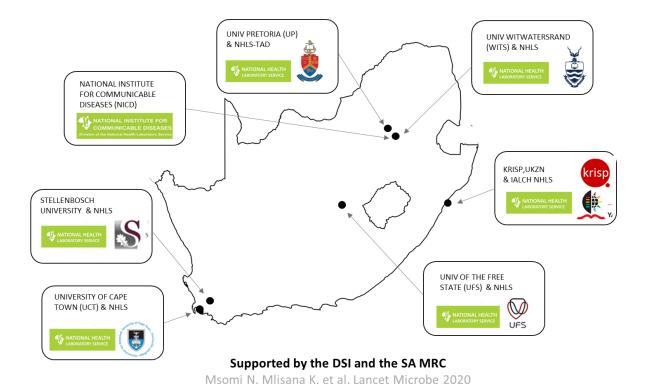


GS-SA Network for Genomic Surveillance in South Africa (NGS-SA)

SARS-CoV-2 Sequencing Update 24 March 2023

























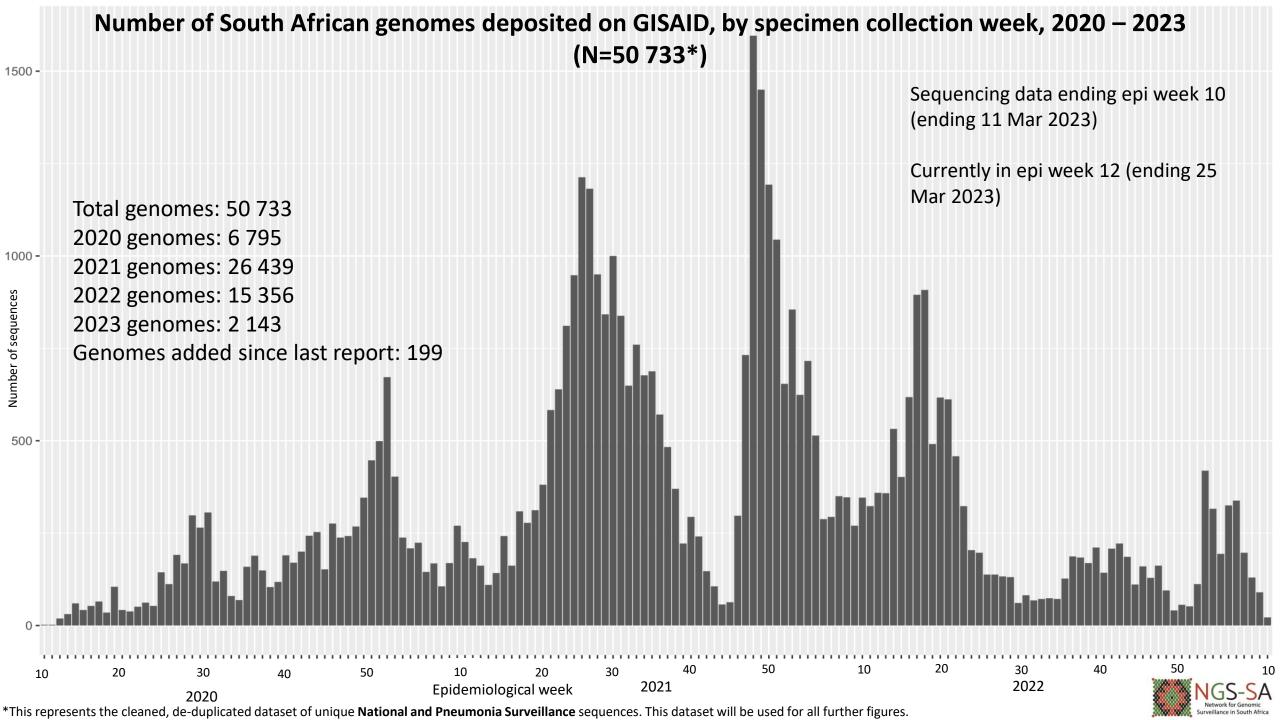
The genomic data presented here are based on South African SARS-CoV-2 sequence data downloaded from GISAID (www.gisaid.org) on 23 March 2023 at 07h15



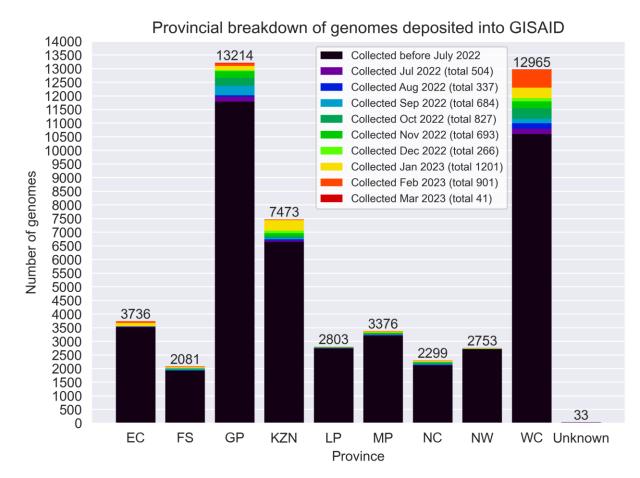
Data license: https://www.gisaid.org/registration/terms-of-use/

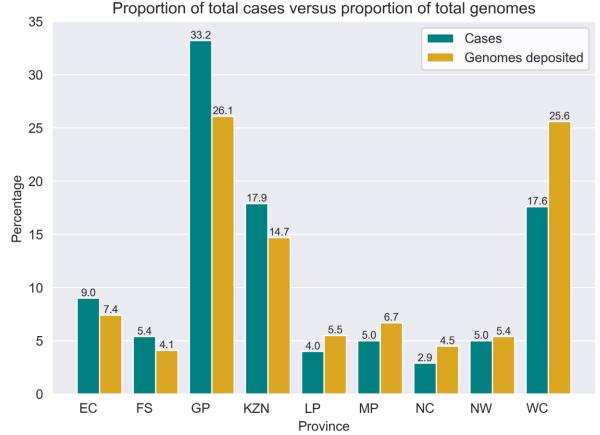
Elbe, S., and Buckland-Merrett, G. (2017) Data, disease and diplomacy: GISAID's innovative contribution to global health. Global Challenges, 1:33-46. DOI: 10.1002/gch2.1018 PMCID: 31565258

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



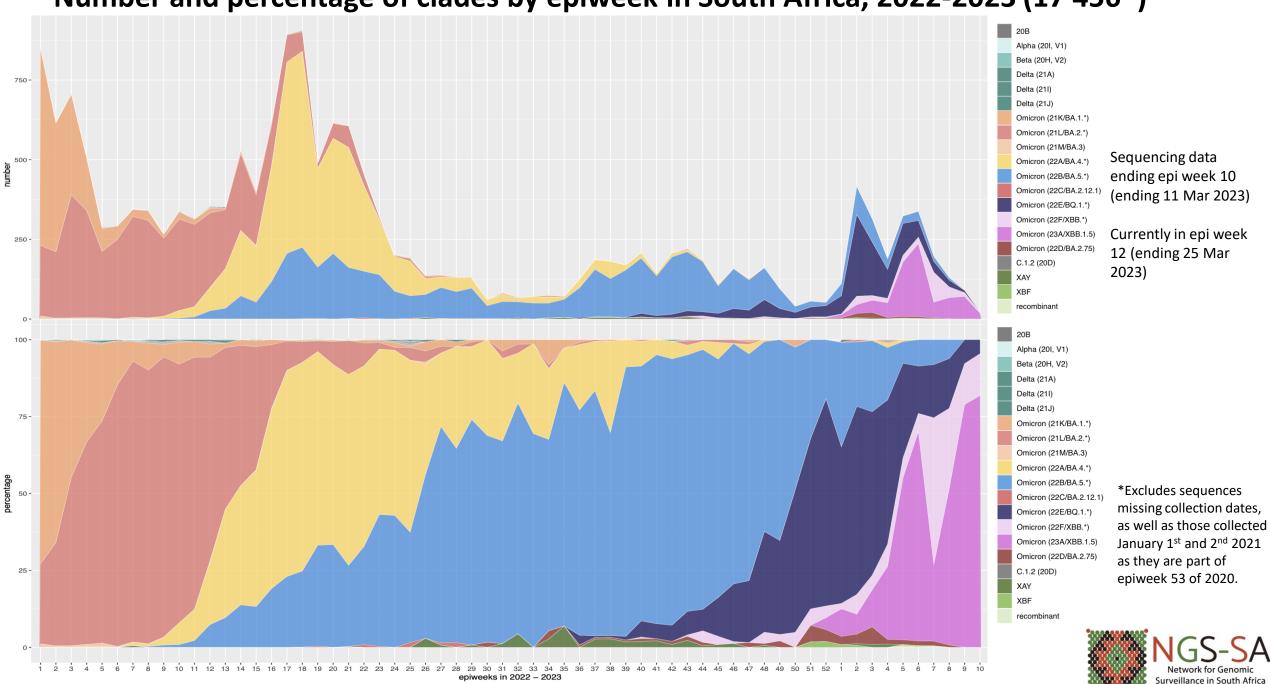
GISAID genomes vs total cases, 2020 - 2023 (N=50 733)





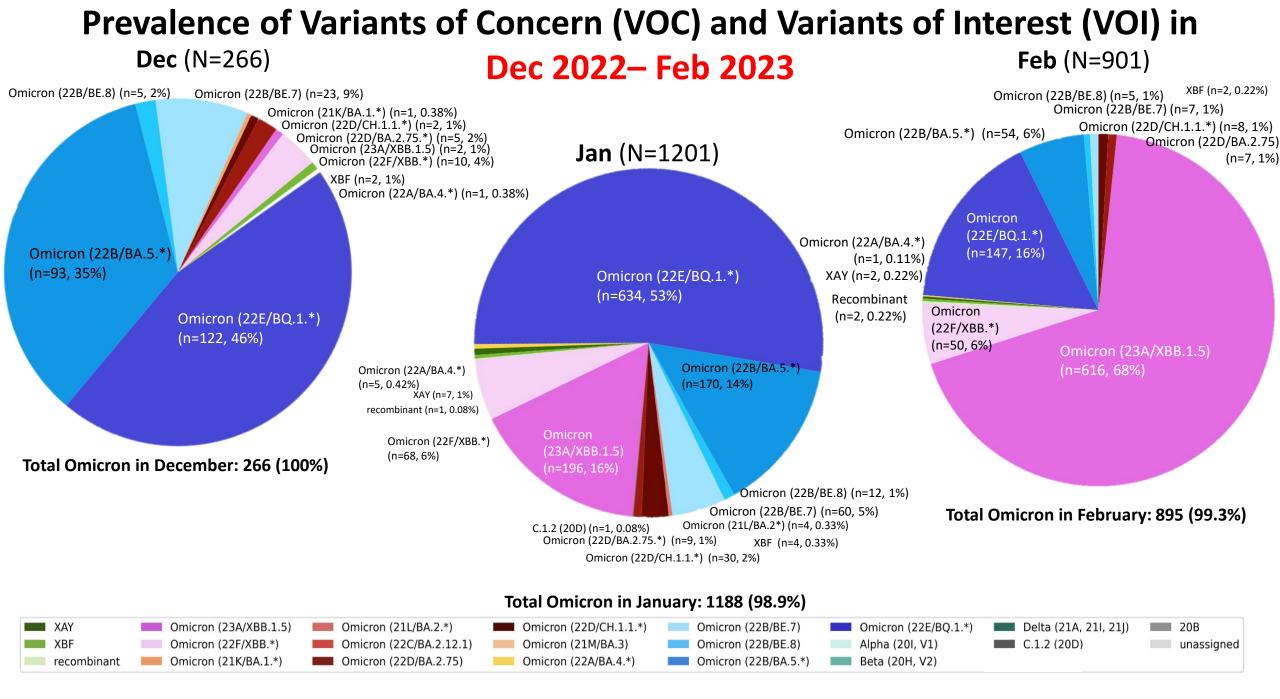


Number and percentage of clades by epiweek in South Africa, 2022-2023 (17 456*)

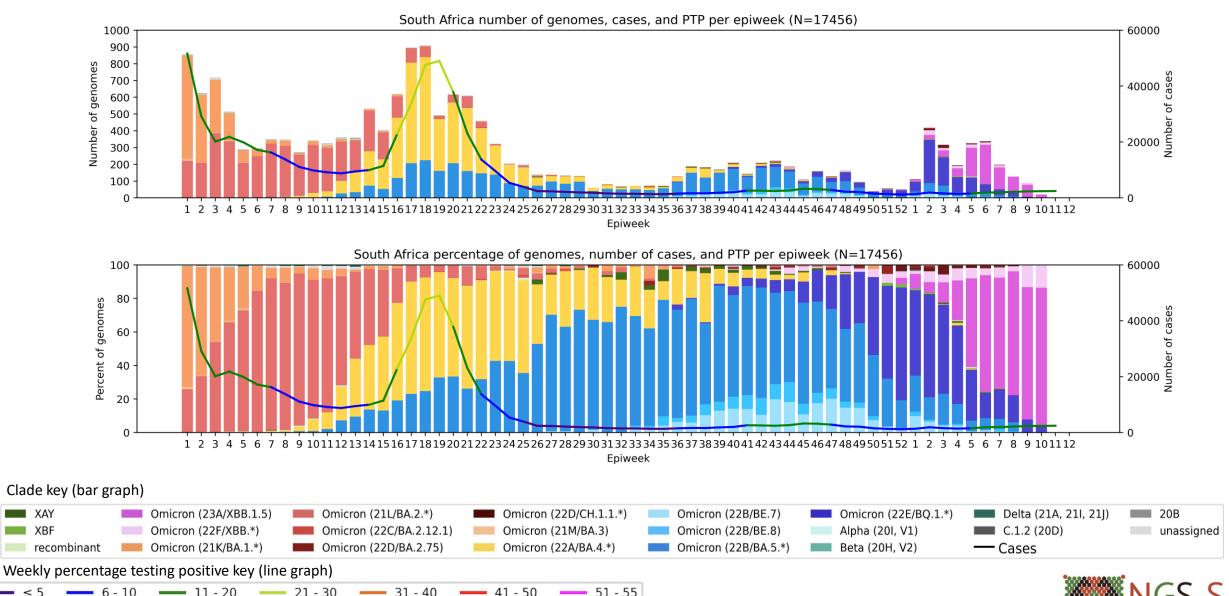


Detection Rates: Beta, Delta, C.1.2, recombinants, and Omicron Percentage 25 **Total sequences** 1515 2555 337 1201 Month Omicron Omicron Omicron Omicron Omicron (22C/BA.2.12.1) (23A/XBB.1.5) (21L/BA.2.*) (22A/BA.4)(22E/BQ.1.*) Omicron Omicron Omicron Omicron Delta (21K/BA.1.*) (21M/BA.3) (22B/BA.5)(22D/BA.2.75.*)

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



South Africa, 2022-2023, n = 17 456*



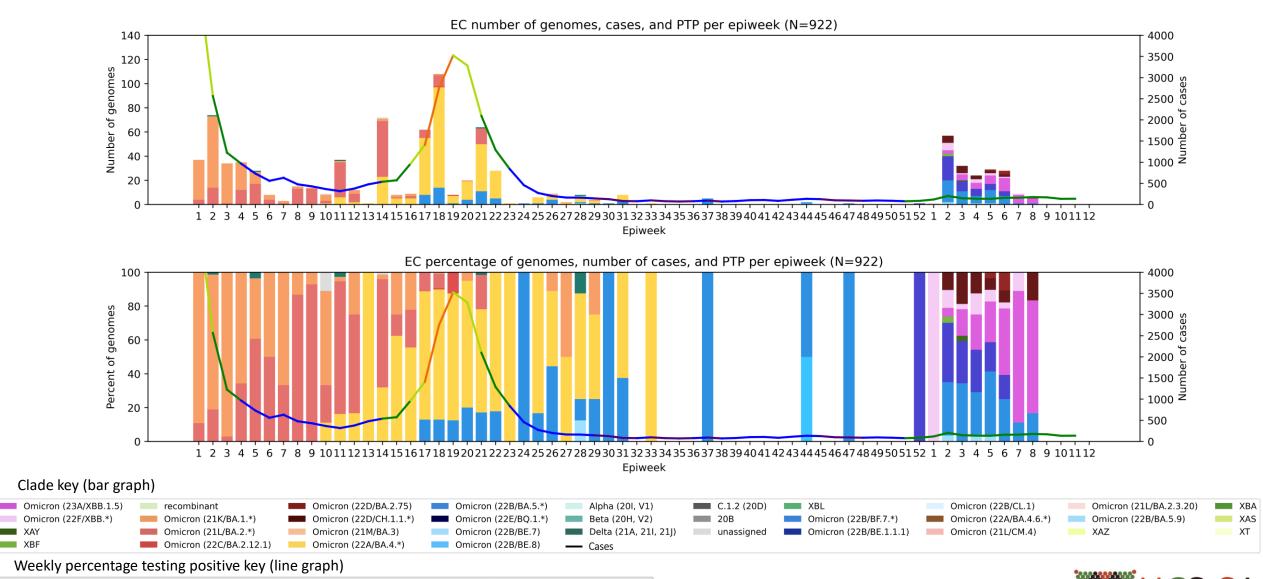
^{*}Excludes sequences missing collection dates. Lineages of particular interest (mainly WHO Omicron subvariants under monitoring) are separate from the main clade groupings.

^{*}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.



Eastern Cape Province, 2022-2023, n = 922

Genomes added since last report: 23*



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

41 - 50

- 51 - 55

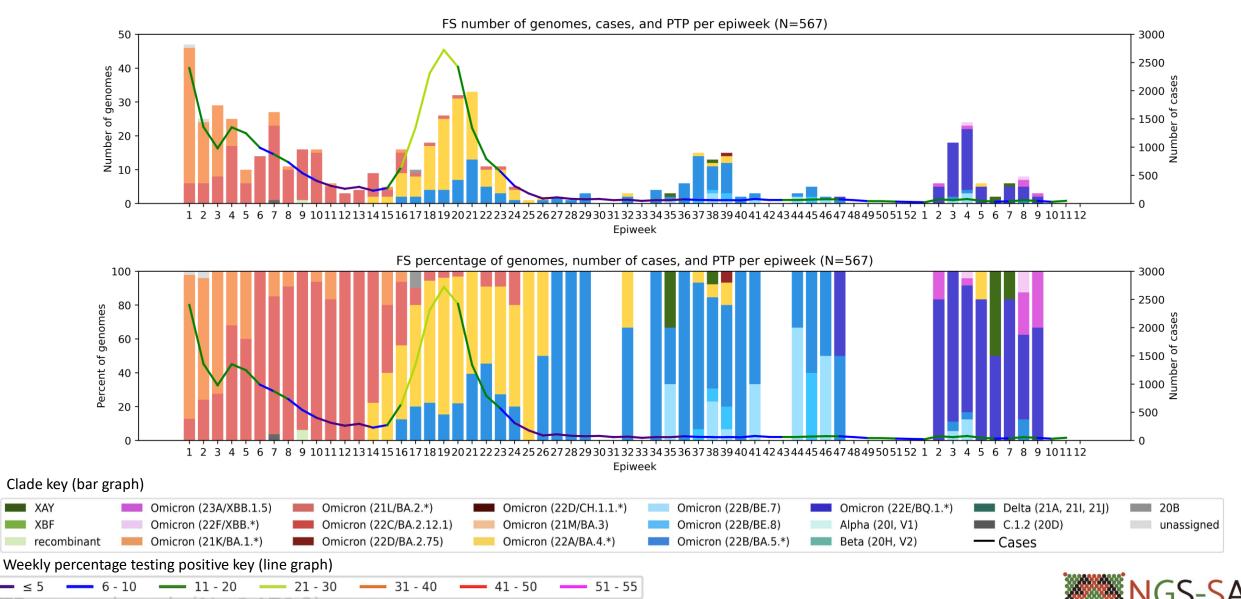
XAY

6 - 10

— 11 - 20

Free State Province, 2022-2023, n = 567

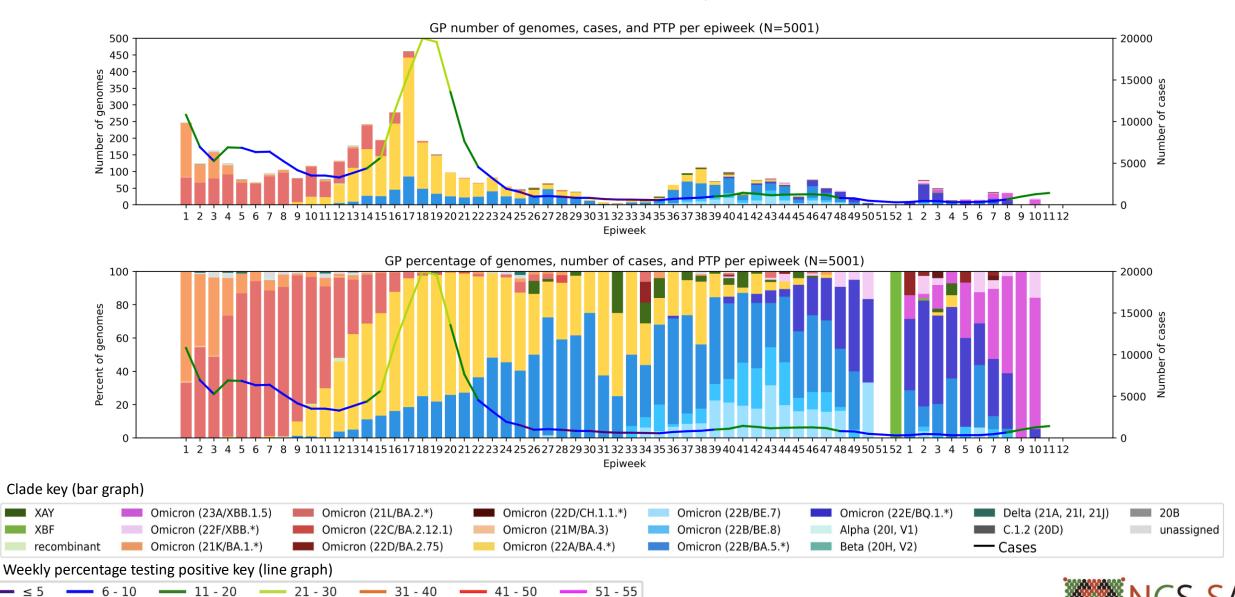
Genomes added since last report: 11*



^{*}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 = 5001

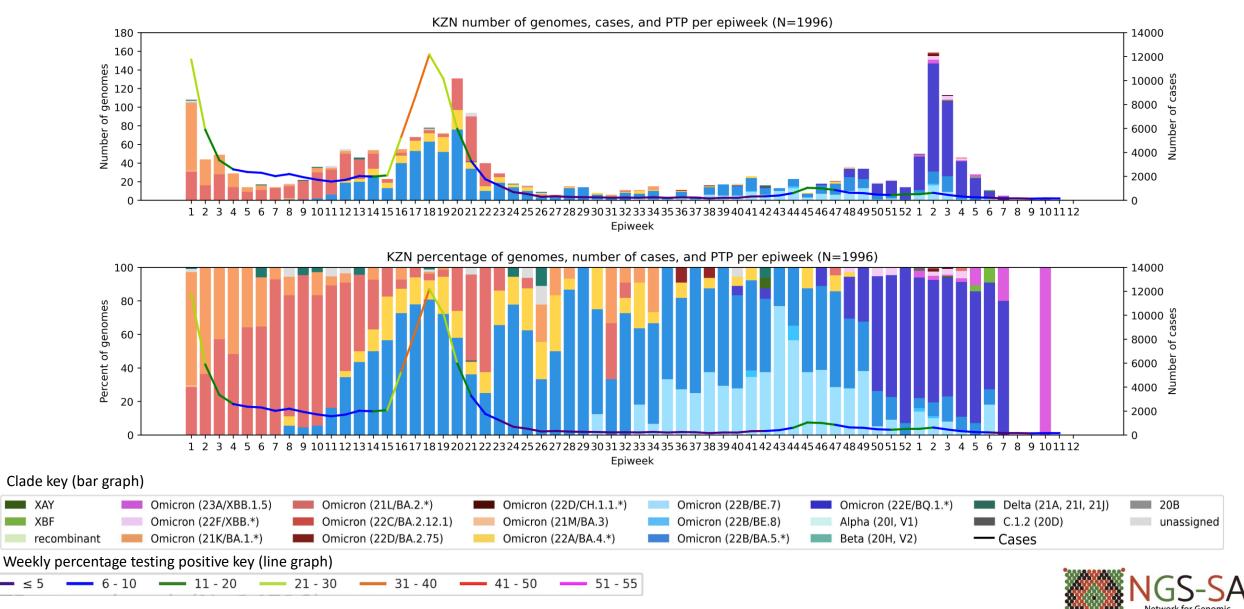
Genomes added since last report: 58*



^{*}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 = 1996

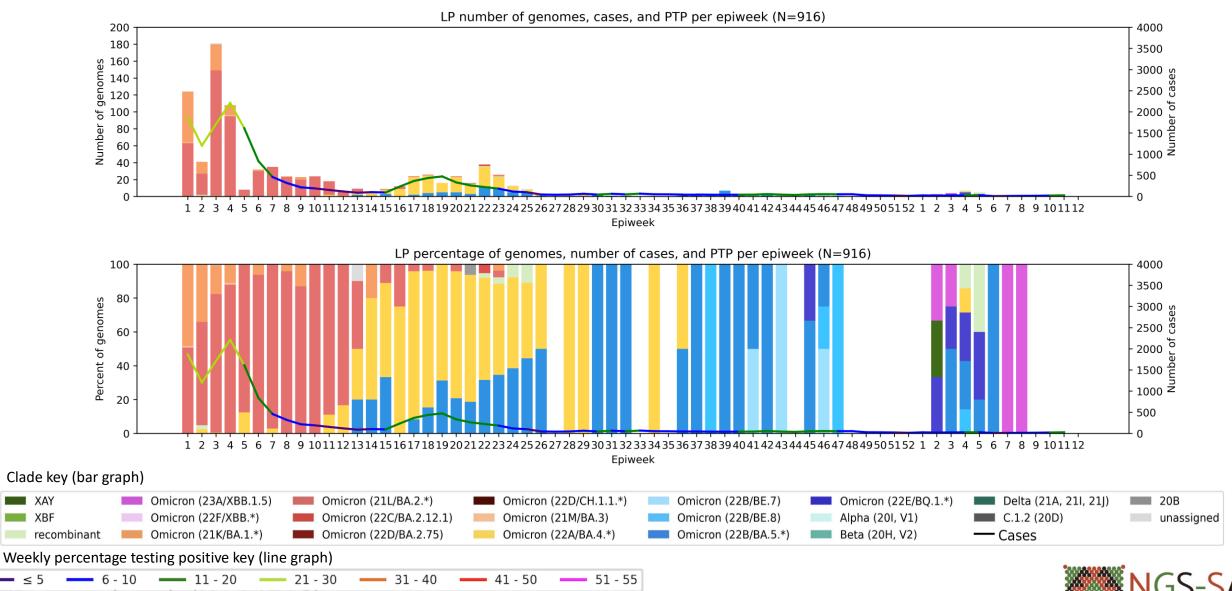
Genomes added since last report: 5*



^{*}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 = 916

Genomes added since last report: 1*

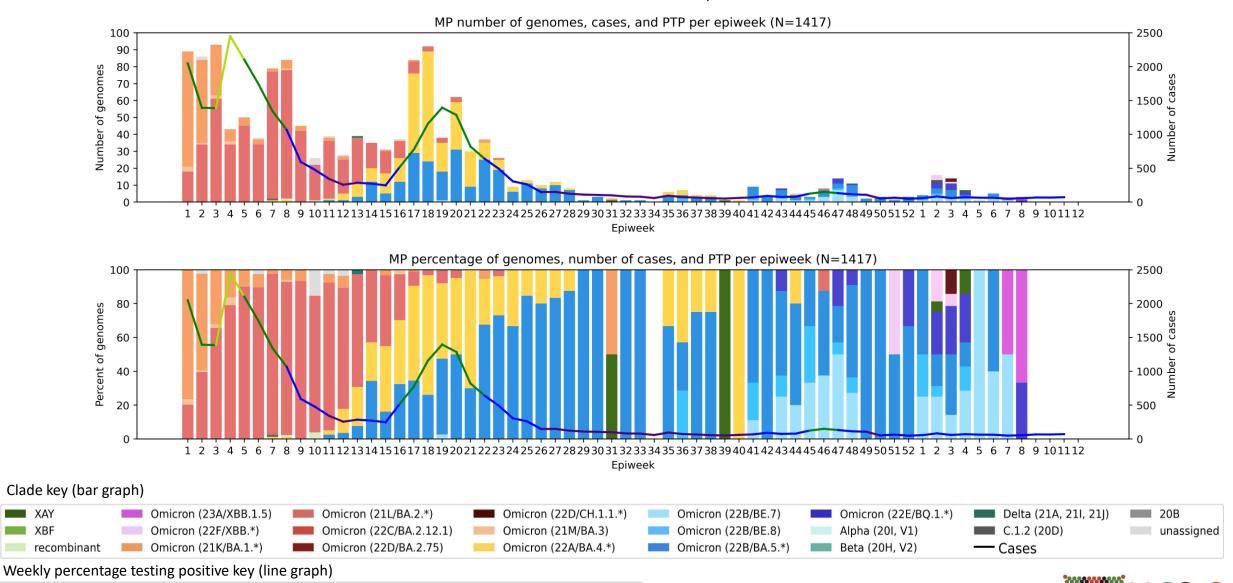


Network for Genomic Surveillance in South Africa

^{*}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 = 1417

Genomes added since last report: 3*



—— 31 - 40

41 - 50

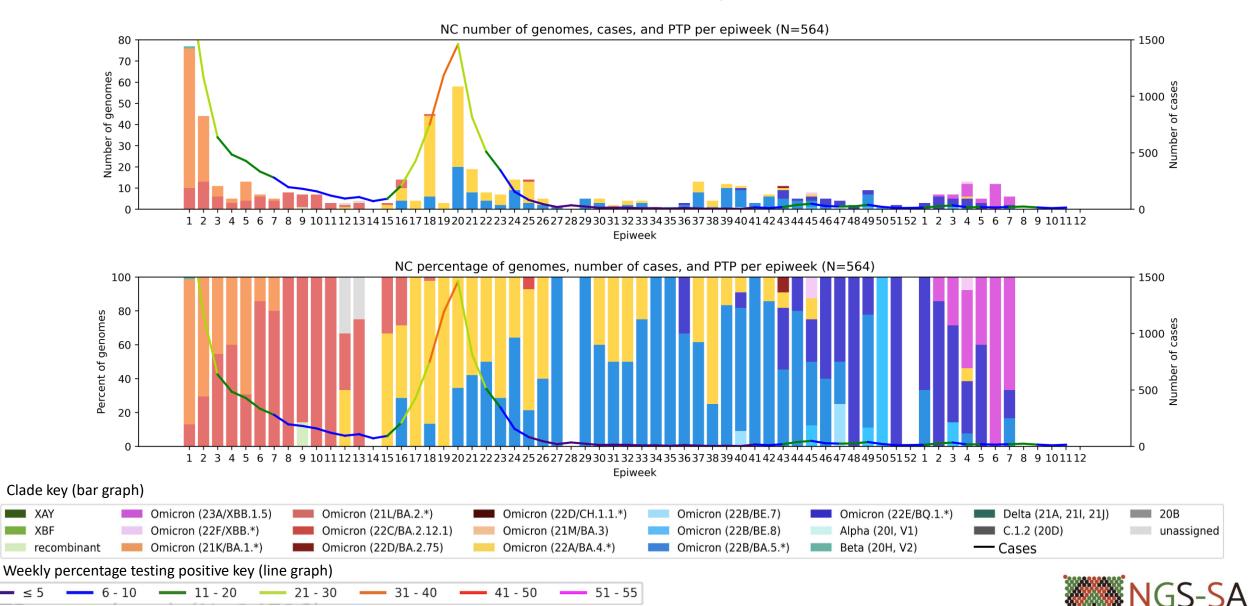
- 51 - 55

— 11 - 20

^{*}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 = 564

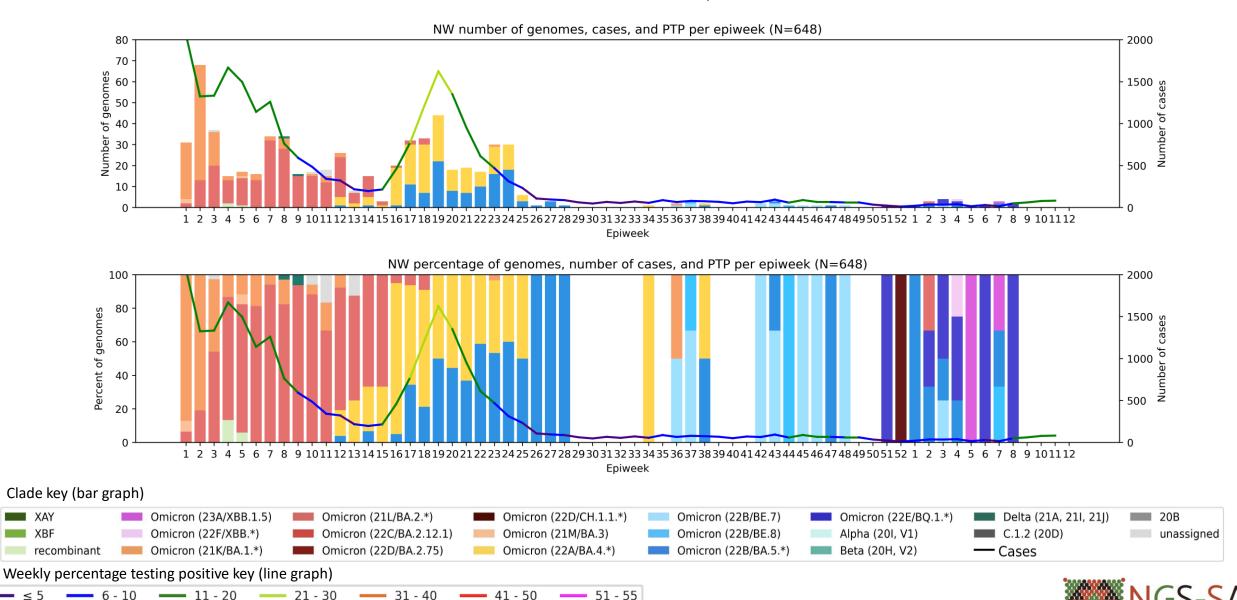
Genomes added since last report: 4*



^{*}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 = 648

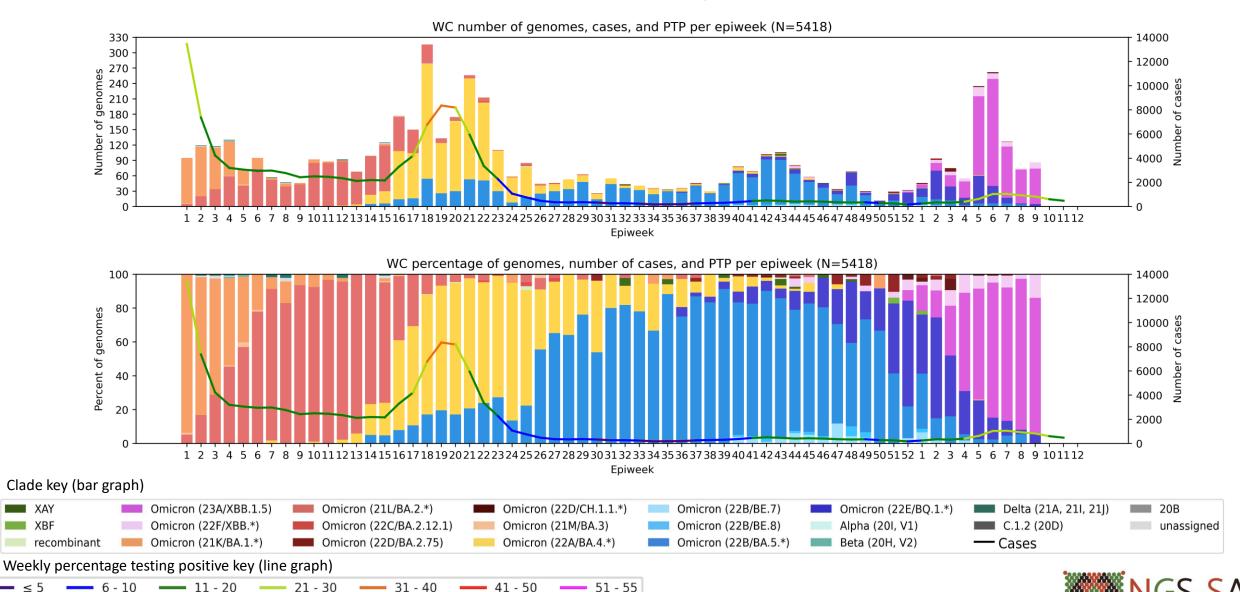
Genomes added since last report: 4*



^{*}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 = 5418

Genomes added since last report: 90*



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

Summary

Sequencing update

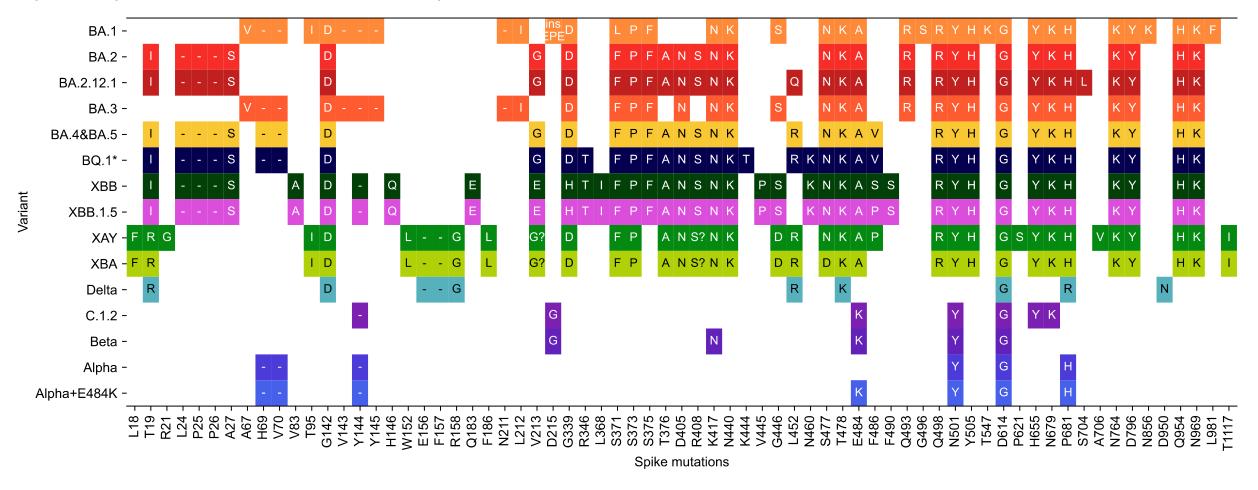
All provinces have sequences for January and February 2023. March sequences are from the Free State,
 Western Cape, KwaZulu-Natal and Gauteng

Variant of Concern Omicron in South Africa

- Omicron continued to dominate in December (100%), January (99%), February (99%) and makes up 100% of March sequences
- 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 is the dominant lineage in February (55%) and March (78%)
- BA.2.75.* continued to be detected at a low prevalence in December, January and February (≤2%)



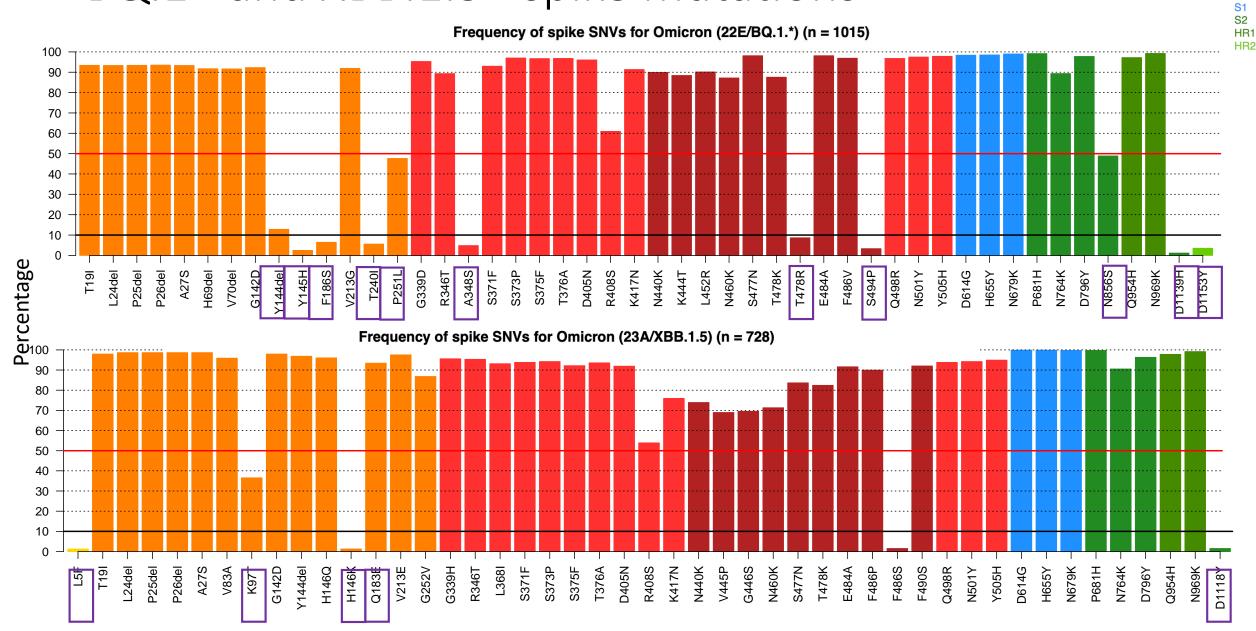
Spike protein mutation* profile of Variants of Interest and Concern



- 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



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



NTD RBD RBM

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 Kerri Francois

Dr Cherise Naicker

Dr Joedene Chetty



Dr Khanyi Msomi Dr Neli Ngcaba Dr Kerusha Govender Dr Tshepiso Mosito Dr Pravi Moodlev Mr Malcolm Ellapen Dr Aabida Khan Mr Kubendran Reddy Dr Lili Gounder The COVID-19 Bench team

University of KwaZulu-Natal & Africa Health Research Institute

Eduan Wilkinson

Sureshnee Pillav

Jennifer Giandhari

Emmanuel James San



KRISP at UKZN: Tulio de Oliveira Richard Lessels Houriivah Tegally

AHRT AFRICA
RESEARCH
RESEARCH
RESEARCH
RESEARCH

Alex Sigal Sandile Cele Willem Hanekom

University of Cape Town, NHLS & Western Cape Government



NHLS-UCT

(U. California)

Carolyn Williamson Nei-yuan Hsiao Diana Hardie Kruger Marais Stephen Korsman Zivaad Valley-Omar

health

WCG-UCT

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

NHLS Greenpoint Annabel Enoch

eceived funding from the European Union's Horizon Europe Research and Innovation Actions under grant No



 $\Lambda \Lambda$

EDCTP

Samrce



ZARV research program/UP

Zoonotic arbo and respiratory virus

program

Centre for Viral Zoonoses Department Medical Virology/ NHLS

Tshwane Academic division

University of Pretoria

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)

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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

health

Cheryl Cohen

Centre for HIV and STIs

Brent Oosthuysen Penny Moore Lynn Morris

NICD Groups

AFRICA CDC

Sequencing Core Facility

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







Thokozani Mkhize Diagnostic laboratory staff

UFS



University of the

Free State

Dominique Goedhals

Emmanuel Ogunbavo

Makgotso Maotoana

NHLS Division of Virology

Lutfiyya Mohamed

Sabeehah Vawda

Felicity Burt

Armand Bester

Martin Myaga

Peter Mwangi

Milton Mogotsi

UCT, IDM and CIDRI-Africa

Deelan Doolabh Arash Iranzadeh Lynn Tyers Innocent Mudau Nokuzola Mbhele Fezokuhle Khumalo Thabang Serakge Bruna Galvão Linda Boloko Arghavan Alisoltani

Robert Wilkinson Darren Martin Nicola Mulder Wendy Burgers Ntobeko Ntusi Rageema Joseph

CAPE TOWN HYTN Sean Wasserman



Jinal Bhiman Cathrine Scheepers Constantinos Kurt Wibmer Thandeka Movo **Tandile Hermanus** Frances Ayres Zanele Molaudzi **Bronwen Lambson Tandile Hermanus** Mashudu Madzivhandila Prudence Kgagudi

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











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







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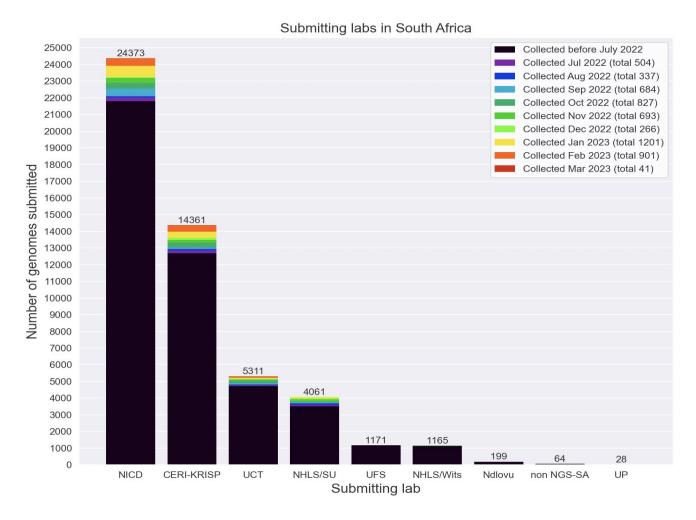








South African genomes submitted per submitting lab, 2020 - 2022 (N=50 733)



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



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.	05-01-2022	11-Jan-2023
		XBB + S:F486P		

Omicron subvariants under monitoring

Pango lineage [#] (+ mutation)	GISAID clade	Nextstrain clade	Relationship to circulating VOC lineages	Spike genetic features	Earliest documented samples
BQ.1 ^{\$}	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 [§]	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§	GRA	22D	BA.2 sublineage	BA.2.75 + S:L452R, S:F486S	27-07-2022
XBB ^μ	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

[#] includes descendent lineages

^{\$} additional mutation outside the spike protein: ORF1a: Q556K, L3829F, ORF1b: Y264H, M1156I, N1191S, N: E136D, ORF9b: P10F

[§] additional mutations outside of the spike protein: ORF1a: S1221L, P1640S, N4060S, ORF1b: G662S, E: T11A

μ 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 (≥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.)