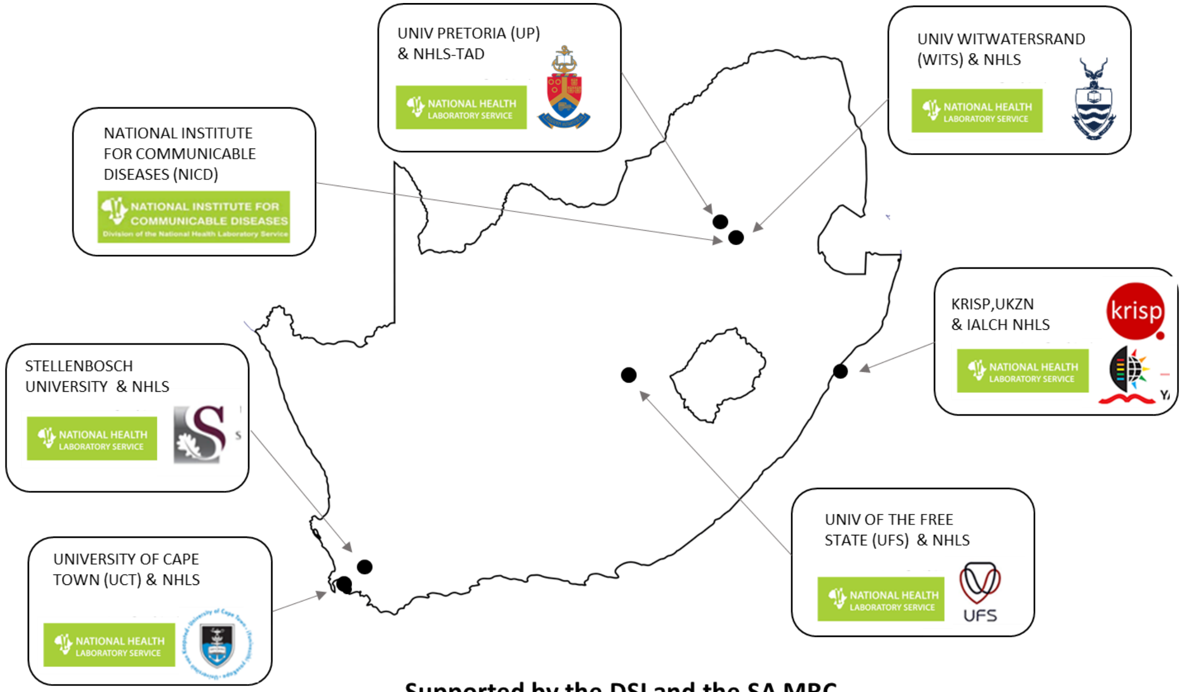
NGS-SA
Network for Genomic
Surveillance in South Africa

Network for Genomic Surveillance in South Africa (NGS-SA)

SARS-CoV-2 Sequencing Update

5 May 2023



UNIV PRETORIA (UP) & NHLS-TAD

UNIV WITWATERSRAND (WITS) & NHLS

NATIONAL INSTITUTE FOR COMMUNICABLE DISEASES (NICD)

KRISP,UKZN & IALCH NHLS

STELLENBOSCH UNIVERSITY & NHLS

UNIV OF THE FREE STATE (UFS) & NHLS

UNIVERSITY OF CAPE TOWN (UCT) & NHLS

Supported by the DSI and the SA MRC

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

Prepared by the National Institute for Communicable Diseases (NICD) of the National Health Laboratory (NHLS) on behalf of the Network for Genomics Surveillance in South Africa (NGS-SA)

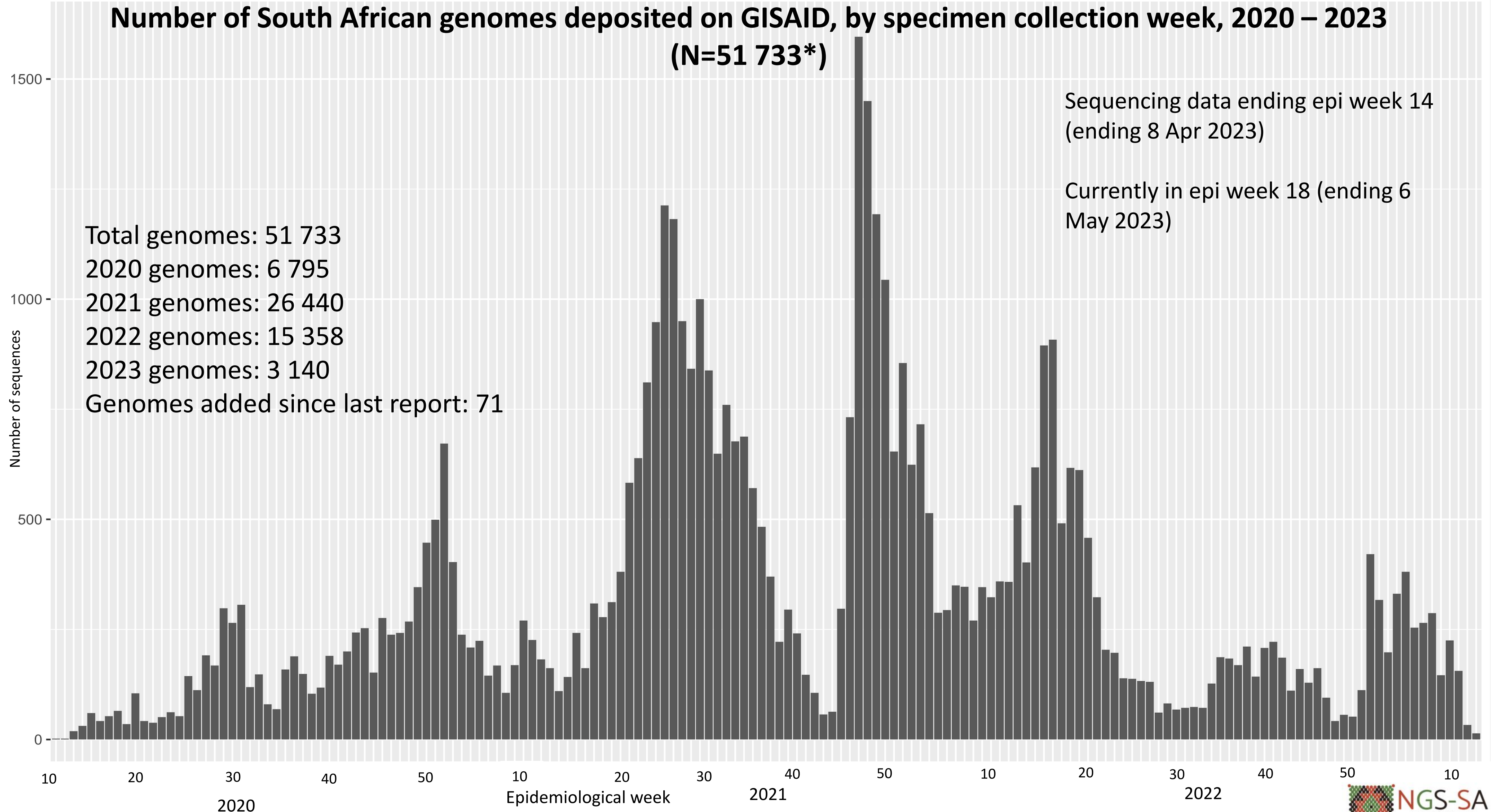
The genomic data presented here are based on South African SARS-CoV-2 sequence data downloaded from GISAID (www.gisaid.org) on 05 May 2023 at 08h44

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=51 733*)



Sequencing data ending epi week 14 (ending 8 Apr 2023)

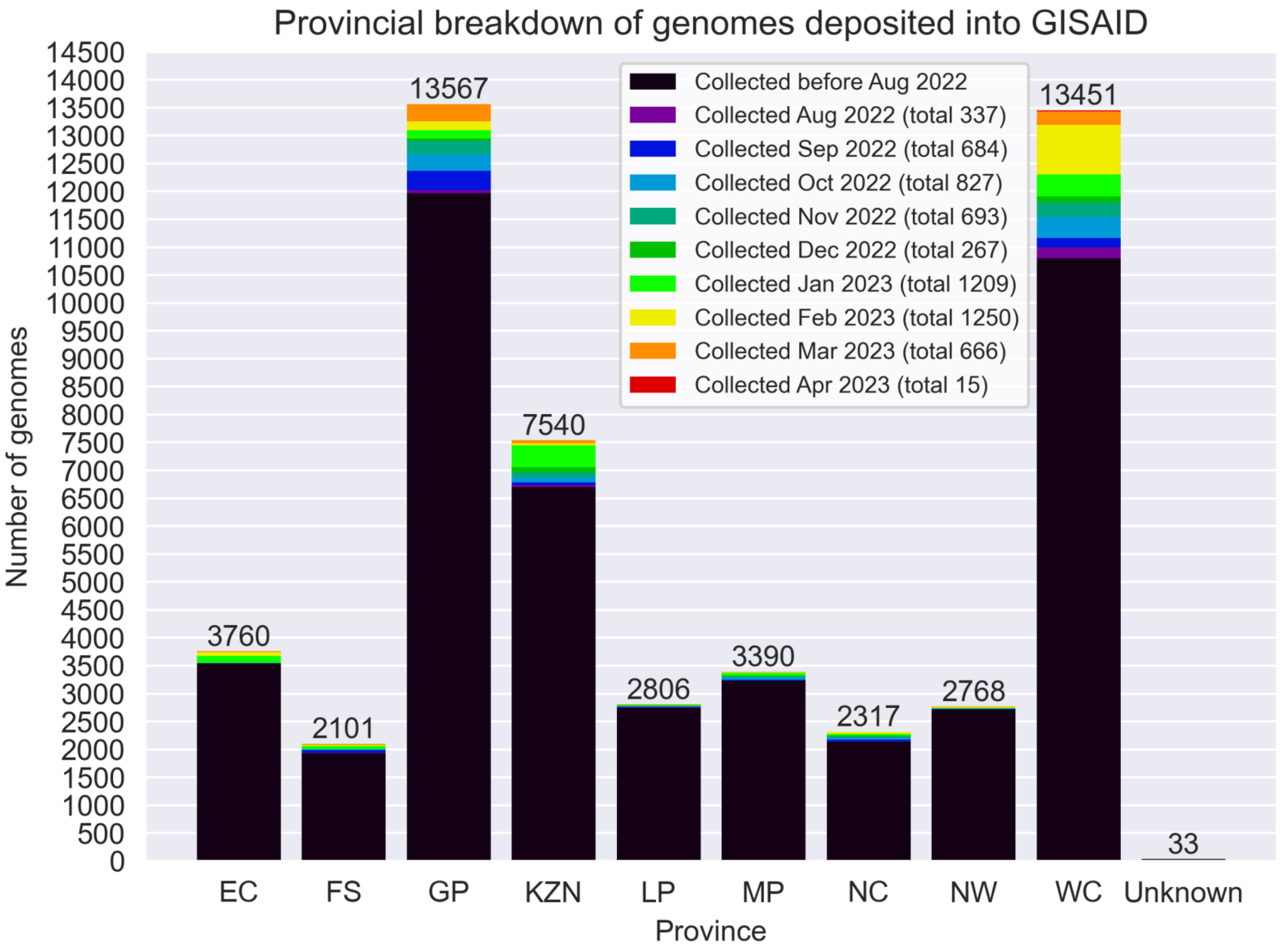
Currently in epi week 18 (ending 6 May 2023)

Total genomes: 51 733
2020 genomes: 6 795
2021 genomes: 26 440
2022 genomes: 15 358
2023 genomes: 3 140
Genomes added since last report: 71

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

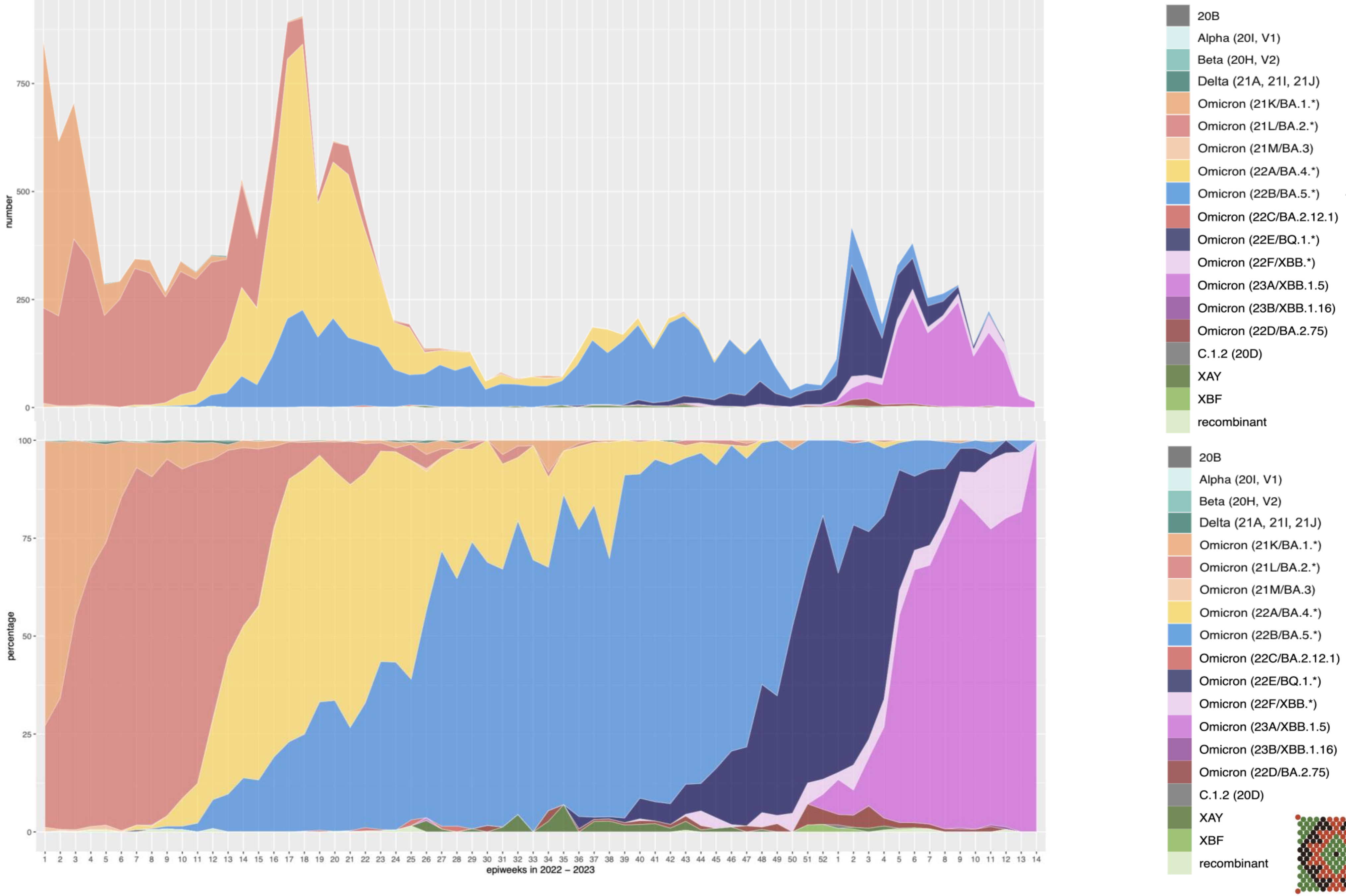
NGS-SA
Network for Genomic Surveillance in South Africa

GISAID genomes vs total cases, 2020 – 2023 (N= 51 733)



NGS-SA
Network for Genomic
Surveillance in South Africa

Number and percentage of clades by epiweek in South Africa, 2022-2023 (18 455*)



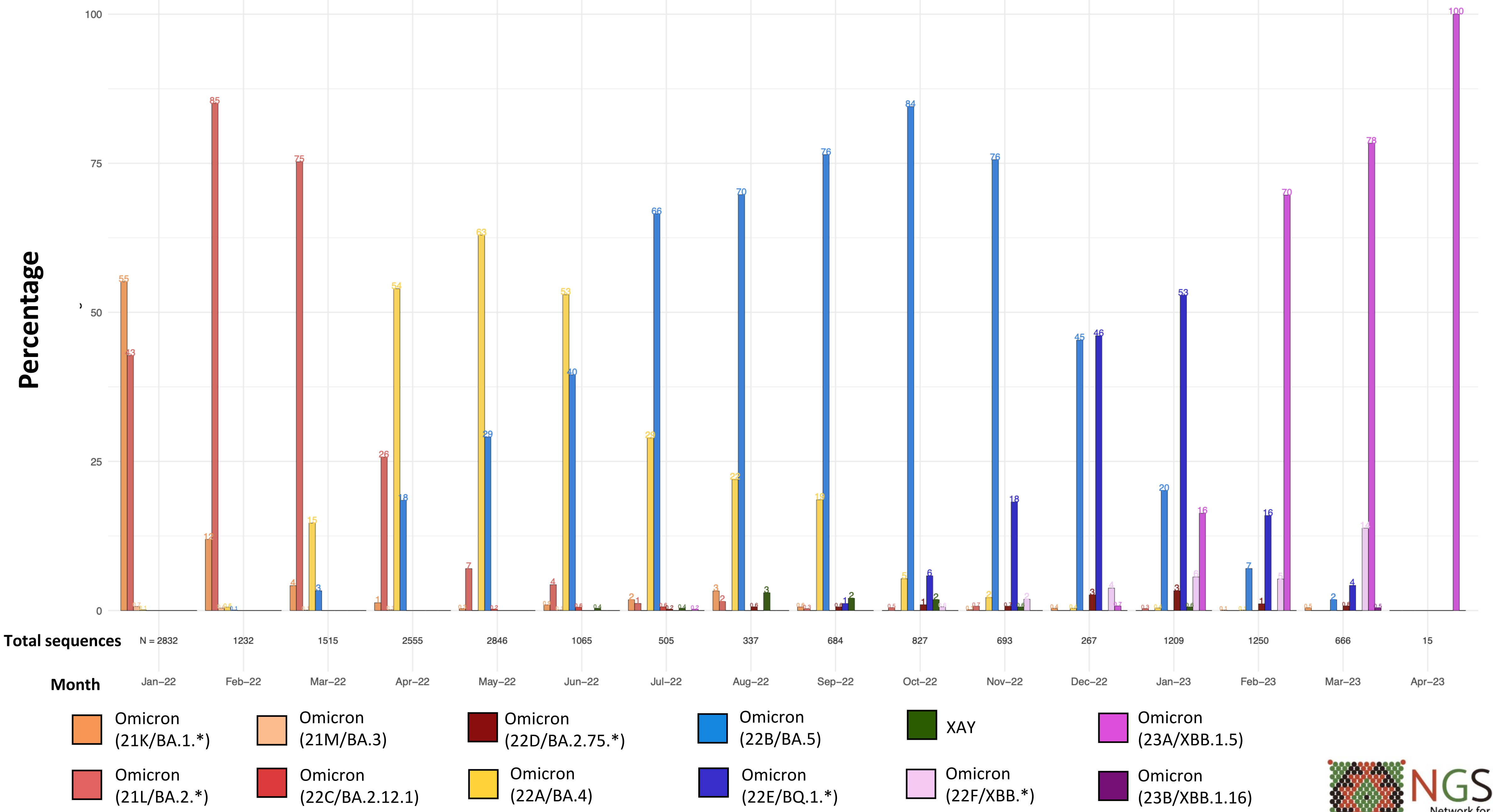
Sequencing data ending epi week 14 (ending 8 Apr 2023)

Currently in epi week 18 (ending 6 May 2023)

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

NGS-SA
Network for Genomic Surveillance in South Africa

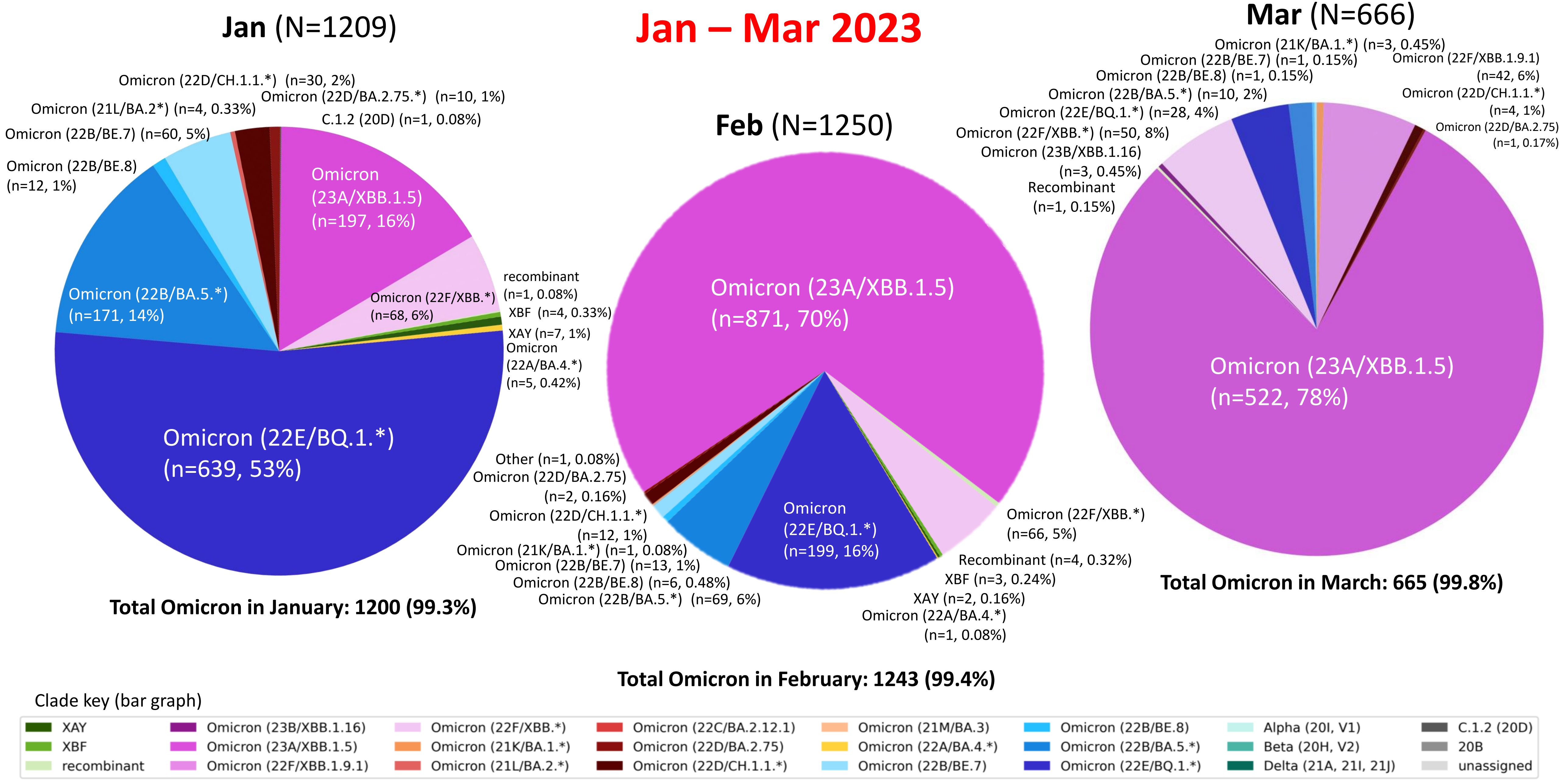
Detection Rates: Omicron and recombinants



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

NGS-SA
Network for Genomic Surveillance in South Africa

Prevalence of Variants of Concern (VOC) and Variants of Interest (VOI) in Jan – Mar 2023



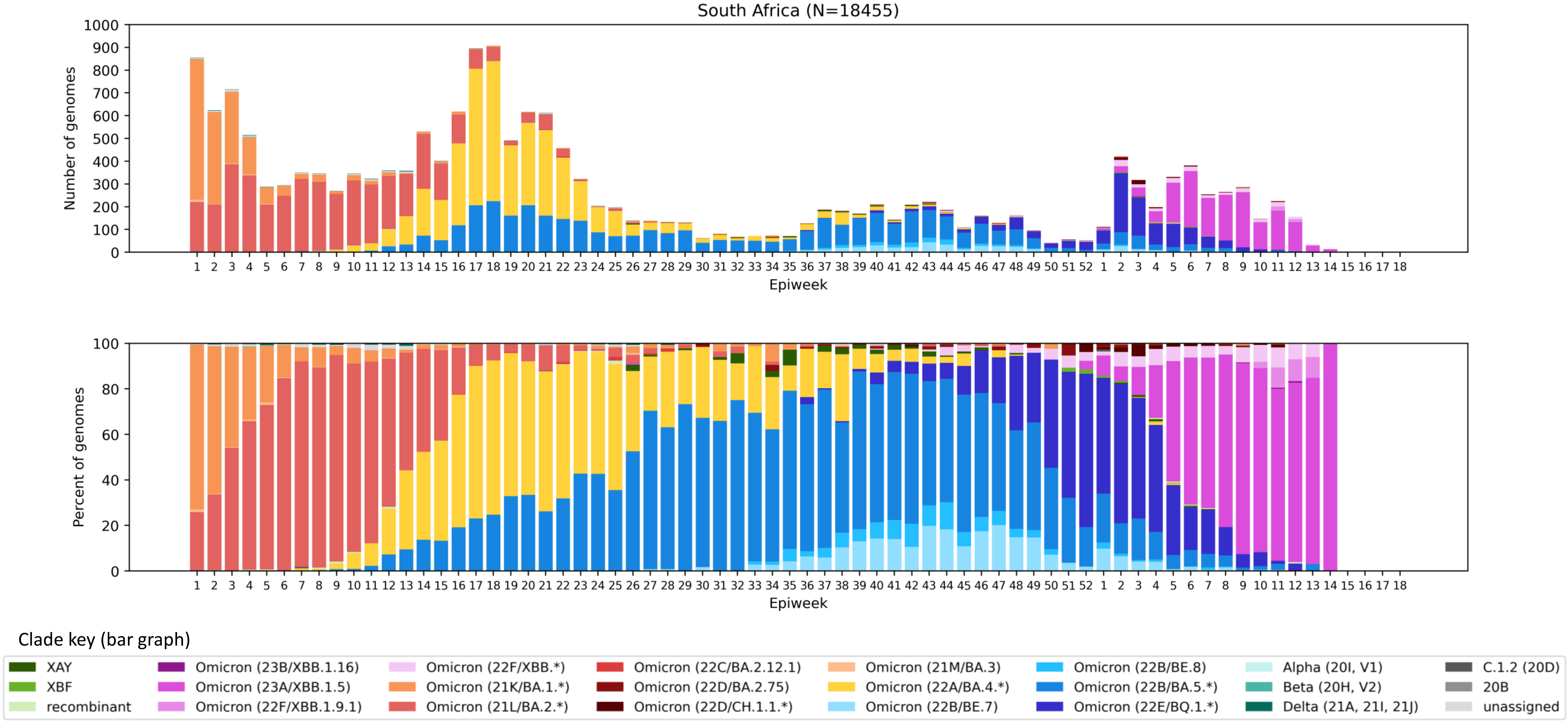
Total Omicron in January: 1200 (99.3%)

Total Omicron in February: 1243 (99.4%)

Total Omicron in March: 665 (99.8%)

Note: XBF is an Omicron-Omicron recombinant and so is counted in the total number of Omicrons.

South Africa, 2022-2023, n = 18 455*



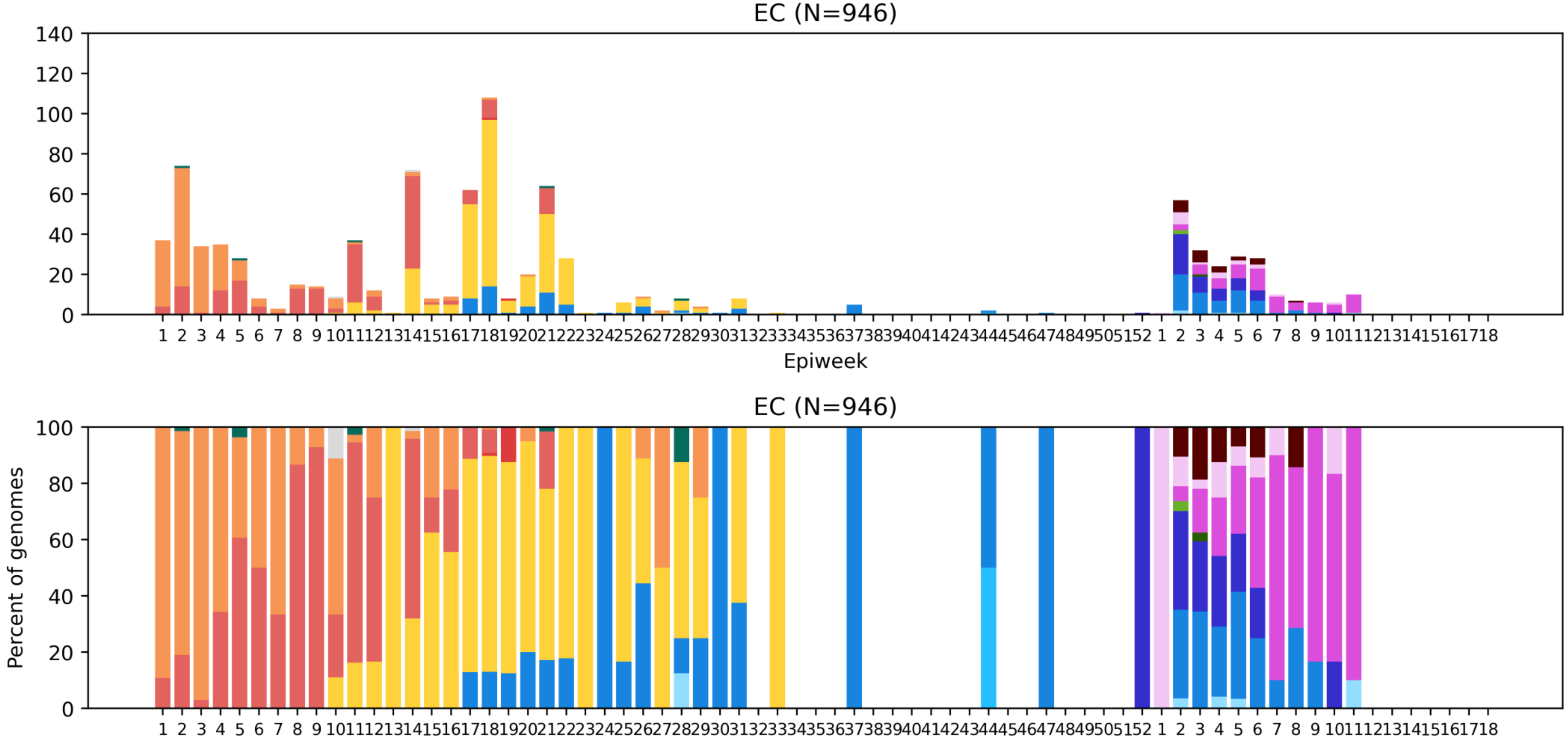
*Excludes sequences missing collection dates. Lineages of particular interest (mainly WHO Omicron subvariants under monitoring) are separate from the main clade groupings.

#Recombinants include all recombinant lineages (viruses consisting of segments of two different lineages) detected in South Africa at low levels. Currently it consists of XT, XAS, XAZ, XBA, XBF.

NGS-SA
Network for Genomic Surveillance in South Africa

Eastern Cape Province, 2022-2023, n = 946

Genomes added since last report: 0*



Clade key (bar graph)

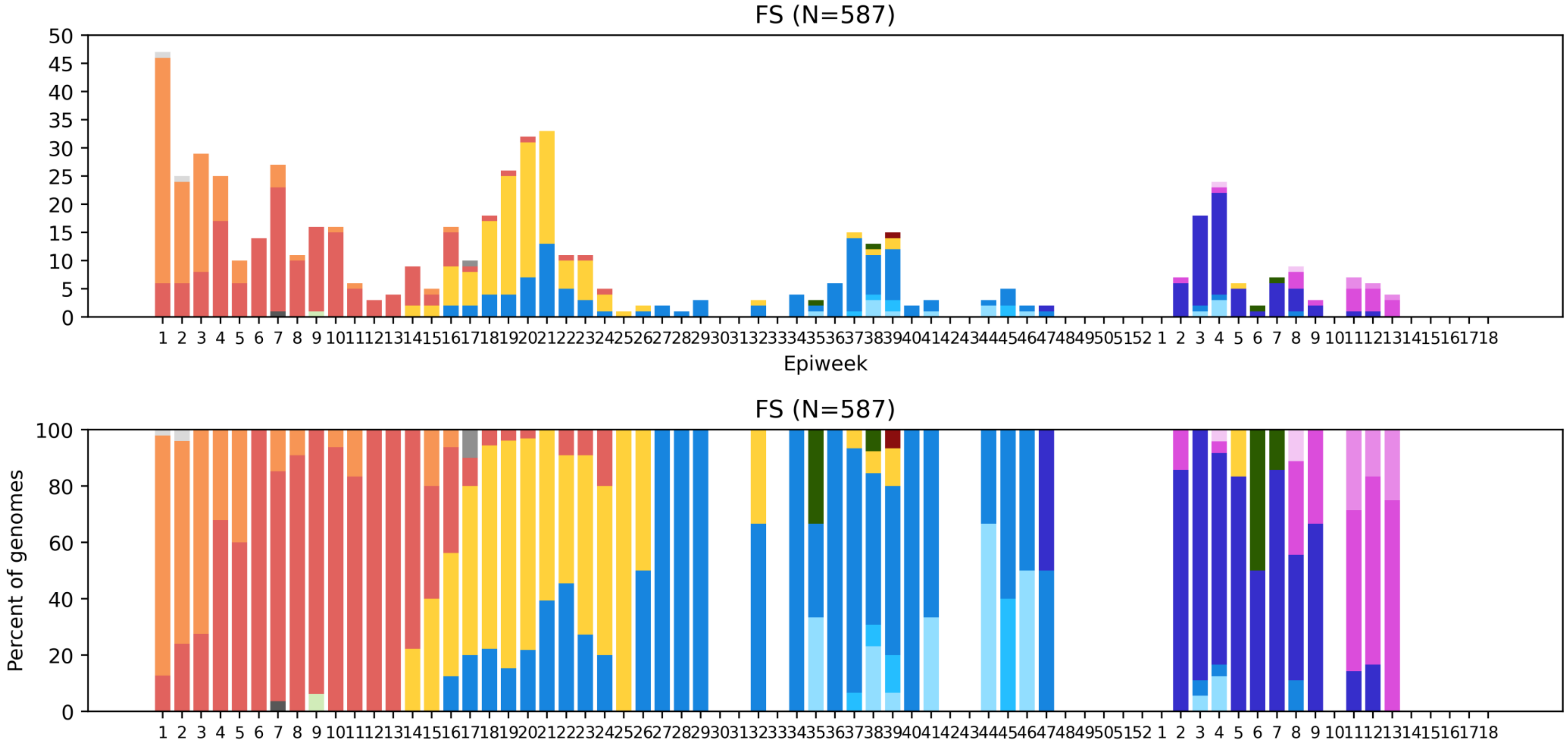
- 🟩 XAY
- 🟩 XBF
- 🟩 recombinant
- 🟪 Omicron (23B/XBB.1.16)
- 🟪 Omicron (23A/XBB.1.5)
- 🟪 Omicron (22F/XBB.1.9.1)
- 🟪 Omicron (22F/XBB.*)
- 🟧 Omicron (21K/BA.1.*)
- 🟥 Omicron (21L/BA.2.*)
- 🟥 Omicron (22C/BA.2.12.1)
- 🟥 Omicron (22D/BA.2.75)
- 🟫 Omicron (22D/CH.1.1.*)
- 🟧 Omicron (21M/BA.3)
- 🟨 Omicron (22A/BA.4.*)
- 🟦 Omicron (22B/BE.7)
- 🟦 Omicron (22B/BE.8)
- 🟦 Omicron (22B/BA.5.*)
- 🟦 Omicron (22E/BQ.1.*)
- 🟩 Alpha (20I, V1)
- 🟩 Beta (20H, V2)
- 🟩 Delta (21A, 21I, 21J)
- ⬛ C.1.2 (20D)
- ⬛ 20B
- ⬜ unassigned

NGS-SA
Network for Genomic
Surveillance in South Africa

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

Free State Province, 2022-2023, n = 587

Genomes added since last report: 0*



Clade key (bar graph)

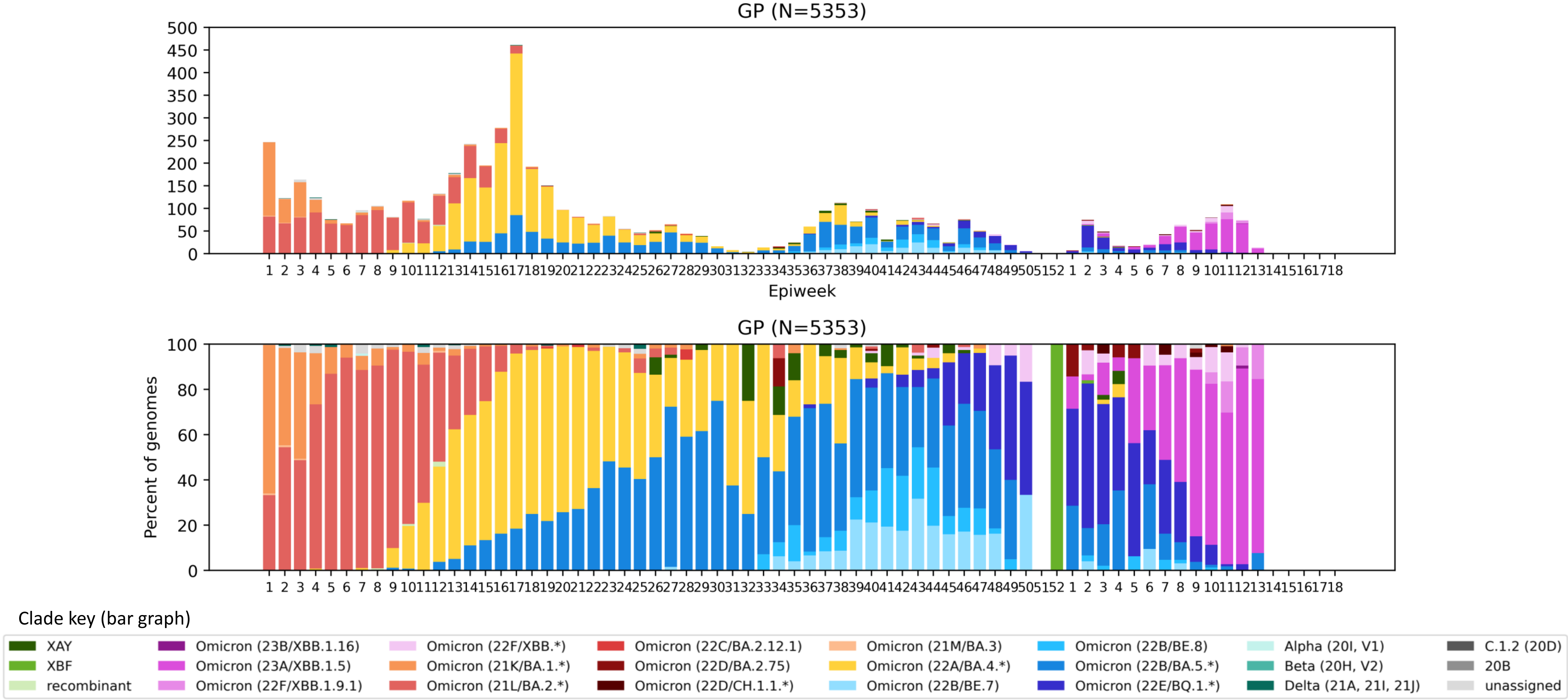
- 🟩 XAY
- 🟩 XBF
- 🟩 recombinant
- 🟪 Omicron (23B/XBB.1.16)
- 🟪 Omicron (23A/XBB.1.5)
- 🟪 Omicron (22F/XBB.1.9.1)
- 🟪 Omicron (22F/XBB.*)
- 🟧 Omicron (21K/BA.1.*)
- 🟥 Omicron (21L/BA.2.*)
- 🟥 Omicron (22C/BA.2.12.1)
- 🟥 Omicron (22D/BA.2.75)
- 🟫 Omicron (22D/CH.1.1.*)
- 🟧 Omicron (21M/BA.3)
- 🟨 Omicron (22A/BA.4.*)
- 🟦 Omicron (22B/BE.7)
- 🟦 Omicron (22B/BE.8)
- 🟦 Omicron (22B/BA.5.*)
- 🟦 Omicron (22E/BQ.1.*)
- 🟩 Alpha (20I, V1)
- 🟩 Beta (20H, V2)
- 🟩 Delta (21A, 21I, 21J)
- ⬛ C.1.2 (20D)
- ⬛ 20B
- ⬜ unassigned

NGS-SA
Network for Genomic
Surveillance in South Africa

*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 = 5353

Genomes added since last report: 11*

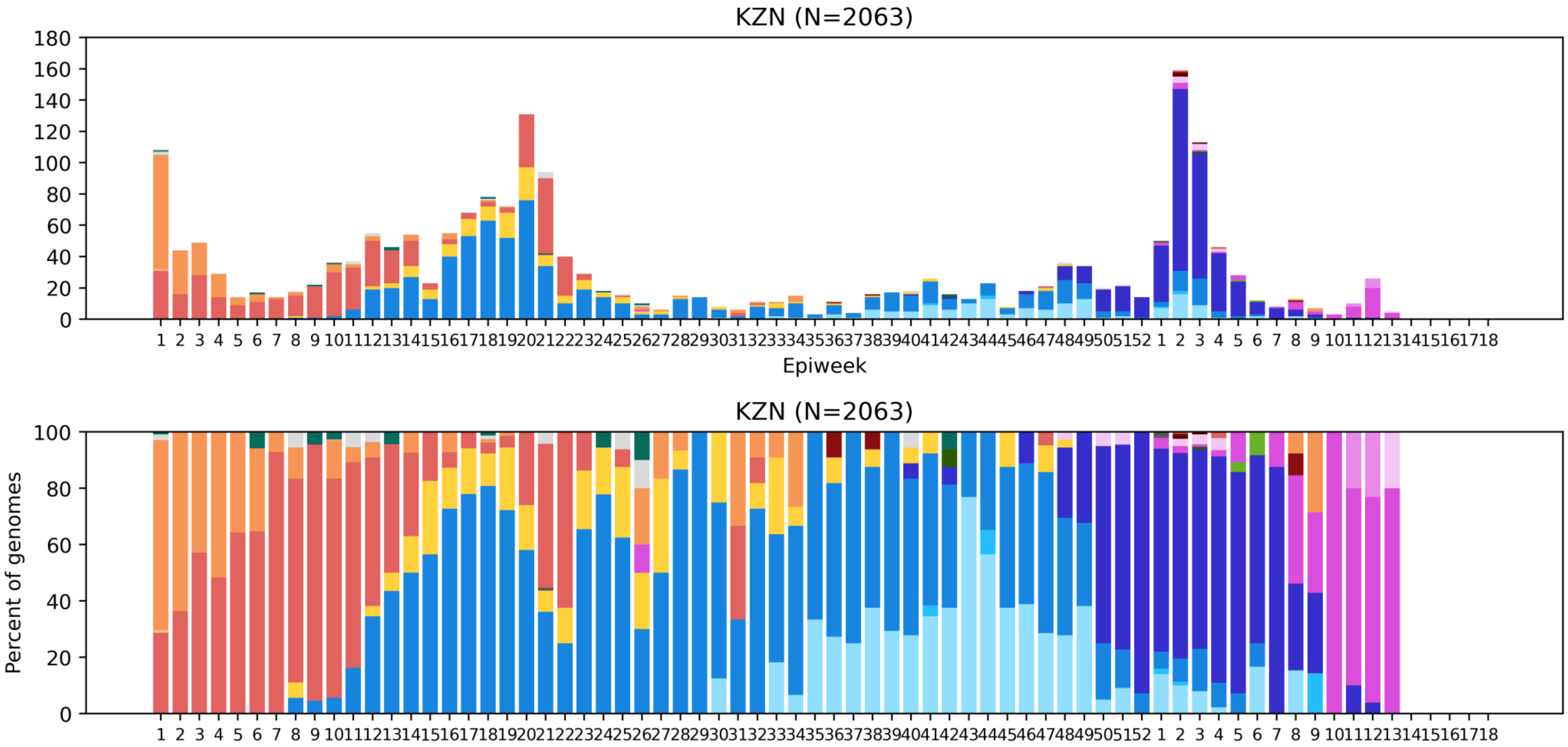


NGS-SA
Network for Genomic Surveillance in South Africa

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

KwaZulu-Natal Province, 2022-2023, n = 2063

Genomes added since last report: 0*



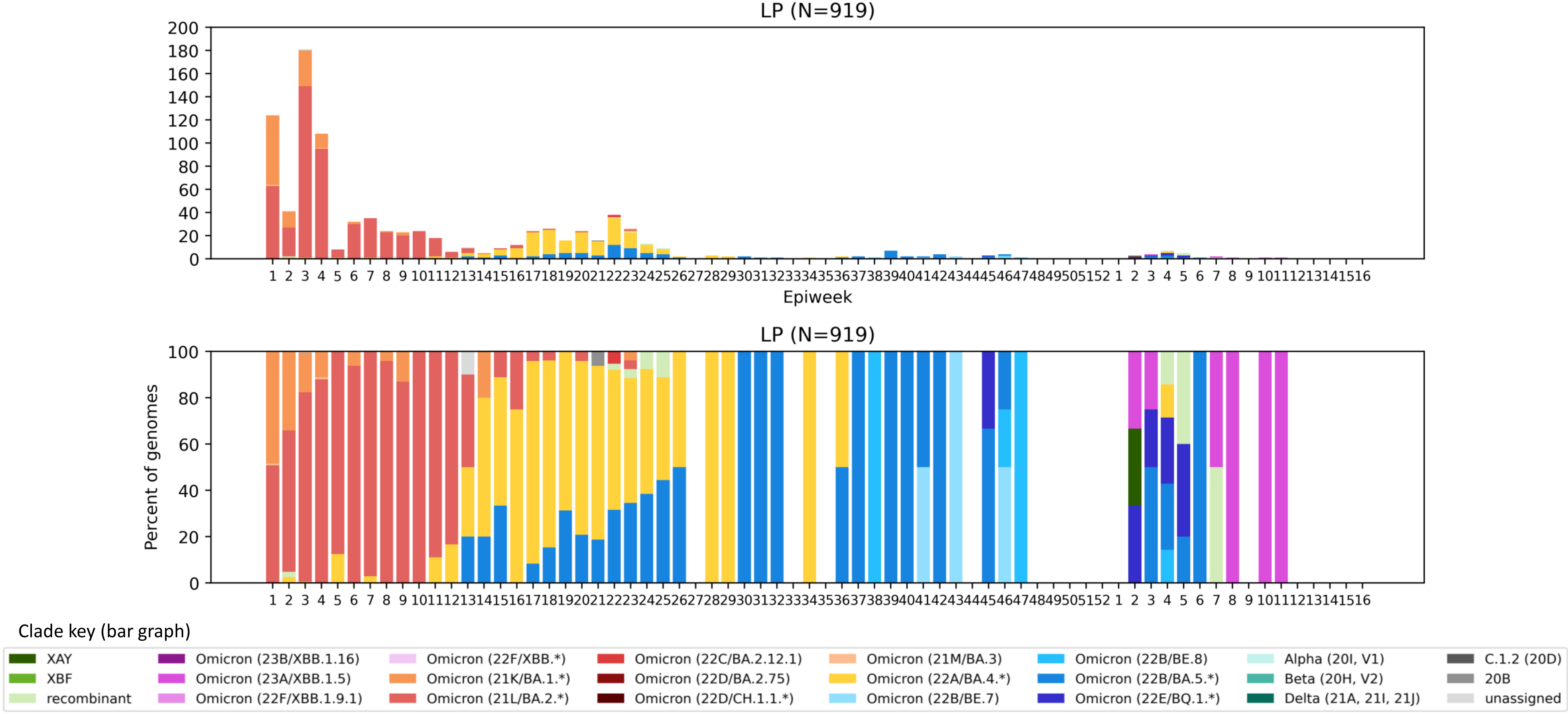
Clade key (bar graph)

XAY	Omicron (23B/XBB.1.16)	Omicron (22F/XBB.*)	Omicron (22C/BA.2.12.1)	Omicron (21M/BA.3)	Omicron (22B/BE.8)	Alpha (20I, V1)	C.1.2 (20D)
XBF	Omicron (23A/XBB.1.5)	Omicron (21K/BA.1.*)	Omicron (22D/BA.2.75)	Omicron (22A/BA.4.*)	Omicron (22B/BA.5.*)	Beta (20H, V2)	20B
recombinant	Omicron (22F/XBB.1.9.1)	Omicron (21L/BA.2.*)	Omicron (22D/CH.1.1.*)	Omicron (22B/BE.7)	Omicron (22E/BQ.1.*)	Delta (21A, 21I, 21J)	unassigned

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

Limpopo Province, 2022-2023, n = 919

Genomes added since last report: 0*


LP (N=919)

LP (N=919)

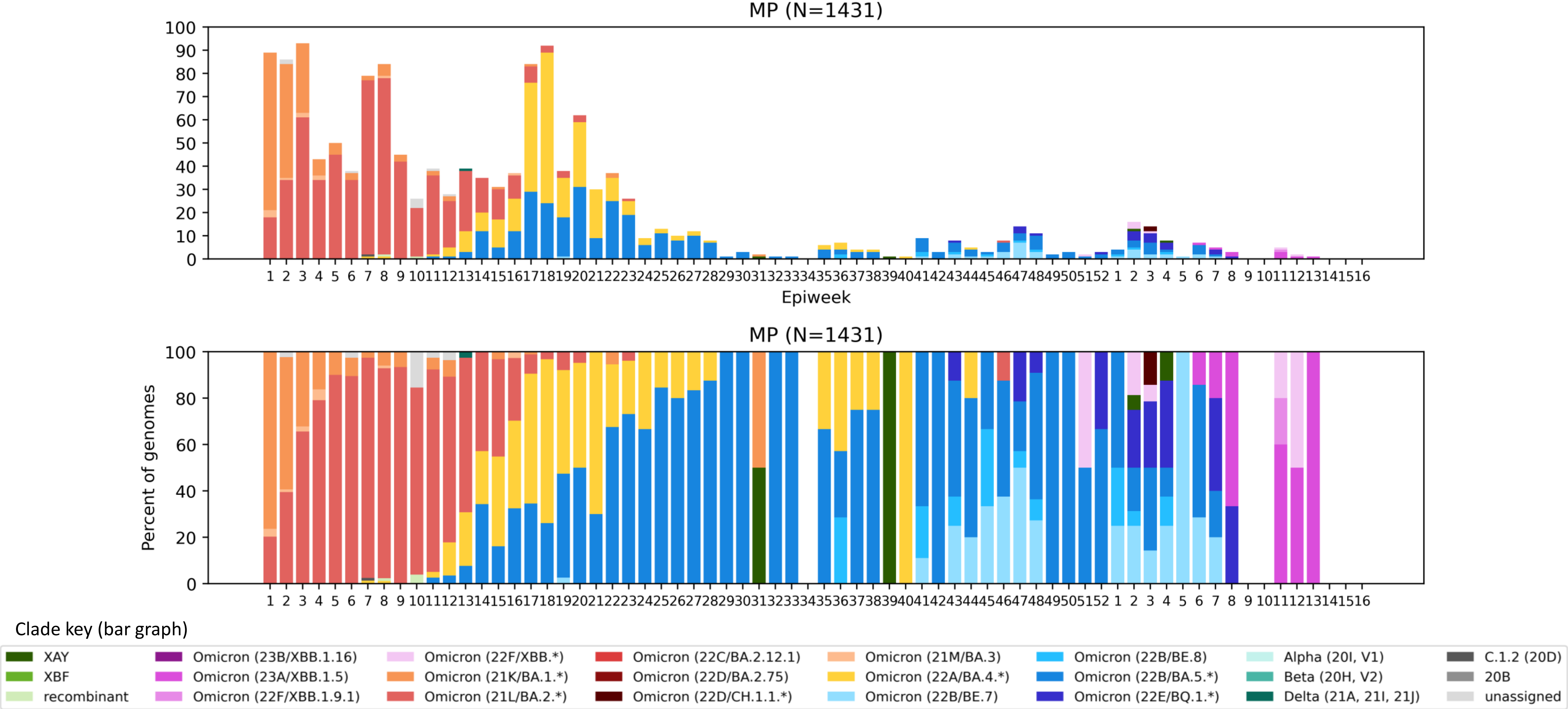
Clade key (bar graph)

NGS-SA
Network for Genomic Surveillance in South Africa

*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 = 1431

Genomes added since last report: 0*

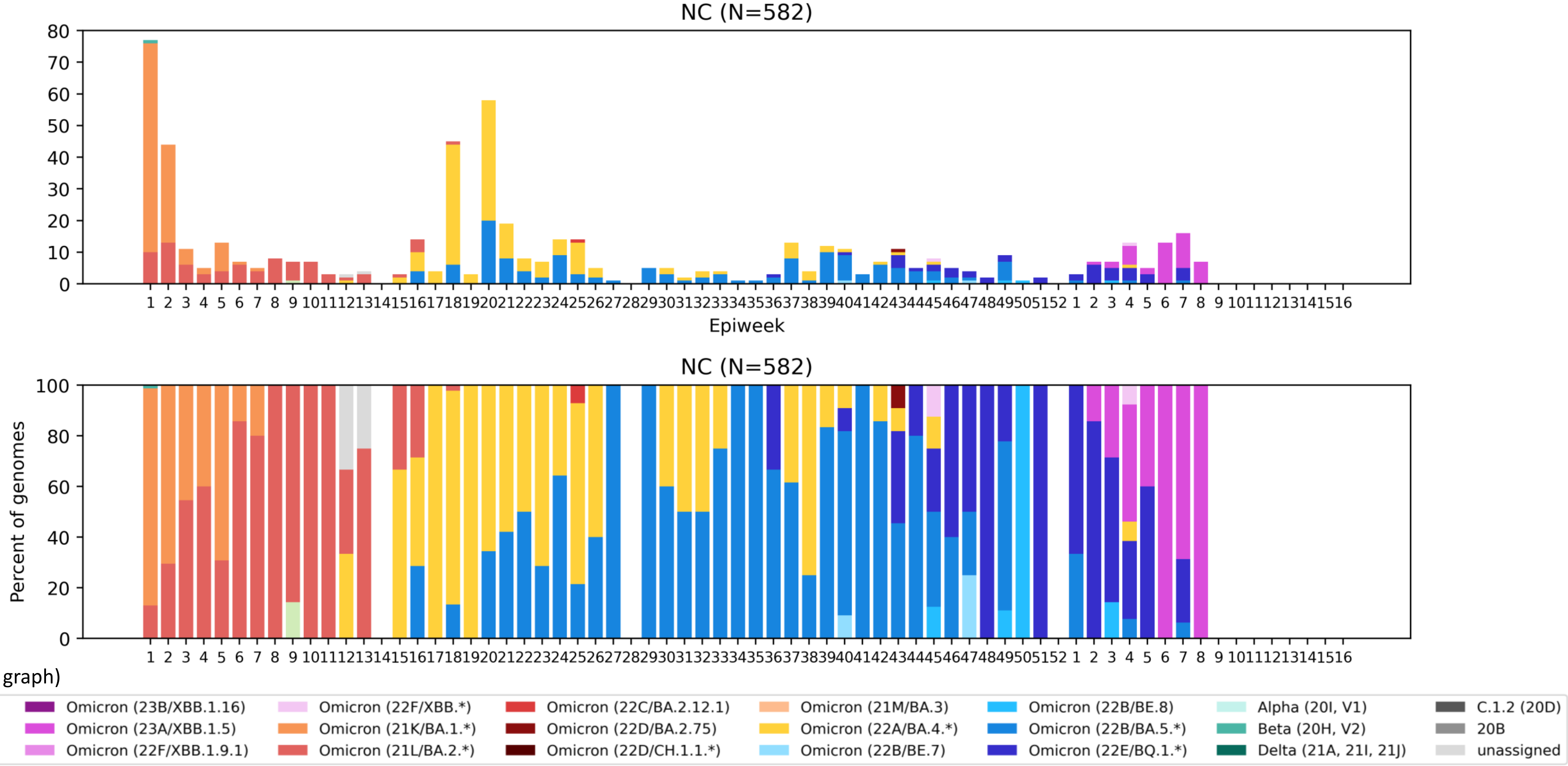


NGS-SA
Network for Genomic
Surveillance in South Africa

*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 = 582

Genomes added since last report: 0*

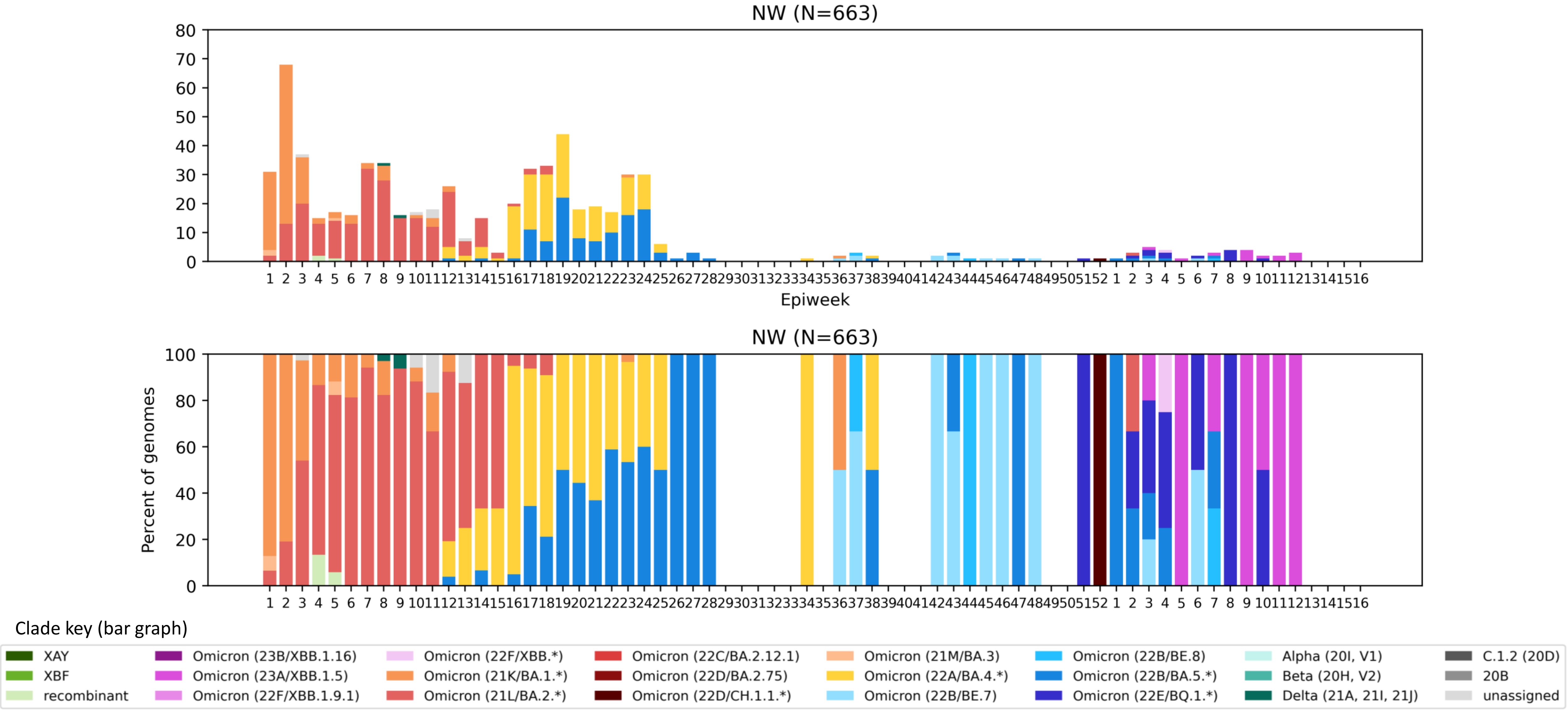


NGS-SA
Network for Genomic
Surveillance in South Africa

*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 = 663

Genomes added since last report: 0*

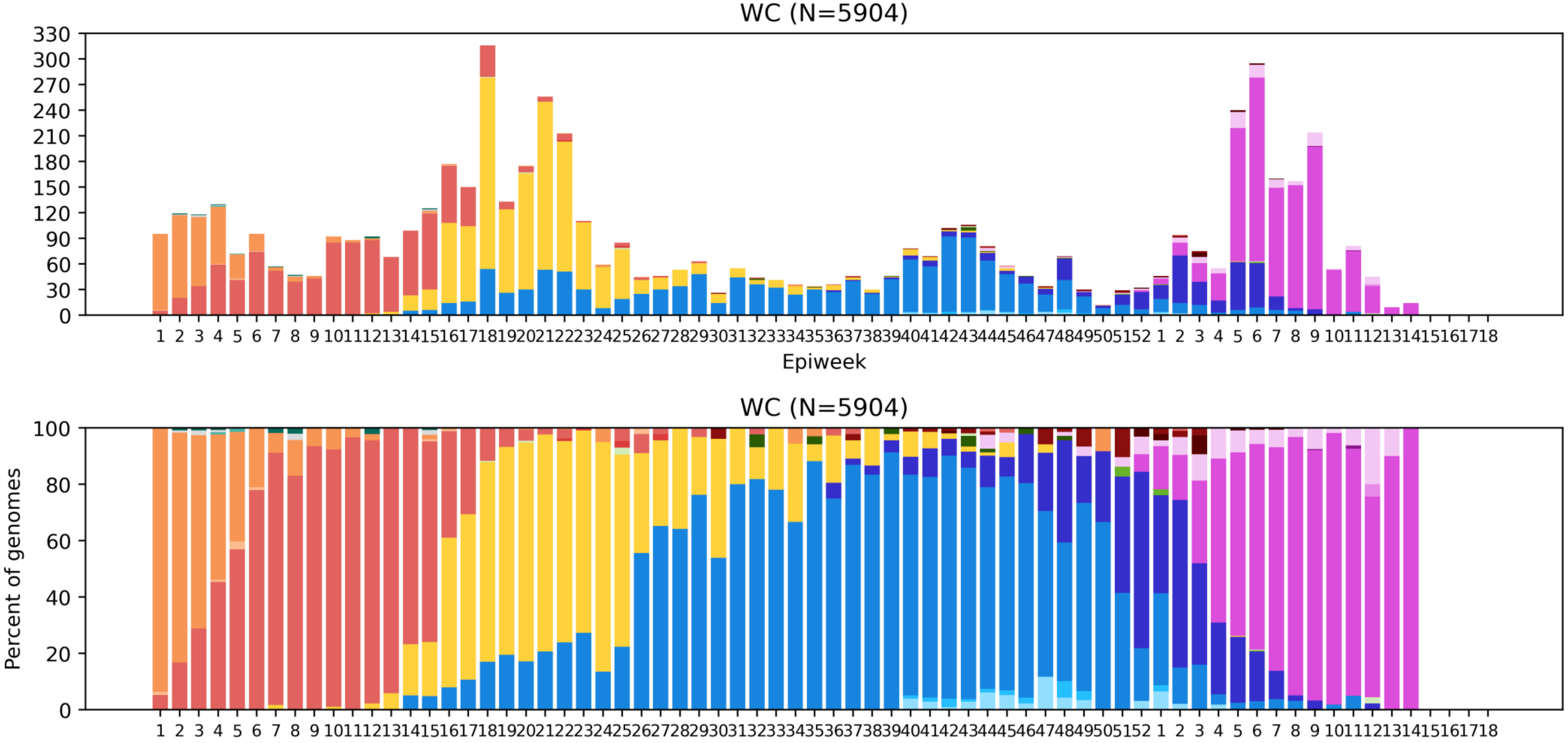


NGS-SA
Network for Genomic Surveillance in South Africa

*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 = 5904

Genomes added since last report: 60*



Clade key (bar graph)

- ■ XAY
- ■ XBF
- ■ recombinant
- ■ Omicron (23B/XBB.1.16)
- ■ Omicron (23A/XBB.1.5)
- ■ Omicron (22F/XBB.1.9.1)
- ■ Omicron (22F/XBB.*)
- ■ Omicron (21K/BA.1.*)
- ■ Omicron (21L/BA.2.*)
- ■ Omicron (22C/BA.2.12.1)
- ■ Omicron (22D/BA.2.75)
- ■ Omicron (22D/CH.1.1.*)
- ■ Omicron (21M/BA.3)
- ■ Omicron (22A/BA.4.*)
- ■ Omicron (22B/BE.7)
- ■ Omicron (22B/BE.8)
- ■ Omicron (22B/BA.5.*)
- ■ Omicron (22E/BQ.1.*)
- ■ Alpha (20I, V1)
- ■ Beta (20H, V2)
- ■ Delta (21A, 21I, 21J)
- ■ C.1.2 (20D)
- ■ 20B
- ■ unassigned

NGS-SA
Network for Genomic
Surveillance in South Africa

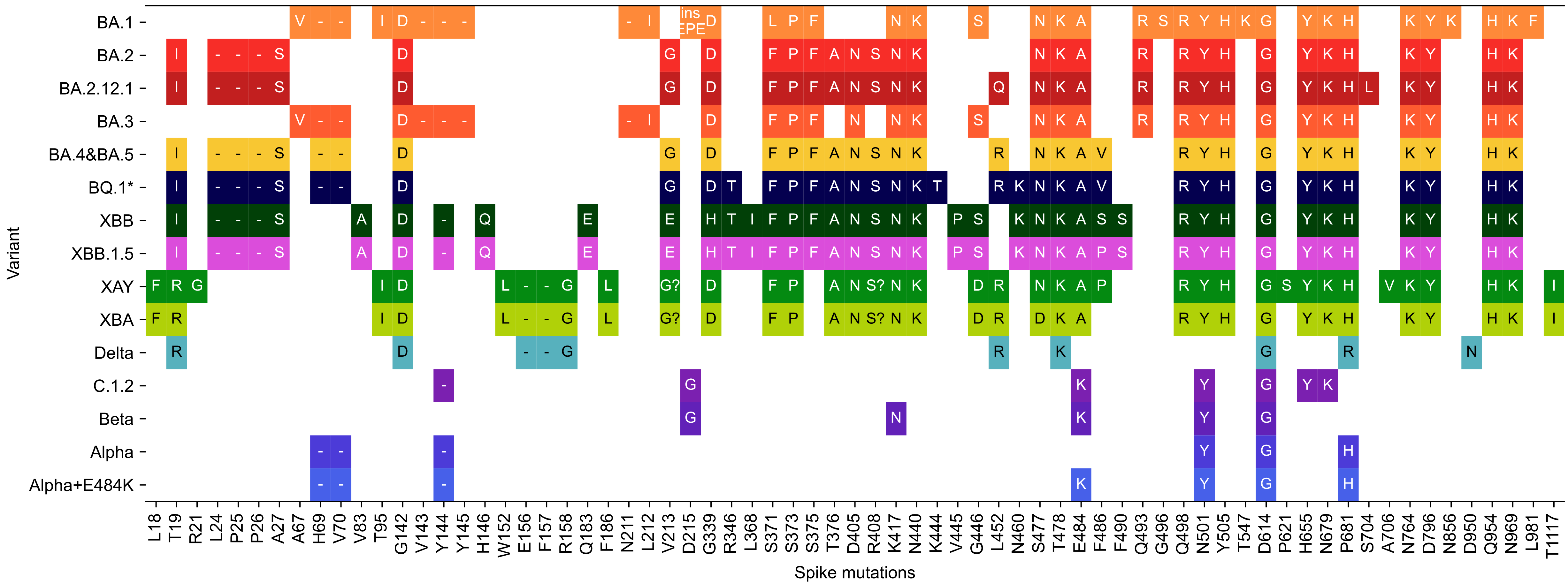
*May include genomes from 2020 and 2021 which are not pictured here and are not included in the slide total.

Summary

- **Sequencing update**
  - All provinces have sequences for January and February 2023. March sequences are from all provinces, except the Northern Cape. April sequence data is from the Western Cape

- **Variant of Concern Omicron in South Africa**
  - Omicron continued to dominate in January (99.3%), February (99.4%) and March (99.8%)
  - BQ.1 and sub-lineages were the dominant Omicron lineage in December (46%) and January (53%)
  - XBB.1.5 was detected in December 2022 (0.8%) and January 2023 (16%), and was the dominant lineage in February (70%) and March (78%). It is currently dominant in April (100%), although this is based on low numbers (n=15)
  - XBB.1.16 has been detected at a low prevalence in March (<1%) in Gauteng (n=1) and the Western Cape (n=2)
  - BA.2.75.* continued to be detected at a low prevalence in January through March (≤1%)

Spike protein mutation* profile of Variants of Interest and Concern



Variant	L18	T19	R21	L24	P25	P26	A27	A67	H69	V70	V83	T95	G142	V143	Y144	Y145	H146	W152	E156	F157	R158	Q183	F186
BA.1								V	-	-		I	D	-	-	-							
BA.2		I		-	-	-	S						D										
BA.2.12.1		I		-	-	-	S						D										
BA.3								V	-	-			D	-	-	-							
BA.4&BA.5		I		-	-	-	S		-	-			D										
BQ.1*		I		-	-	-	S		-	-			D										
XBB		I		-	-	-	S				A		D		-		Q					E	
XBB.1.5		I		-	-	-	S				A		D		-		Q					E	
XAY	F	R	G									I	D					L	-	-	G		L
XBA	F	R										I	D					L	-	-	G		L
Delta		R											D						-	-	G		
C.1.2															-								
Beta																							
Alpha									-	-					-								
Alpha+E484K									-	-					-								

Variant	N211	L212	V213	D215	G339	R346	L368	S371	S373	S375	T376	D405	R408	K417	N440	K444	V445	G446	L452	N460	S477	T478	E484
BA.1	-	I		ins EPE	D			L	P	F				N	K			S			N	K	A
BA.2			G		D			F	P	F	A	N	S	N	K						N	K	A
BA.2.12.1			G		D			F	P	F	A	N	S	N	K				Q		N	K	A
BA.3	-	I			D			F	P	F		N		N	K			S			N	K	A
BA.4&BA.5			G		D			F	P	F	A	N	S	N	K				R		N	K	A
BQ.1*			G		D	T		F	P	F	A	N	S	N	K	T			R	K	N	K	A
XBB			E		H	T	I	F	P	F	A	N	S	N	K		P	S		K	N	K	A
XBB.1.5			E		H	T	I	F	P	F	A	N	S	N	K		P	S		K	N	K	A
XAY			G?		D			F	P		A	N	S?	N	K			D	R		N	K	A
XBA			G?		D			F	P		A	N	S?	N	K			D	R		D	K	A
Delta																			R			K	
C.1.2				G																			K
Beta				G										N									K
Alpha																							
Alpha+E484K																							K

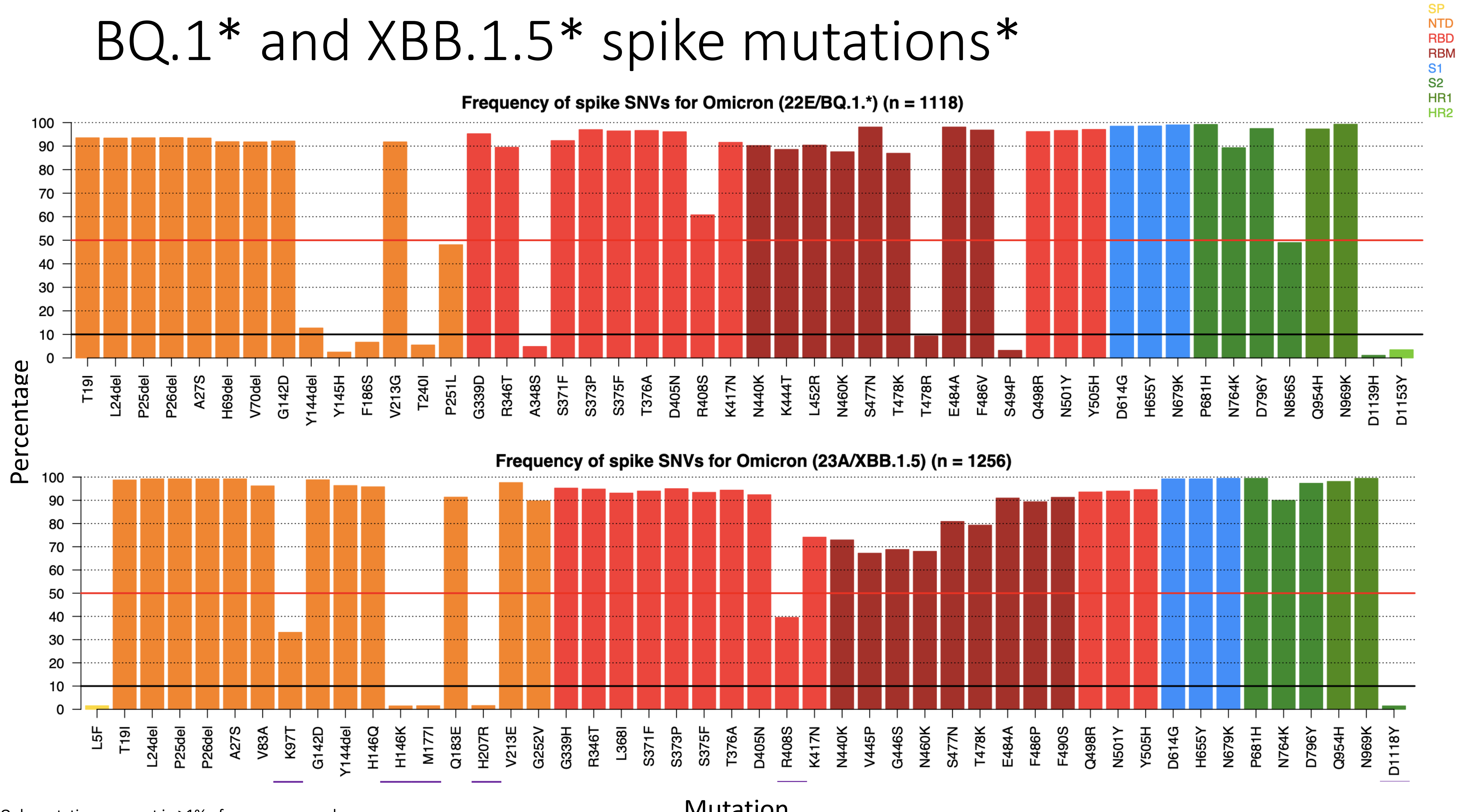
Variant	F486	F490	Q493	G496	Q498	N501	Y505	T547	D614	P621	H655	N679	P681	S704	A706	N764	D796	N856	D950	Q954	N969	L981	T1117
BA.1			R	S	R	Y	H	K	G		Y	K	H			K	Y	K		H	K	F	
BA.2			R		R	Y	H		G		Y	K	H			K	Y			H	K		
BA.2.12.1			R		R	Y	H		G		Y	K	H	L		K	Y			H	K		
BA.3			R		R	Y	H		G		Y	K	H			K	Y			H	K		
BA.4&BA.5	V				R	Y	H		G		Y	K	H			K	Y			H	K		
BQ.1*	V				R	Y	H		G		Y	K	H			K	Y			H	K		
XBB	S	S			R	Y	H		G		Y	K	H			K	Y			H	K		
XBB.1.5	P	S			R	Y	H		G		Y	K	H			K	Y			H	K		
XAY	P				R	Y	H		G	S	Y	K	H		V	K	Y			H	K		I
XBA					R	Y	H		G		Y	K	H			K	Y			H	K		I
Delta									G				R						N				
C.1.2						Y			G		Y	K											
Beta						Y			G														
Alpha						Y			G				H										
Alpha+E484K						Y			G				H										

- 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 recombinant sequences are pictured

NGS-SA
Network for Genomic Surveillance in South Africa

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



*Only mutations present in ≥1% of sequences are shown.

University of Stellenbosch & NHLS Tygerberg Virology

Susan Engelbrecht
Wolfgang Preiser
Gert van Zyl
Tongai Maponga
Bronwyn Kleinhans
Shannon Wilson
Karabo Phadu
Tania Stander
Kamela Mahlakwane
Mathilda Claassen
Diagnostic laboratory staff

UKZN-Inkosi Albert Luthuli Central Hospital

Dr Khanyi Msomi
Dr Kerusha Govender
Dr Pravi Moodley
Dr Aabida Khan
Dr Lili Gounder
Dr Kerri Francois
Dr Cherise Naicker
Dr Joedene Chetty

Dr Neli Ngcaba
Dr Tshepiso Mosito
Mr Malcolm Ellapen
Mr Kubendran Reddy
The COVID-19 Bench team

University of KwaZulu-Natal & Africa Health Research Institute

KRISP at UKZN:
Tulio de Oliveira
Richard Lessels
Houriiyah Tegally
Eduan Wilkinson
Jennifer Giandhari
Sureshnee Pillay
Emmanuel James San

AHRI
Alex Sigal
Sandile Cele
Willem Hanekom

University of Cape Town, NHLS & Western Cape Government

NHLS-UCT
Carolyn Williamson
Nei-yuan Hsiao
Diana Hardie
Kruger Marais
Stephen Korsman
Ziyaad Valley-Omar

WCG-UCT
Mary-Anne Davies
Hannah Hussey
Andrew Boulle
Masudah Paleker
Theuns Jacobs
Erna Morden

NHLS Greenpoint
Annabel Enoch

This project has received funding from the European Union's Horizon Europe Research and Innovation Actions under grant No. 101046041

UCT, IDM and CIDRI-Africa
Deelan Doolabh
Arash Iranzadeh
Lynn Tyers
Innocent Mudau
Nokuzola Mbhele
Fezokuhle Khumalo
Thabang Serakge
Bruna Galvão
Arghavan Alisoltani (U. California)

Robert Wilkinson
Darren Martin
Nicola Mulder
Wendy Burgers
Ntobeko Ntusi
Rageema Joseph
Sean Wasserman
Linda Boloko

Zoonotic arbo and respiratory virus program Centre for Viral Zoonoses Department Medical Virology/ NHLS Tshwane Academic division University of Pretoria

ZARV research program/UP
Marietjie Venter (Head: ZARV)
Adriano Mendes (Postdoc)
Amy Strydom (Postdoc)
Michaela Davis (MSc, intern medical scientist)
Carien van Niekerk

NHLS Tshwane
Prof Simnikiwe Mayaphi (HOD)

Funders:
GIZ/BMBF: African Network for Improved diagnostics and epidemiology of common and emerging infectious agents (ANDEMIA)
G7 Global Health fund, Robert Koch Institute, Dr Fabian Leendertz

National Institute for Communicable Diseases

Centre for Respiratory Diseases & Meningitis
Anne von Gottberg
Thabo Mohale
Daniel Amoako
Josie Everatt
Boitshoko Mahlangu
Noxolo Ntuli
Anele Mnguni
Amelia Buys
Cardia Fourie
Noluthando Duma
Linda de Gouveia
Jackie Kleynhans
Nicole Wolter
Sibongile Walaza
Mignon du Plessis
Stefano Tempia
Mvuyo Makhasi
Cheryl Cohen

Centre for HIV and STIs
Jinal Bhiman
Cathrine Scheepers
Constantinos Kurt Wibmer
Thandeka Moyo
Tandile Hermanus
Frances Ayres
Zanele Molaudzi
Bronwen Lambson
Tandile Hermanus
Mashudu Madzivhandila
Prudence Kgagudi
Brent Oosthuysen
Penny Moore
Lynn Morris

NICD Groups
NICD COVID-19 response team
NICD SARS-CoV-2 Sequencing Group

Sequencing Core Facility
Zamantungwa Khumalo
Annie Chan
Morne du Plessis
Stanford Kwenda
Phillip Senzo Mtshali
Mushal Allam
Florah Mnyameni
Arshad Ismail

University of the Free State

UFS
Dominique Goedhals
Armand Bester
Martin Myaga
Peter Mwangi
Emmanuel Ogunbayo
Milton Mogotsi
Makgotso Maotoana
Lutfiyya Mohamed

NHLS Division of Virology
Sabeehah Vawda
Felicity Burt
Thokozani Mkhize
Diagnostic laboratory staff

Additional support and collaborators

NHLS
Koeleka Mlisana
Zinhle Makatini
Eugene Elliot
Florette K. Treurnicht
Kathleen Subramoney
Oluwakemi Laguda-Akingba
Shareef Abrahams
Greta Hoyland
Gloria Selabe
Elias Bereda
Jeannette Wadula

Hyrax Biosciences
Simon Travers

Cape Town HVTN Laboratory
Erica Anderson-Nissen
Anneta Naidoo

Ndlovu Research
Hugo Tempelman
CJ Umunnakwe

Lancet
Allison J. Glass
Raquel Viana

Ampath
Terry Marshall
Cindy van Deventer
Eddie Silberbauer

Pathcare Vermaak
Andries Dreyer
Howard Newman
Riaan Writes
Marianne Wolfaardt
Warren Lowman

Bridge-the-Gap
Raymond Rott

Cytespace Africa Laboratories
Christa Viljoen

ARC-OVI
Lia Rotherham

CAPRISA
Salim Abdool Karim
Nigel Garret

UKZN - Big Data
Francesco Pettruccione
Ilya Sinayskiy

University of Oxford
José Lourenço

FioCruz, Brazil
Vagner Fonseca
Marta Giovanetti
Luiz Carlos Junior Alcantara

Africa CDC and Africa PGI
John Nkengasong
Sofonias Tessema

Netcare
Richard Friedland
Craig Murphy
Caroline Maslo
Liza Sitharam

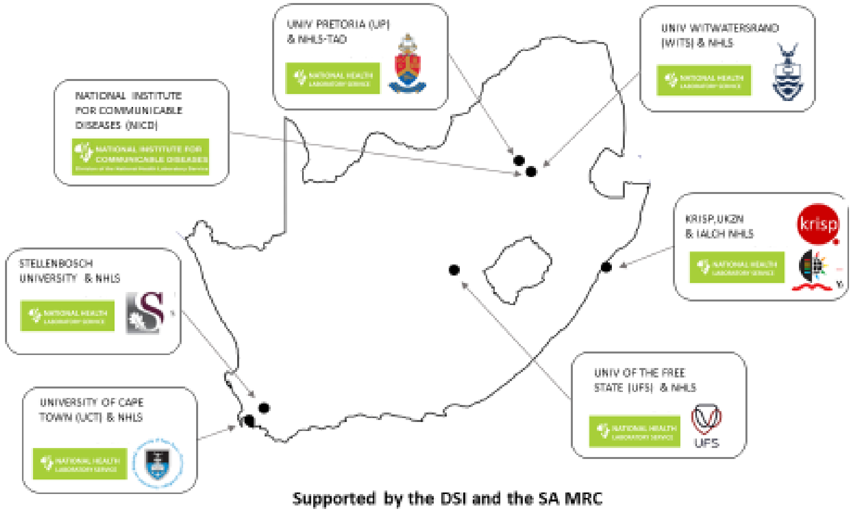
DSI
Glaudina Loots

SA MRC
Glenda Gray

Pathcare N1 City
Jean Maritz
Nadine Cronje
Petra Raimond
Kim Hoek

NGS-SA

Network for Genomic Surveillance in South Africa



Supported by the DSI and the SA MRC

UNIVERSITEIT VAN PRETORIA
UNIVERSITY OF PRETORIA
YUNIBESITHI YA PRETORIA

UNIVERSITEIT
iYUNIVESITHI
STELLENBOSCH
UNIVERSITY

science & innovation
Department:
Science and Innovation
REPUBLIC OF SOUTH AFRICA

UFS

saMRC
advancinglife

NATIONAL INSTITUTE FOR COMMUNICABLE DISEASES
Division of the National Health Laboratory Service

EDCTP

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

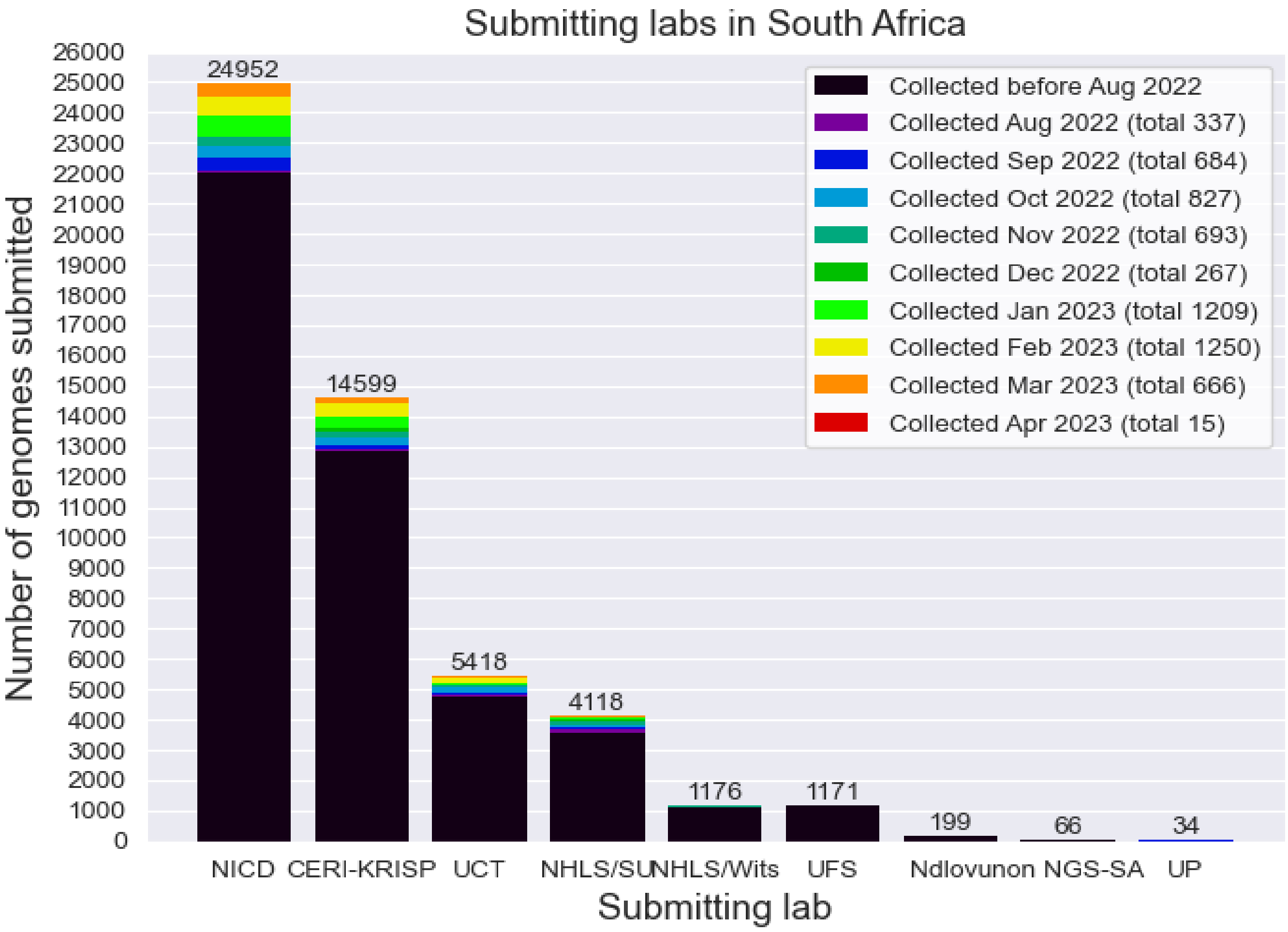
UNIVERSITY OF CAPE TOWN
IYUNIVESITHI YASEKAPA • UNIVERSITEIT VAN KAAPSTAD

WITS
UNIVERSITY

UNIVERSITY OF
KWAZULU-NATAL
INYUVESI
YAKWAZULU-NATALI

NATIONAL HEALTH
LABORATORY SERVICE

South African genomes submitted per submitting lab, 2020 - 2023 (N=51 733)



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.

NGS-SA
Network for Genomic Surveillance in South Africa

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.1 + S:F486P (similar Spike genetic profile as XBB.1.9.1)	05-01-2022	11-Jan-2023 XBB.1.5 Rapid Risk Assessment, 11 January 2023 XBB.1.5 Updated Rapid Risk Assessment, 25 January 2023 XBB.1.5 Updated Risk Assessment, 24 February 2023

https://www.who.int/en/activities/tracking-SARS-CoV-2-variants/ accessed 21 April 2023

Currently circulating variants under monitoring (VUMs)

Pango lineage# (+ mutation)	**Nextstrain clade**	**Spike genetic features**	**Earliest documented samples**	**Date of designation and risk assessments**
BA.2.75	22D	BA.2 + S:K147E, S:W152R, S:F157L, S:I210V, S:G257S, S:D339H, S:G446S, S:N460K, S:Q493R reversion	31-12-2021	06-Jul-2022
CH.1.1	22D	BA.2.75 + S:L452R, S:F486S	27-07-2022	08-Feb-2023
BQ.1	22E	BA.5 + S:R346T, S:K444T, S:N460K	07-02-2022	21-Sep-2022
XBB	22F	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	12-Oct-2022
XBB.1.16	23B	Recombinant of BA.2.10.1 and BA.2.75 sublineages, i.e. BJ1 and BM.1.1.1 XBB.1 + S:E180V, S:K478R and S:F486P	23-01-2023	22-03-2023
XBB.1.9.1	Not assigned	Recombinant of BA.2.10.1 and BA.2.75 sublineages, i.e. BJ1 and BM.1.1.1 XBB.1 + S:F486P (similar Spike genetic profile as XBB.1.5)	05-12-2022	30-03-2022
XBF	Not assigned	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	08-Feb-2023

https://www.who.int/en/activities/tracking-SARS-CoV-2-variants/ accessed 21 April 2023

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.)