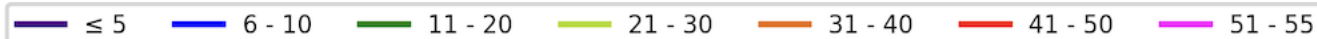
Genomes added since last report: 48\*



## Clade key (bar graph)

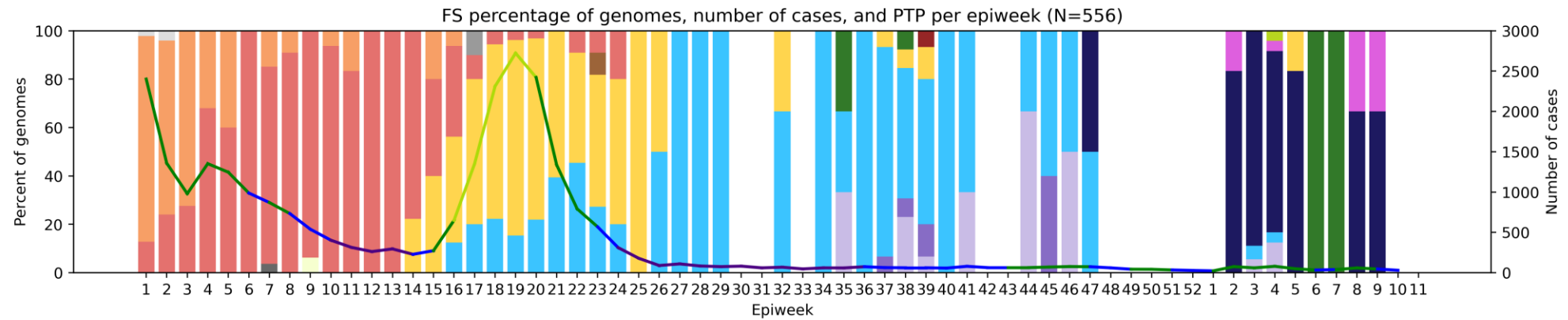
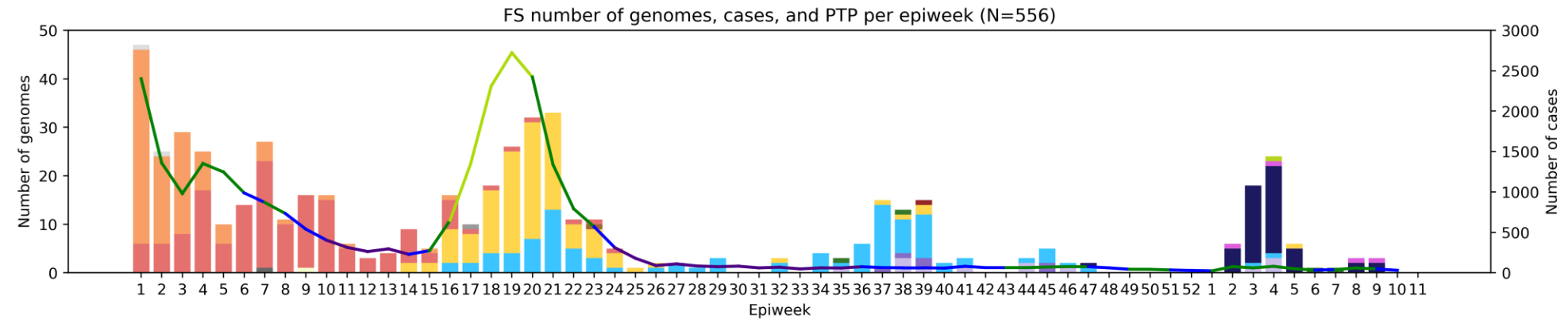


## Weekly percentage testing positive key (line graph)

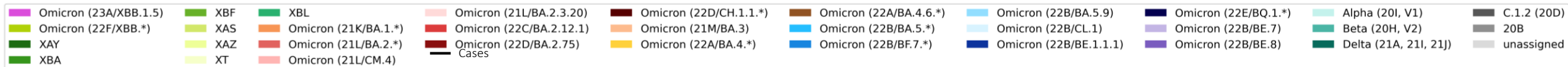


# Free State Province, 2022-2023, n = 556

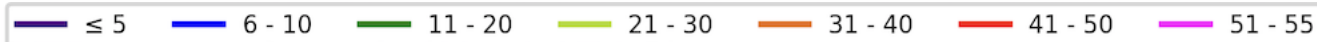
Genomes added since last report: 10\*



## Clade key (bar graph)



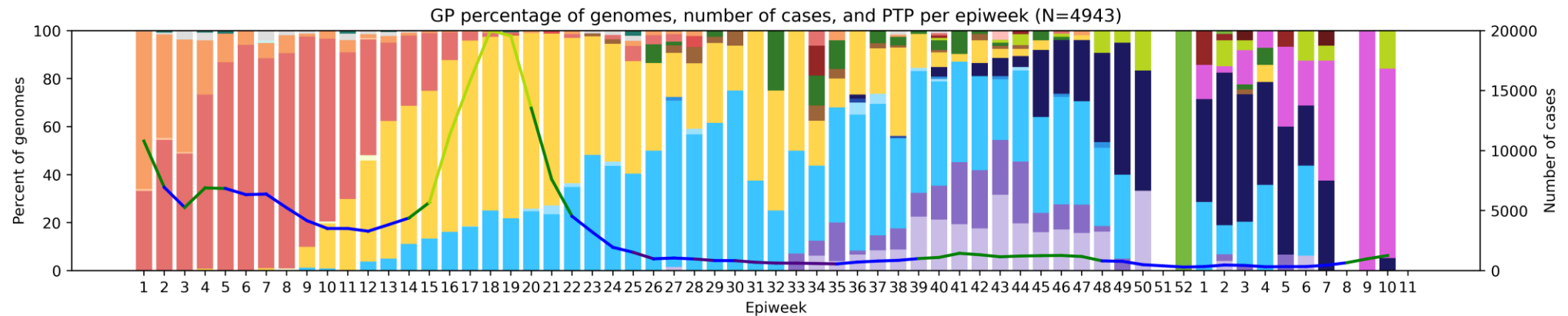
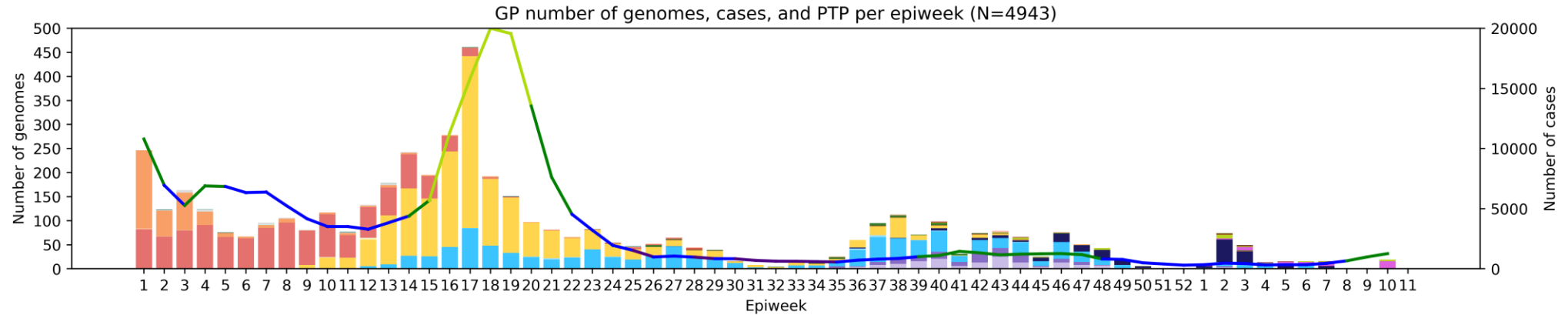
## Weekly percentage testing positive key (line graph)



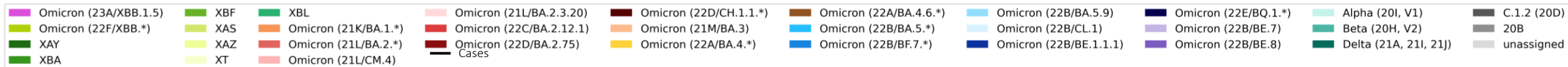
\*May include genomes from 2020 and 2021 which are not pictured here and are not included in the slide total.

# Gauteng Province, 2022-2023, n = 4943

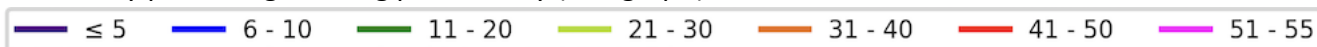
Genomes added since last report: 48\*



## Clade key (bar graph)

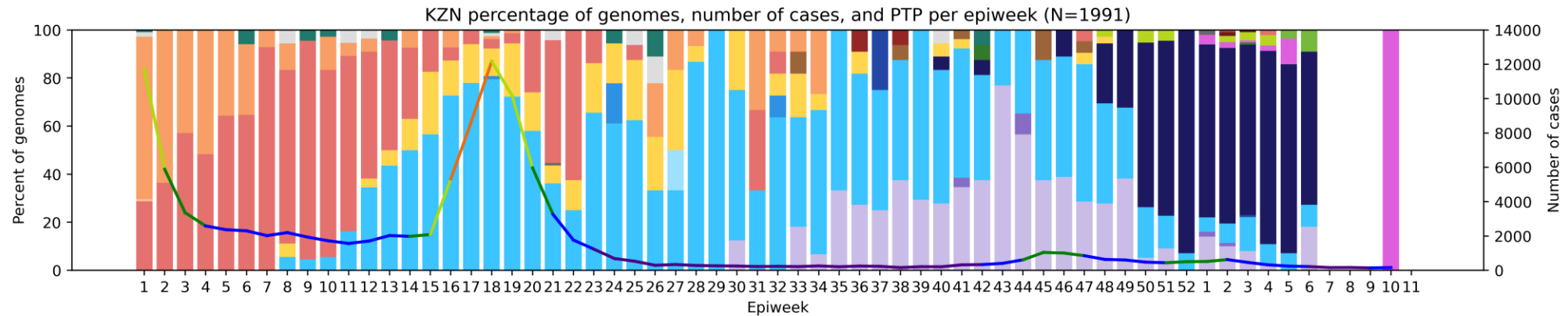
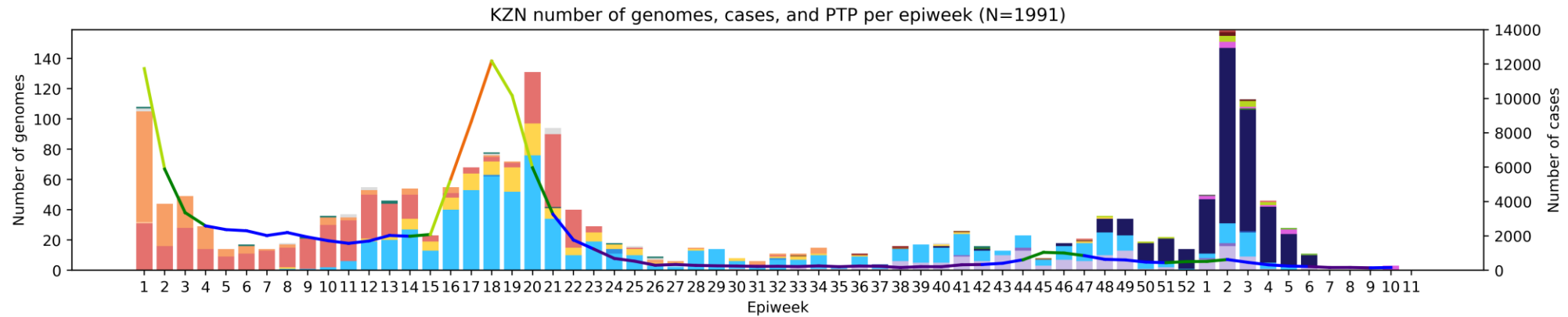


## Weekly percentage testing positive key (line graph)

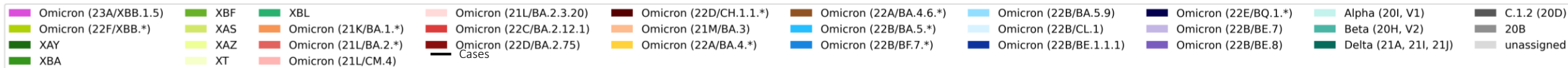


# KwaZulu-Natal Province, 2022-2023, n = 1991

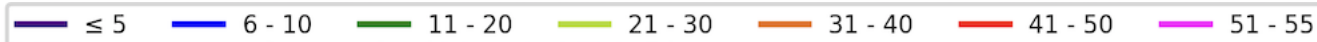
Genomes added since last report: 24\*



## Clade key (bar graph)

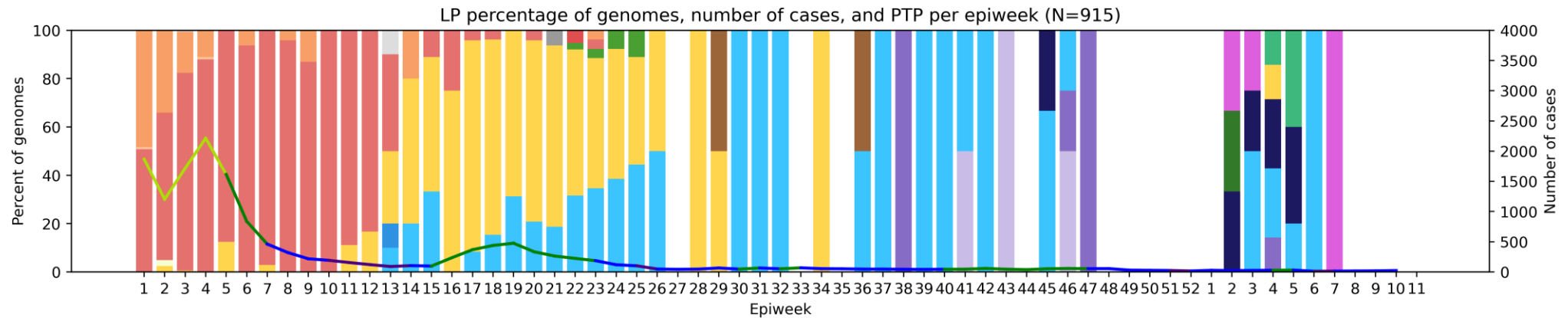
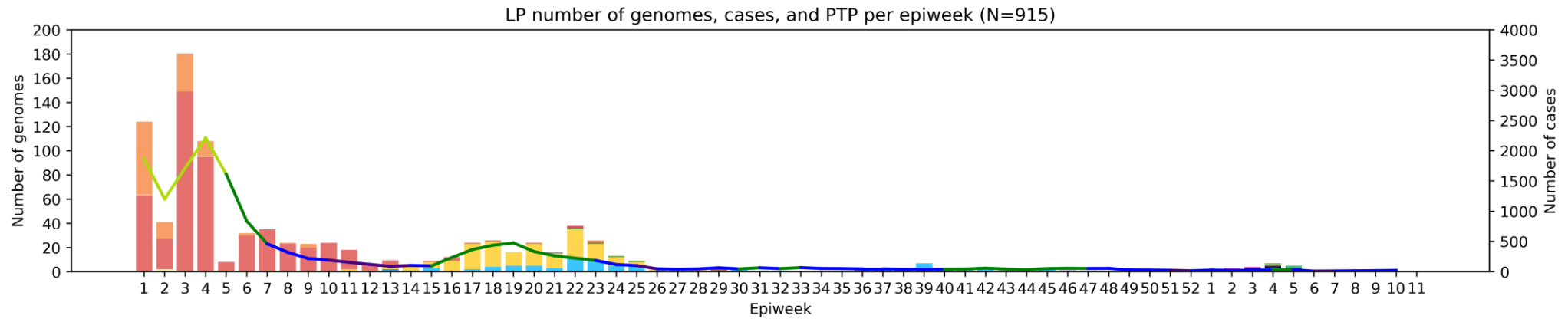


## Weekly percentage testing positive key (line graph)

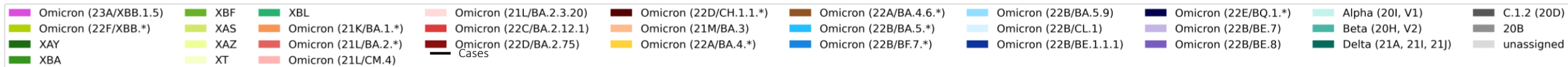


# Limpopo Province, 2022-2023, n = 915

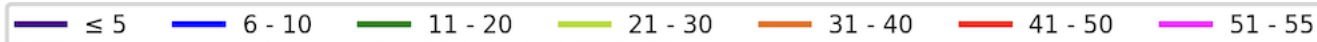
Genomes added since last report: 5\*



## Clade key (bar graph)



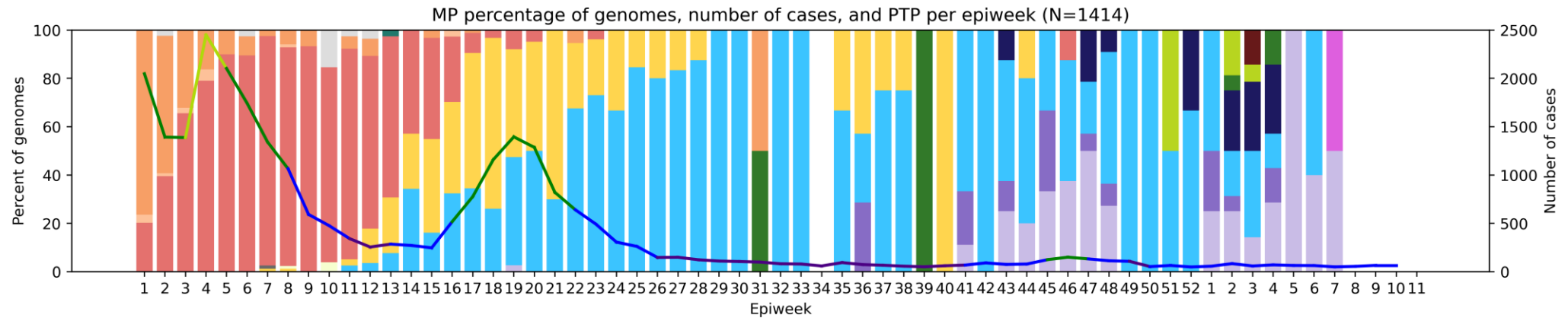
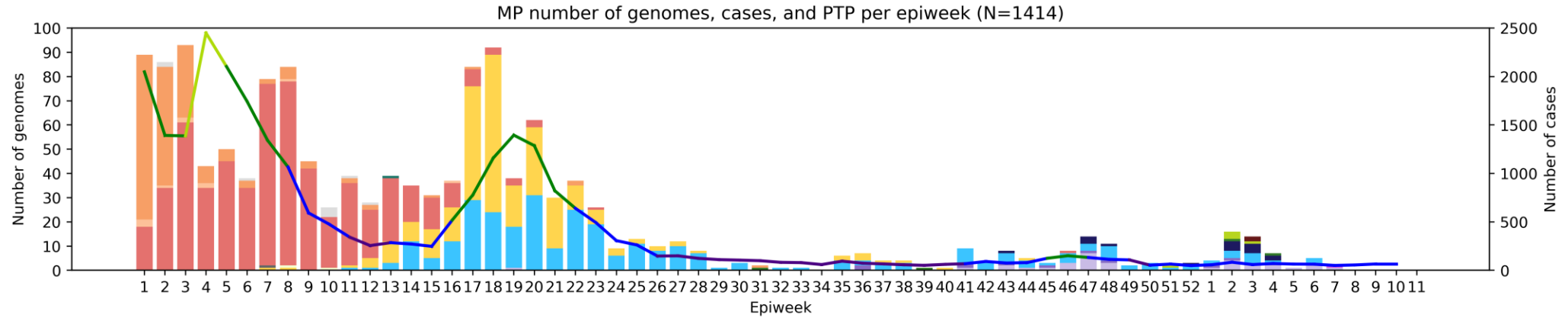
## Weekly percentage testing positive key (line graph)



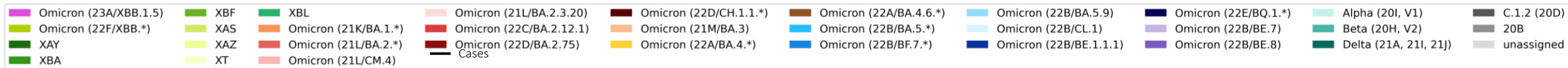
\*May include genomes from 2020 and 2021 which are not pictured here and are not included in the slide total.

# Mpumalanga Province, 2022-2023, n = 1414

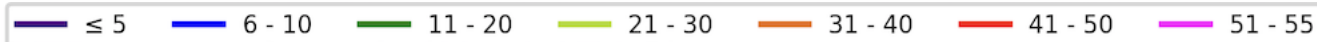
Genomes added since last report: 6\*



## Clade key (bar graph)



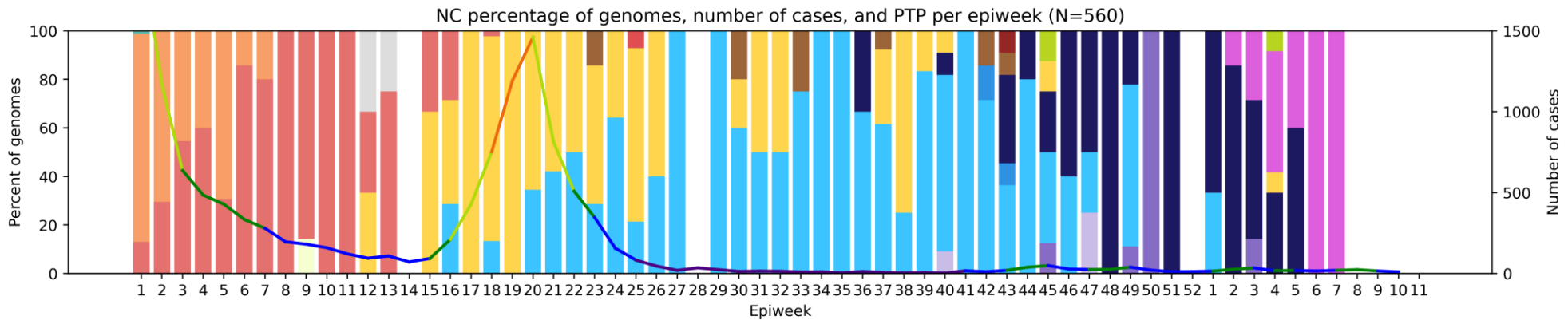
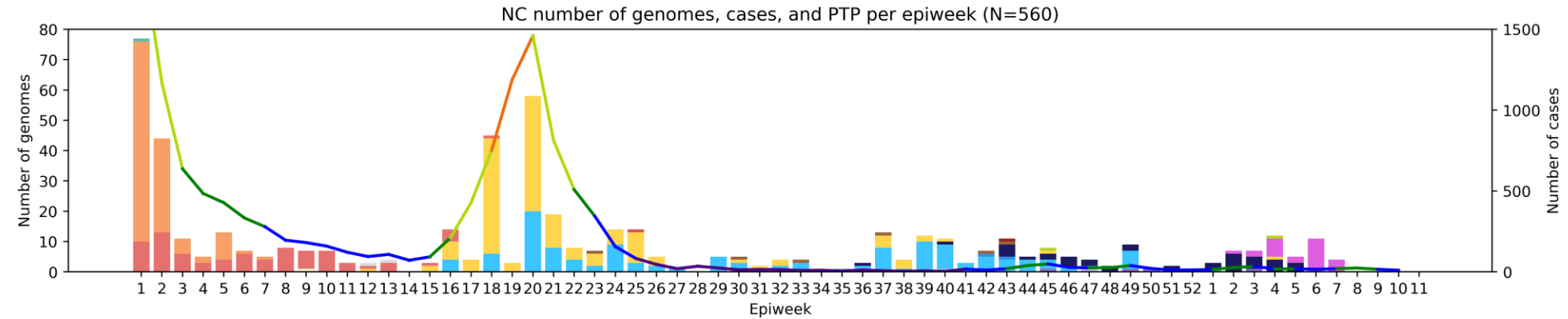
## Weekly percentage testing positive key (line graph)



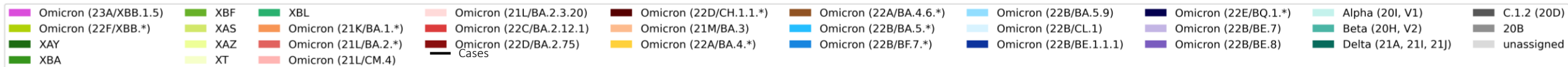
\*May include genomes from 2020 and 2021 which are not pictured here and are not included in the slide total.

# Northern Cape Province, 2022-2023, n = 560

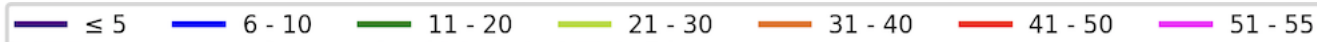
Genomes added since last report: 15\*



## Clade key (bar graph)



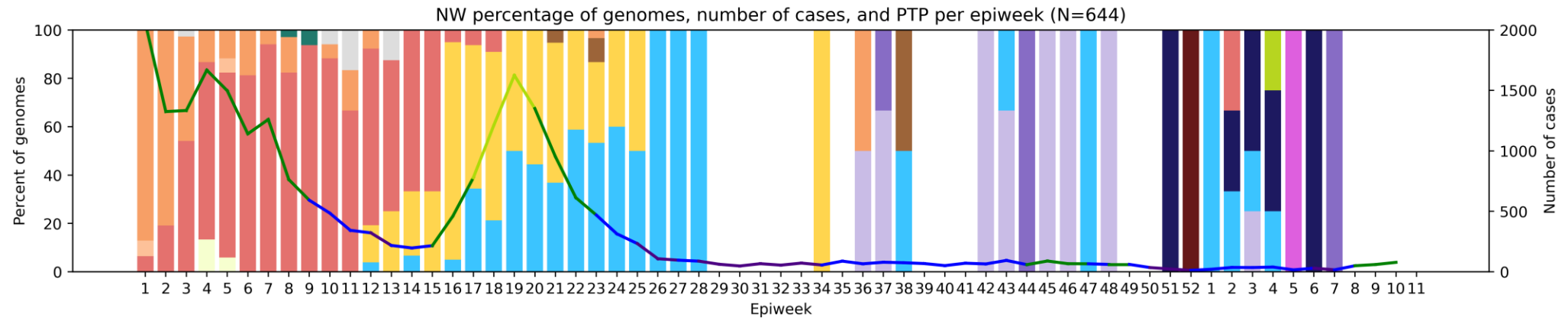
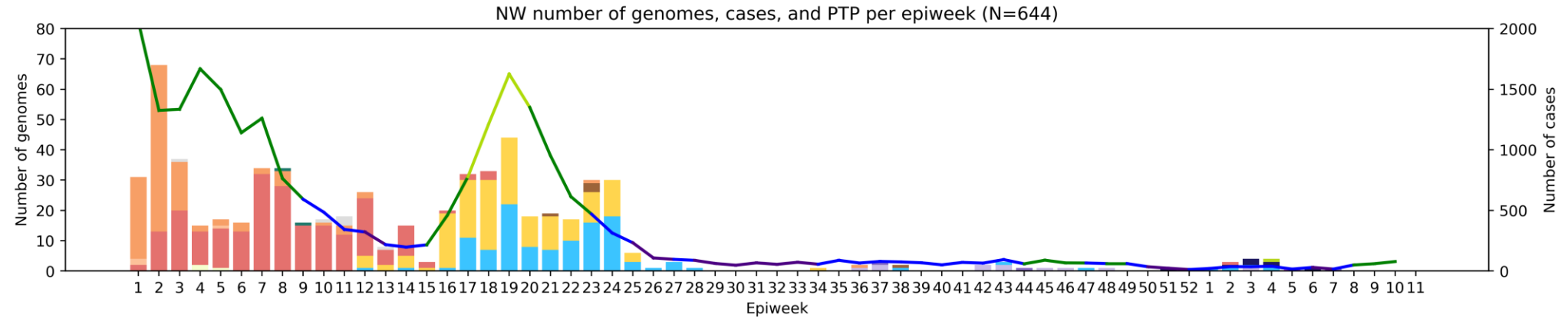
## Weekly percentage testing positive key (line graph)



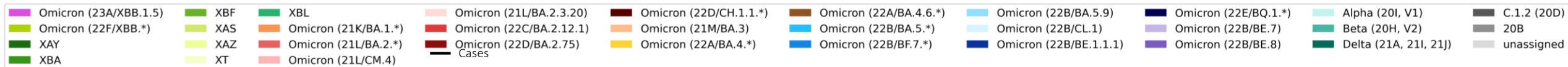
\*May include genomes from 2020 and 2021 which are not pictured here and are not included in the slide total.

# North West Province, 2022-2023, n = 644

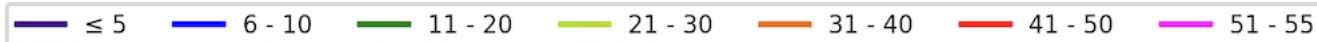
Genomes added since last report: 4\*



## Clade key (bar graph)



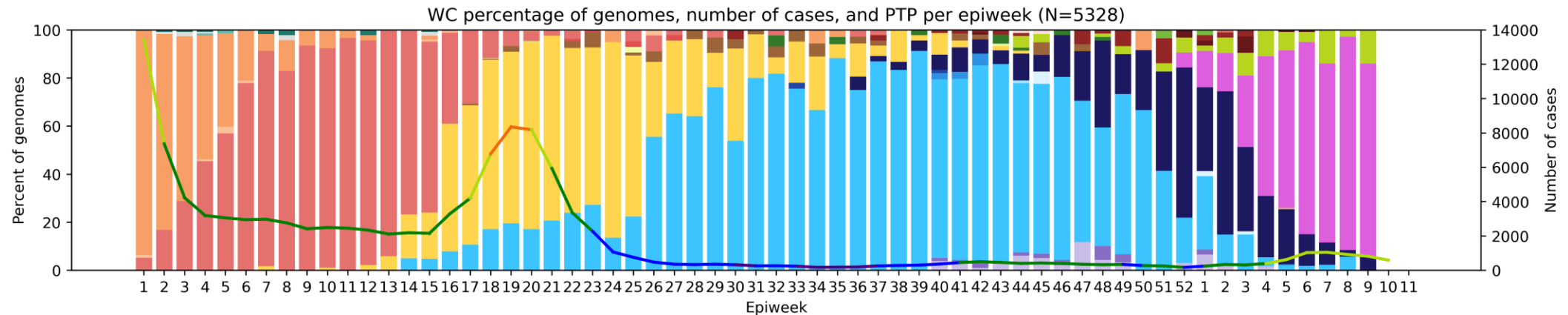
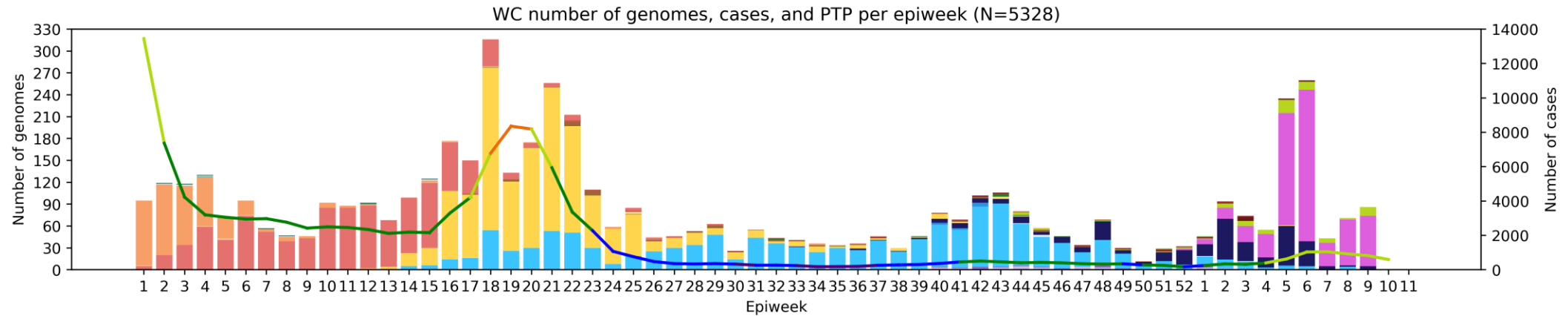
## Weekly percentage testing positive key (line graph)



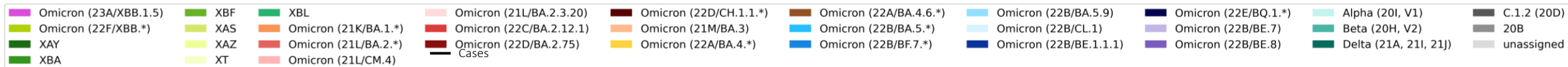
\*May include genomes from 2020 and 2021 which are not pictured here and are not included in the slide total.

# Western Cape Province, 2022-2023, n = 5328

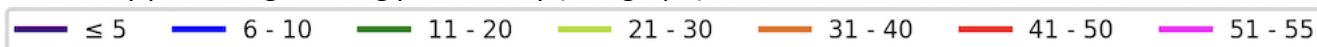
Genomes added since last report: 206\*



## Clade key (bar graph)



## Weekly percentage testing positive key (line graph)



\*May include genomes from 2020 and 2021 which are not pictured here and are not included in the slide total.

# Summary

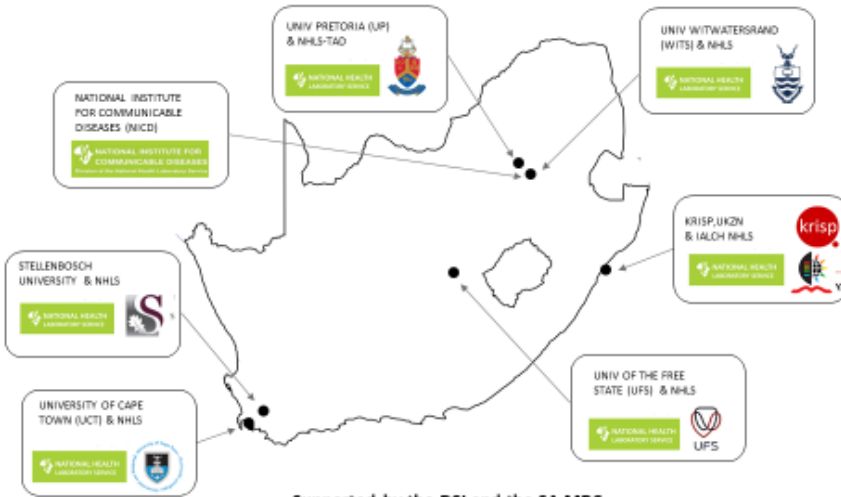
- **Sequencing update**

- All provinces have sequences for January 2023 and all provinces, except the North West, have sequences for February 2023. March sequences are from Free state, Western cape, Kwa-Zulu Natal and Gauteng.

- **Variant of Concern Omicron in South Africa**

- Omicron continued to dominate in December (100%), January (99%) and makes up 100% of February and March sequences
- BQ.1 and sub-lineages were the dominant Omicron lineage in December (46%) and January (54%).
- XBB.1.5 was detected in December 2022 (n=2, 1%) and January 2023 (n=196, 16%), and is the dominant lineage in February 2023 (n=499, 71%) and March 2023 (n=32, 78%).
- BA.2.75.\* continued to be detected at a low prevalence in December, January and February ( $\leq 3\%$ )





Supported by the DSI and the SA MRC



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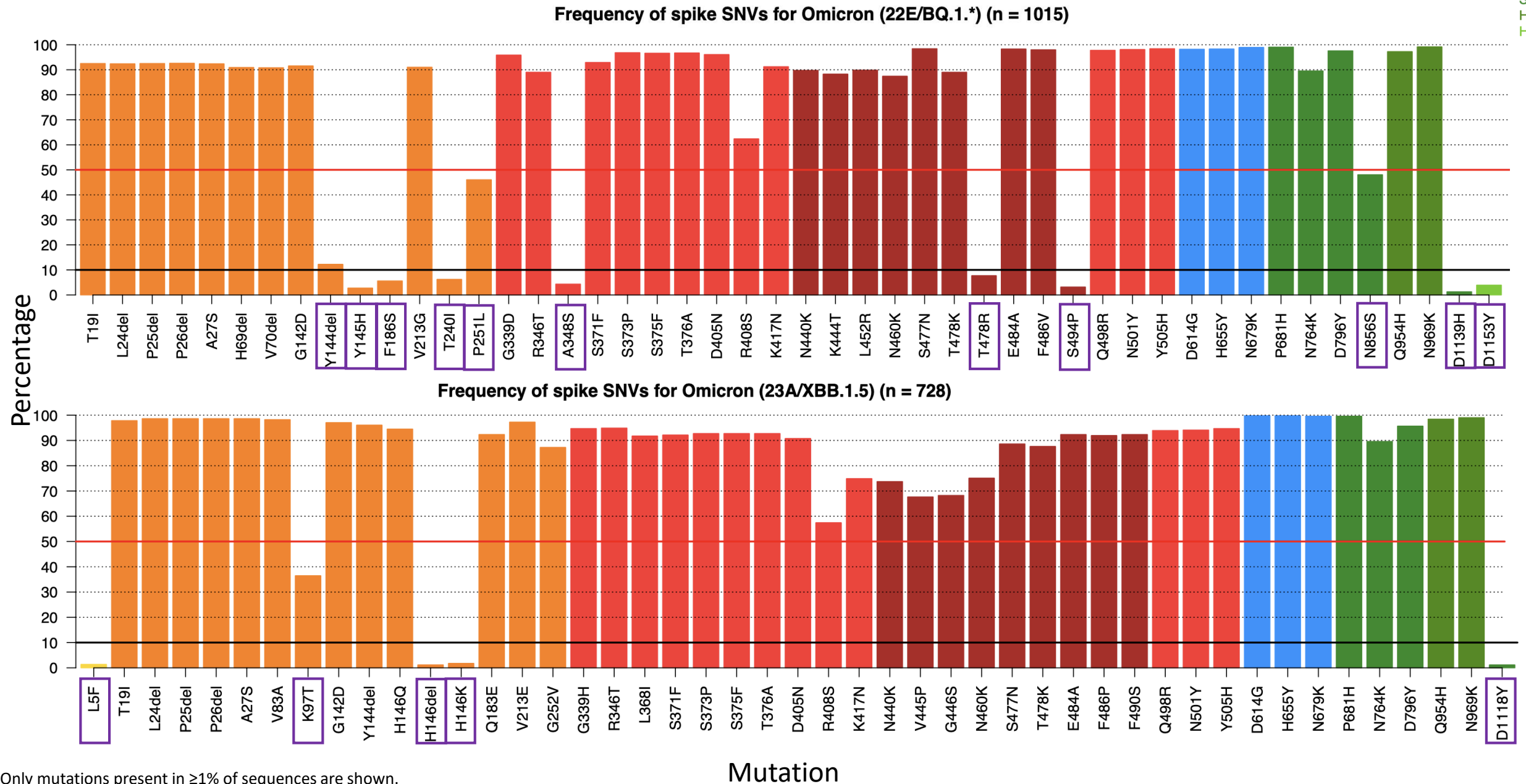
EDCTP

This project (RIA2020EF-3030) is part of the EDCTP2 programme supported by the European Union



# BQ.1\* and XBB.1.5\* spike mutations\*

SP  
NTD  
RBD  
RBM  
S1  
S2  
HR1  
HR2



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NICD SARS-CoV-2 Sequencing Group

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# Currently circulating Variants of Interest (VOI) as of 15 March 2023

Pango lineage●	Nextstrain clade	Genetic features	Earliest documented samples	Date of designation
XBB.1.5	23A	Recombinant of BA.2.10.1 and BA.2.75 sublineages, i.e. BJ1 and BM.1.1.1, with a breakpoint in S1.  XBB + S:F486P	05-01-2022	11-Jan-2023

# Omicron subvariants under monitoring

Pango lineage <sup>#</sup> (+ mutation)	GISAID clade	Nextstrain clade	Relationship to circulating VOC lineages	Spike genetic features	Earliest documented samples
BQ.1 <sup>§</sup>	GRA	22E	BA.5 sublineage	BQ.1 and BQ.1.1: BA.5 + S:R346T, S:K444T, S:N460K	07-02-2022
BA.2.75 <sup>§</sup>	GRA	22D	BA.2 sublineage	BA.2.75: BA.2 + S:K147E, S:W152R, S:F157L, S:I210V, S:G257S, S:D339H, S:G446S, S:N460K, S:Q493R reversion	31-12-2021
CH.1.1 <sup>§</sup>	GRA	22D	BA.2 sublineage	BA.2.75 + S:L452R, S:F486S	27-07-2022
XBB <sup>μ</sup>	GRA	22F	Recombinant of BA.2.10.1 and BA.2.75 sublineages, i.e. BJ1 and BM.1.1.1, with a breakpoint in S1	BA.2+ S:V83A, S:Y144-, S:H146Q, S:Q183E, S:V213E, S:G252V, S:G339H, S:R346T, S:L368I, S:V445P, S:G446S, S:N460K, S:F486S, S:F490S	13-08-2022
XBF	GRA		Recombinant of BA.5.2.3 and CJ.1 (BA.2.75.3 sublineage)	BA.5 + S:K147E, S:W152R, S:F157L, S:I210V, S:G257S, S:G339H, S:R346T, S:G446S, S:N460K, S:F486P, S:F490S	27-07-2022

<sup>#</sup> includes descendent lineages

<sup>§</sup> additional mutation outside the spike protein: ORF1a: Q556K, L3829F, ORF1b: Y264H, M1156I, N1191S, N: E136D, ORF9b: P10F

<sup>§</sup> additional mutations outside of the spike protein: ORF1a: S1221L, P1640S, N4060S, ORF1b: G662S, E: T11A

<sup>μ</sup> additional mutations outside of the spike protein: ORF1a: K47R, ORF1b: G662S, S959P, E: T11A, ORF8: G8\*

# 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 ( $\geq 14$  days after vaccine), especially if hospitalised and clinically severe
- Suspected re-infection ( $\geq 90$  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.)