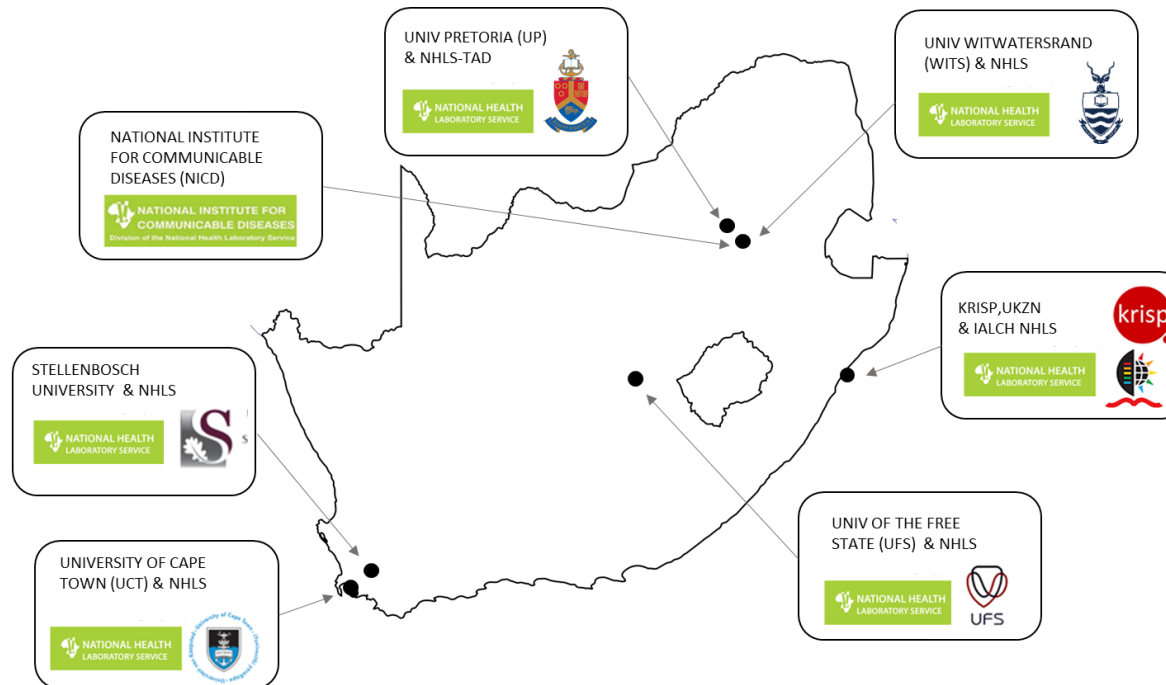


SARS-CoV-2 Sequencing Update 9 September 2021



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 9 September at 08h00



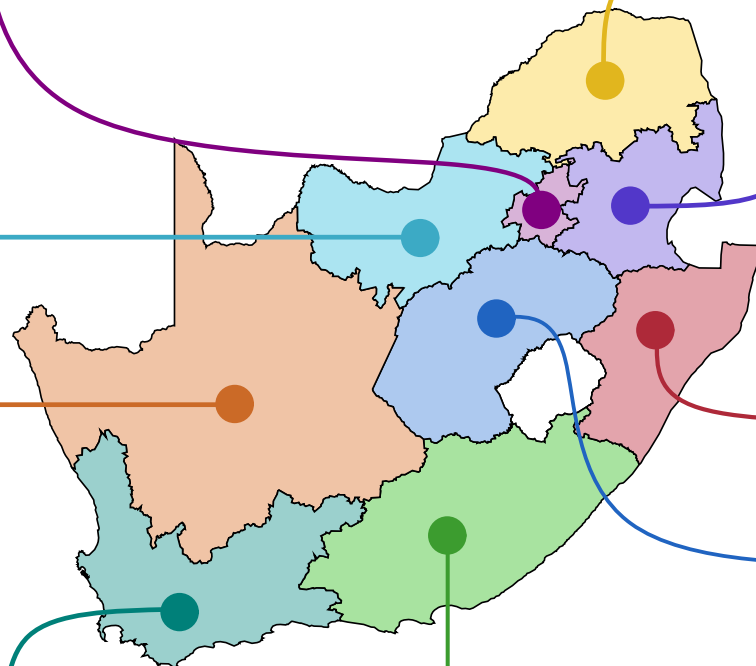
Data license: <https://www.gisaid.org/registration/terms-of-use/>

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

GENOMIC SURVEILLANCE IN THE THIRD WAVE

epiweeks 18 - 36



Gauteng

Genomes Cases
2 564 (30.8%) 483 884 (39.2%)

Genomes deposited in the last week
30 3 1

North West

Genomes Cases
608 (7.3%) 77 234 (6.3%)

Genomes deposited in the last week
33 3 3

Northern Cape

Genomes Cases
227 (2.7%) 43 621 (3.5%)

Genomes deposited in the last week
3 25 7 1

Western Cape

Genomes Cases
1 758 (21.1%) 206 921 (16.8%)

Genomes deposited in the last week
17 99 1 2 7 3

Eastern Cape

Genomes Cases
828 (10.0%) 79 8181 (6.5%)

Genomes deposited in the last week
22 3

Limpopo

Genomes Cases
750 (9.0%) 56 622 (4.6%)

Genomes deposited in the last week
1

Mpumalanga

Genomes Cases
278 (3.3%) 68 346 (5.5%)

Genomes deposited in the last week
23 1 1

KwaZulu-Natal

Genomes Cases
904 (10.9%) 155 108 (12.6%)

Genomes deposited in the last week
82 1

Free State

Genomes Cases
397 (4.8%) 62 630 (5.1%)

Genomes deposited in the last week
1 8 146 2 2 2 2

Alpha (20I, V1) Eta (21D) 20C
Beta (20H, V2) C.1.2 (20D) 20D
Delta (21A) 20A Unassigned
Kappa (21B) 20B Cases

Bar graphs represent genomes sequenced per epiweek, line represents cases by collection date (weeks 18 – 36)

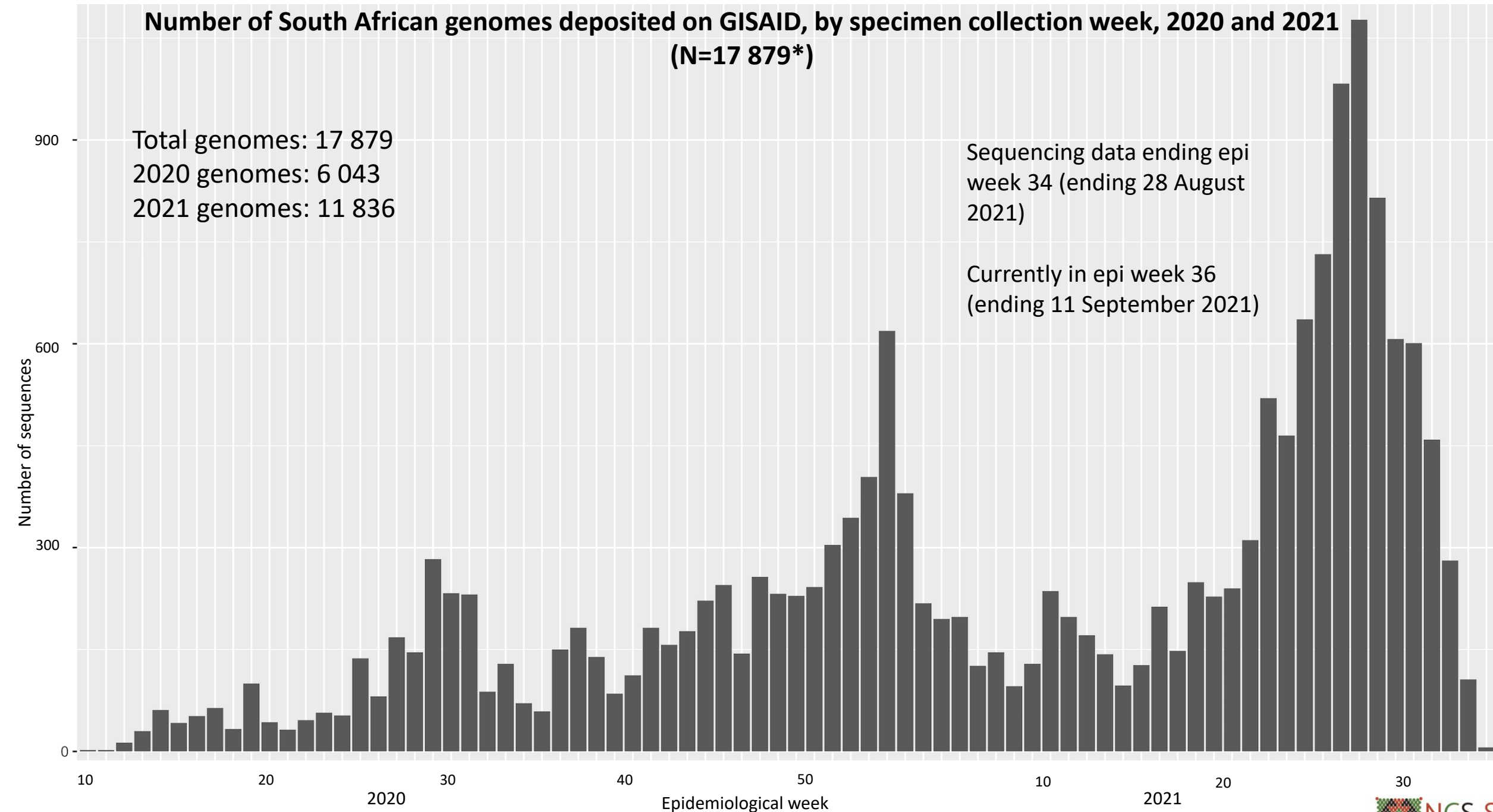
Genomes and cases presented as provincial total (percentage of national total) for epiweeks 18 - 36

Number of South African genomes deposited on GISAID, by specimen collection week, 2020 and 2021 (N=17 879*)

Total genomes: 17 879
2020 genomes: 6 043
2021 genomes: 11 836

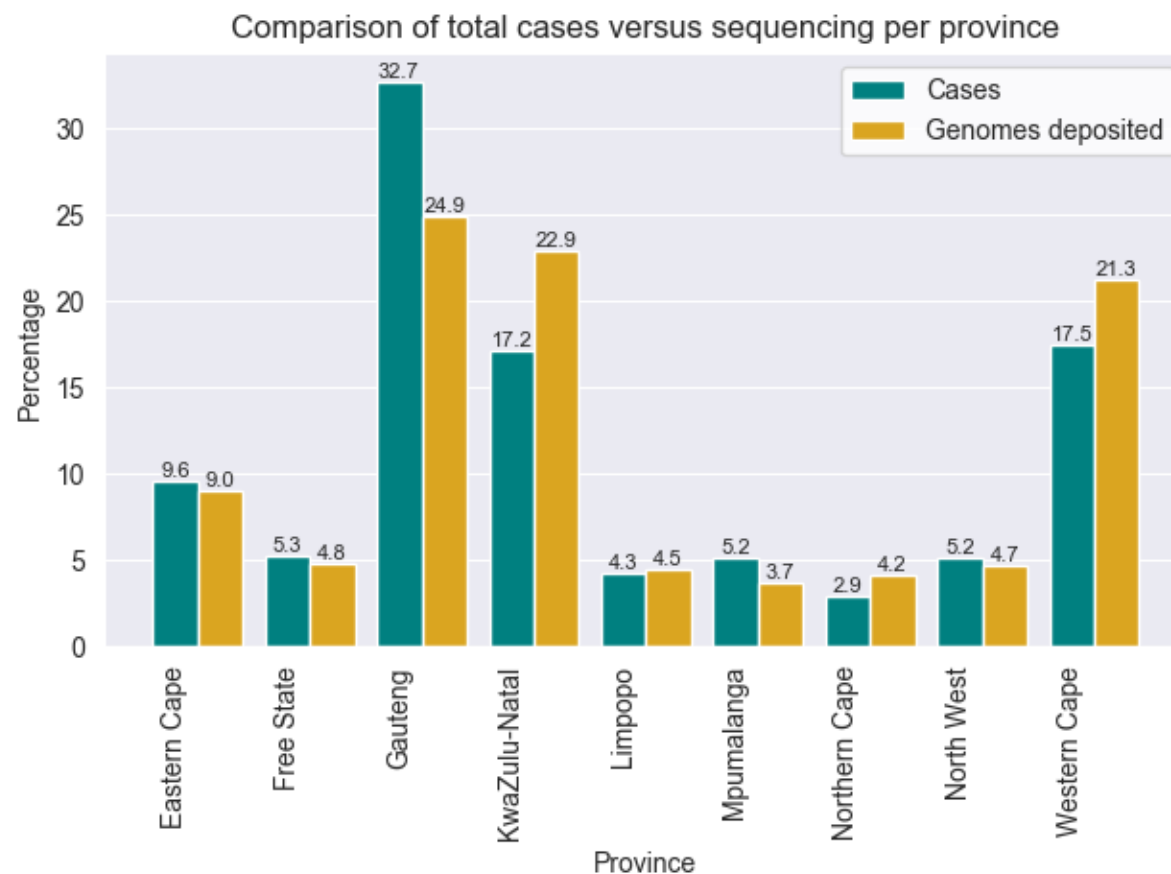
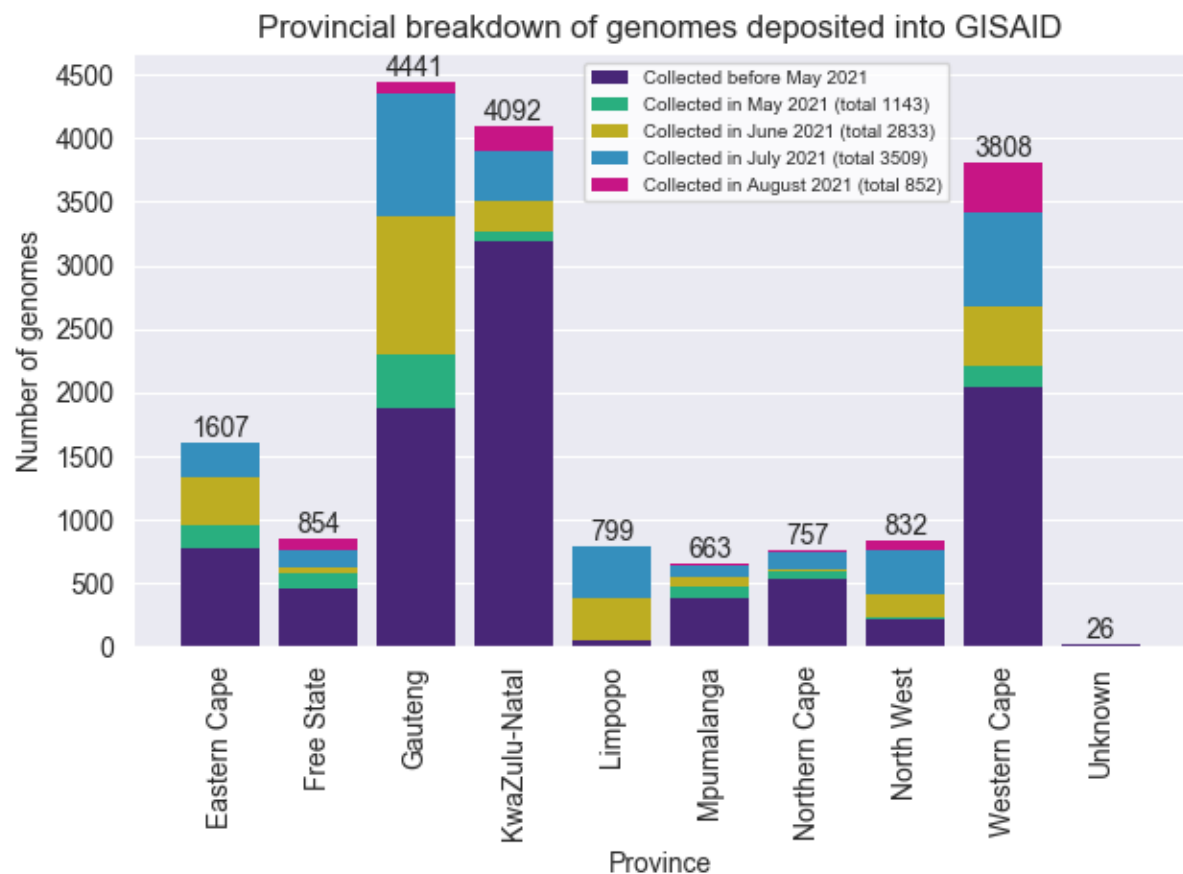
Sequencing data ending epi
week 34 (ending 28 August
2021)

Currently in epi week 36
(ending 11 September 2021)



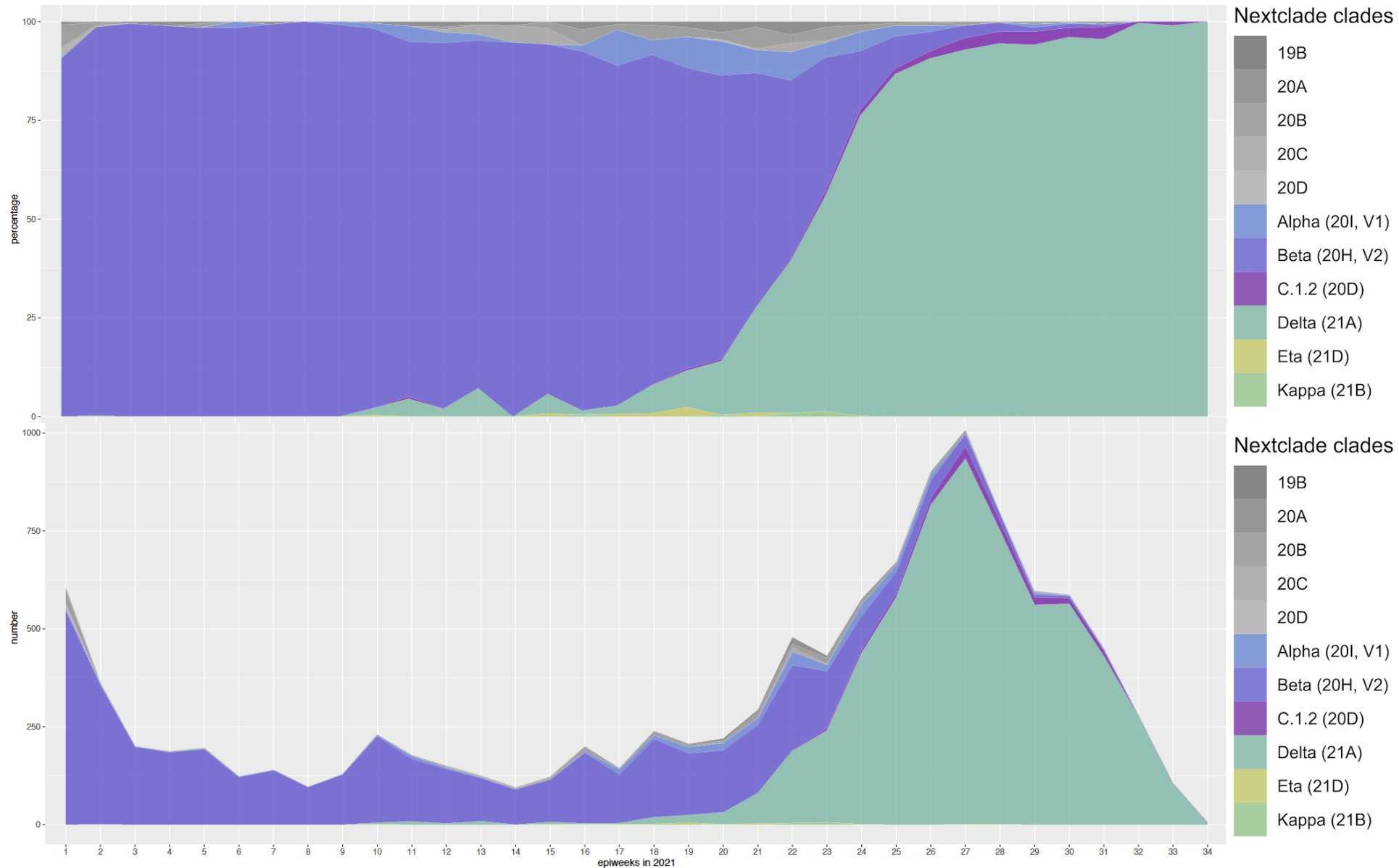
*This represents the cleaned, de-duplicated dataset of unique sequences. This dataset will be used for all further figures.

GISAID genomes vs total cases, 2020 and 2021 (N=17 879)



All provinces, apart from GP, KZN, and WC, have comparable percentage of overall cases and overall sequenced genomes

Distribution and number of clades in South Africa, 2021 (N= 11 836)



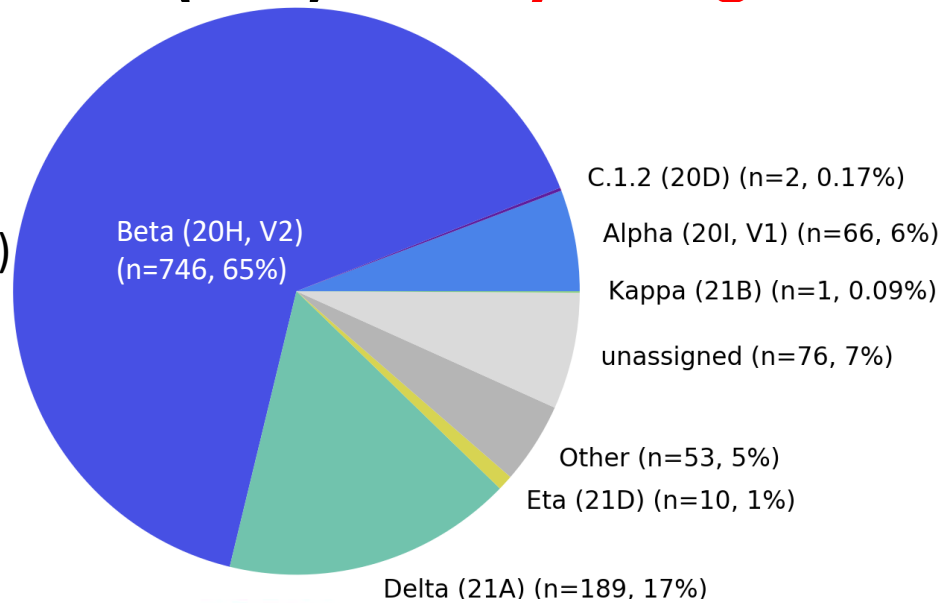
Sequencing data
ending epi week 34
(ending 28 August
2021)

Currently in epi
week 36 (ending 11
September 2021)

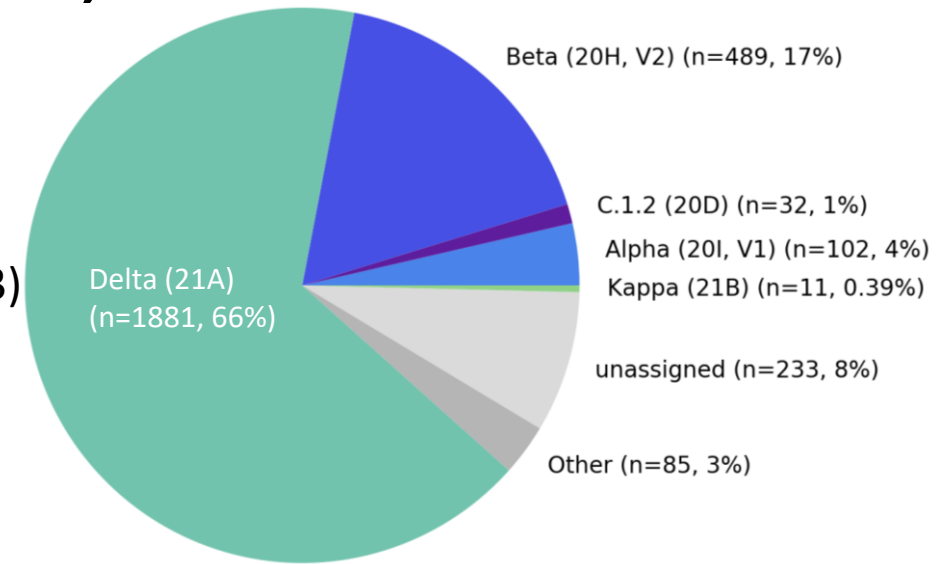
Delta came to dominate by end June at >65% , in July at >85% and in August at >90%
C.1.2 present at <3% frequency since March

Prevalence of Variants of Concern (VOC) and Variants of Interest (VOI) in **May – August 2021** sequences, South Africa

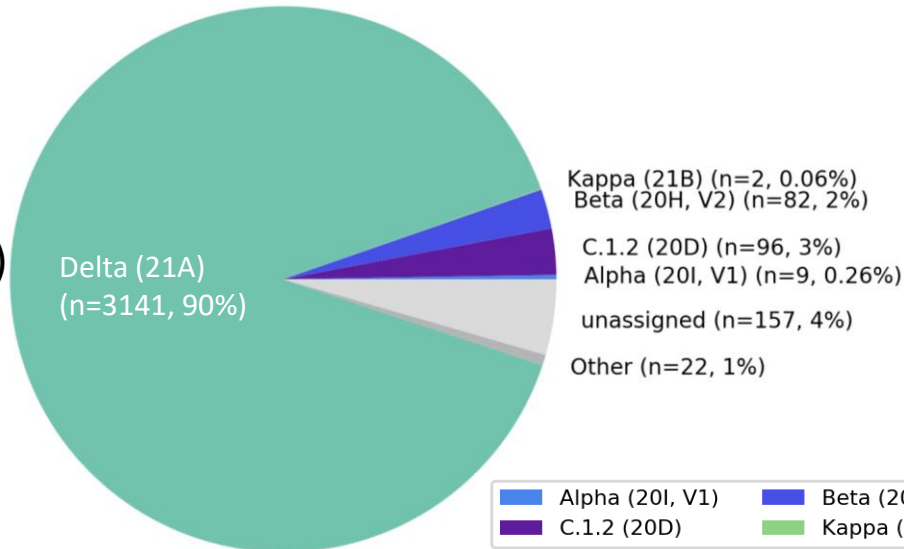
May (N = 1143)



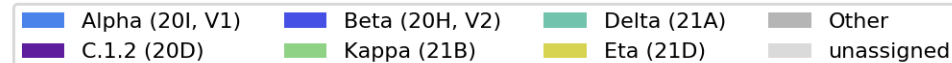
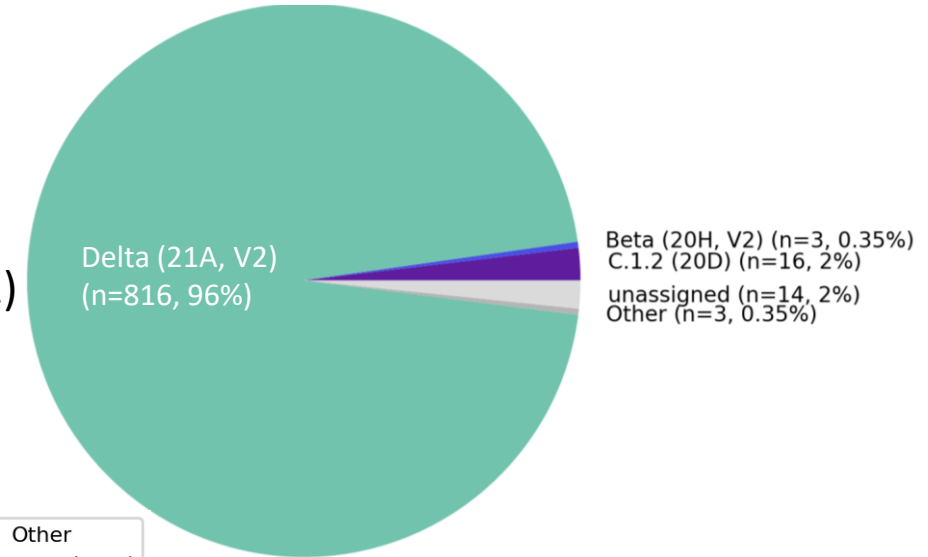
June (N = 2833)



July (N = 3509)

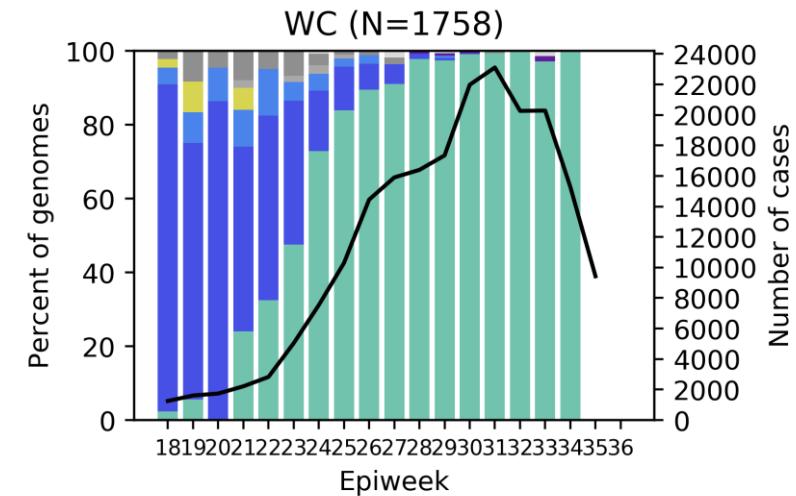
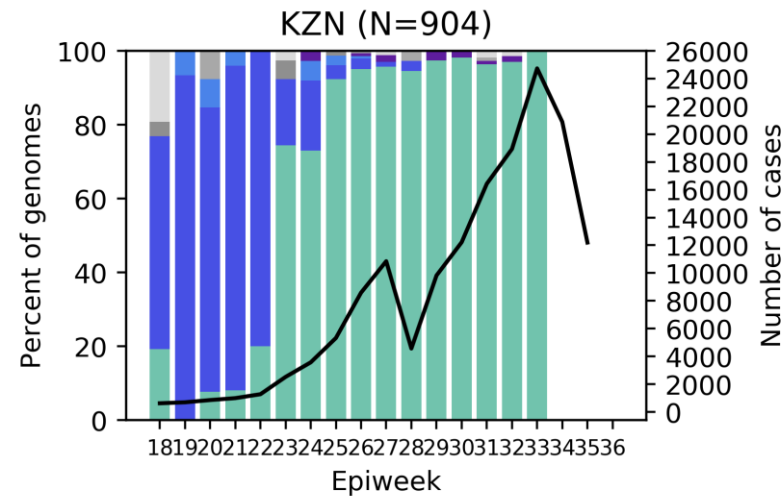
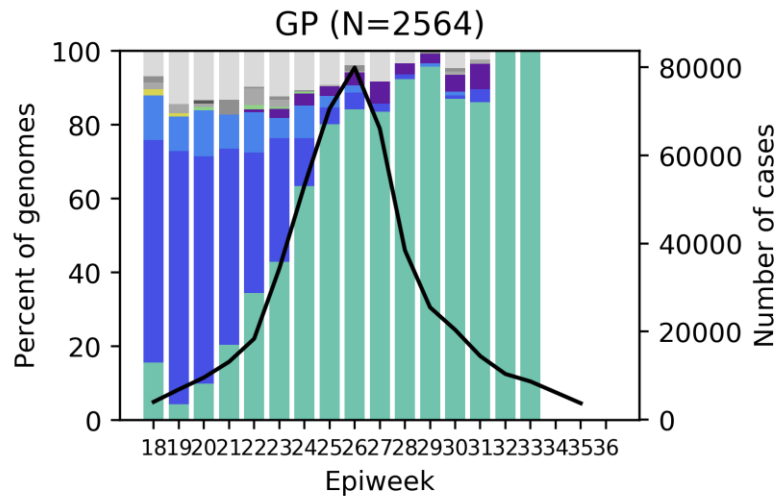
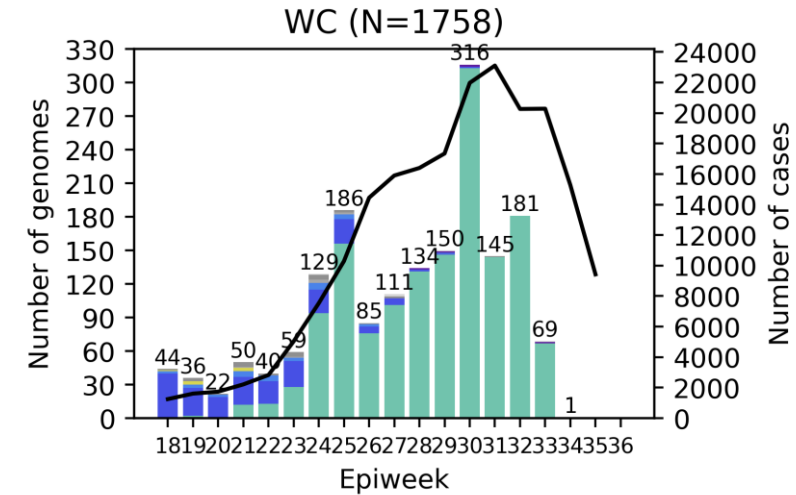
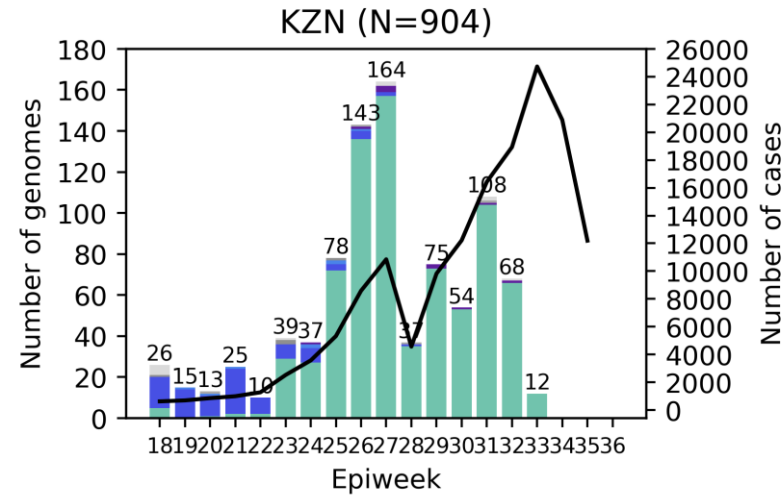
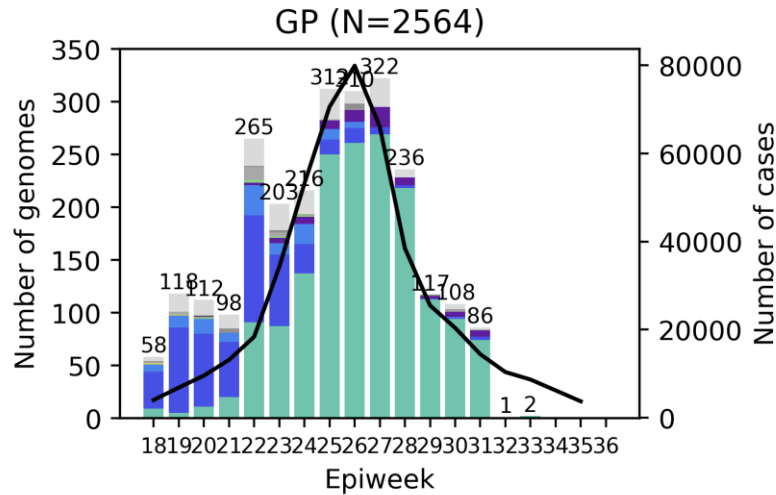


August (N = 852)



Beta variant dominated in May, while the Delta variant dominated in June-August in South Africa

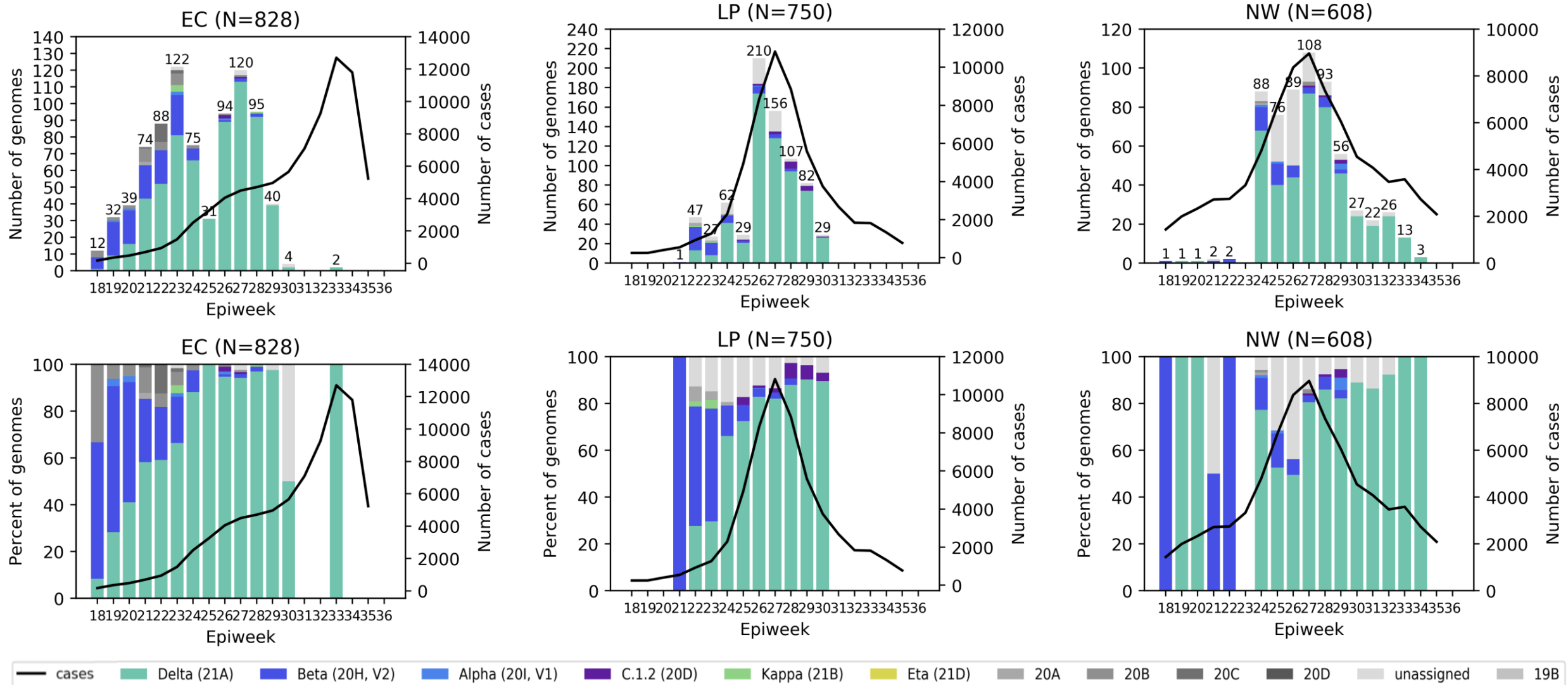
Genomes sequenced from specimens collected in May to mid-August 2021 (epiweeks 18 – 36) from KwaZulu-Natal, Gauteng, Western Cape Provinces



— cases Delta (21A) Beta (20H, V2) Alpha (20I, V1) C.1.2 (20D) Kappa (21B) Eta (21D) 20A 20B 20C 20D unassigned 19B

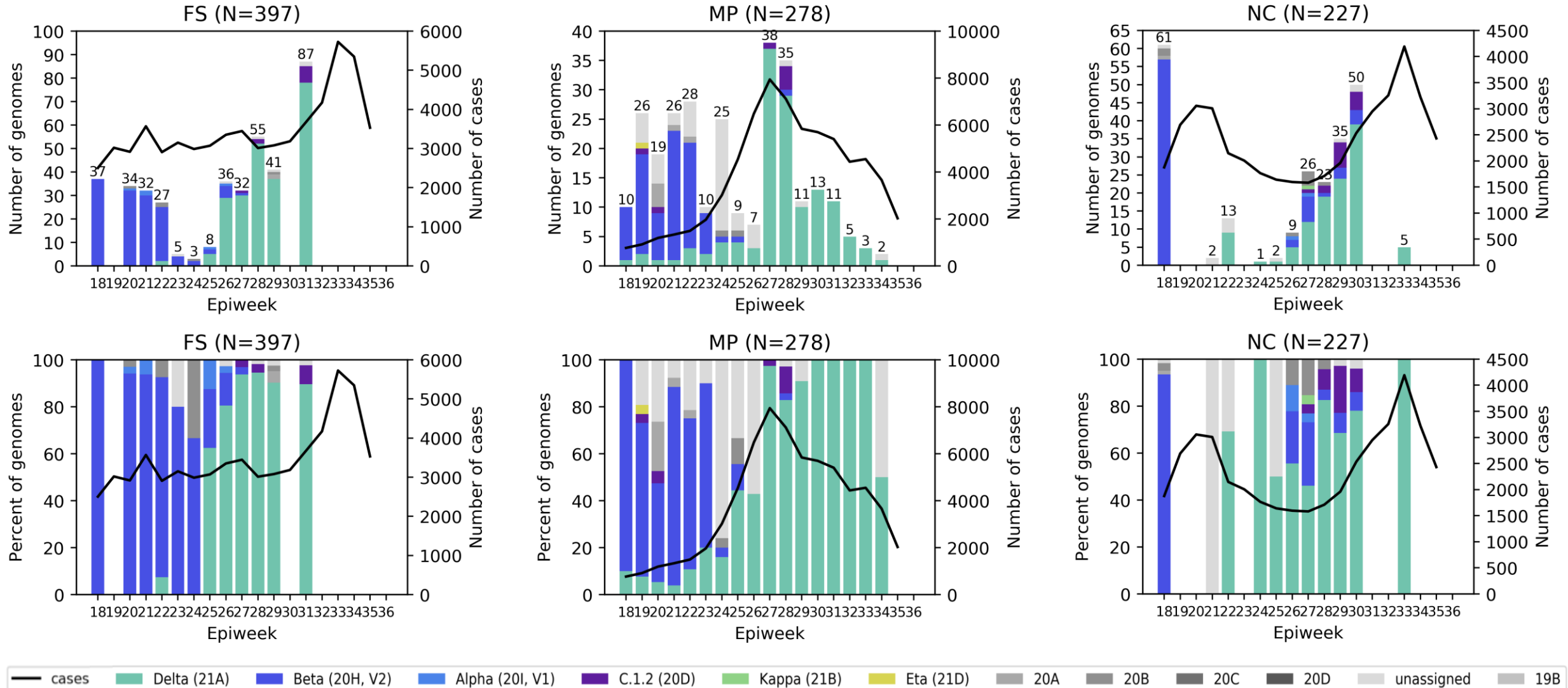
Beta dominates in all provinces in May, however following its detection in week 18, Delta rapidly begins to dominate in all three provinces by mid-June and continues to dominate in July and August

Genomes sequenced from specimens collected in May to mid-August 2021 (epiweeks 18 – 36) from Eastern Cape, Limpopo and North-West Provinces



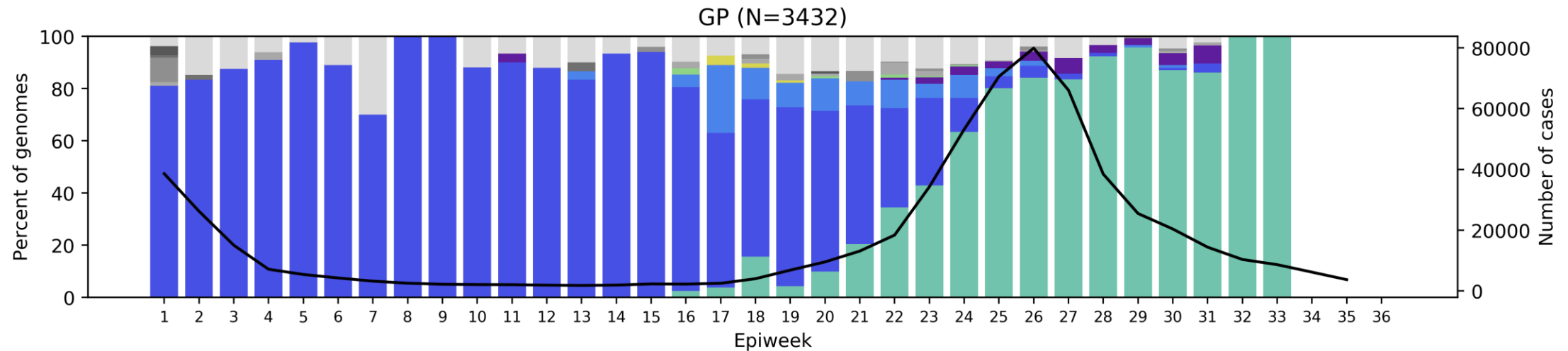
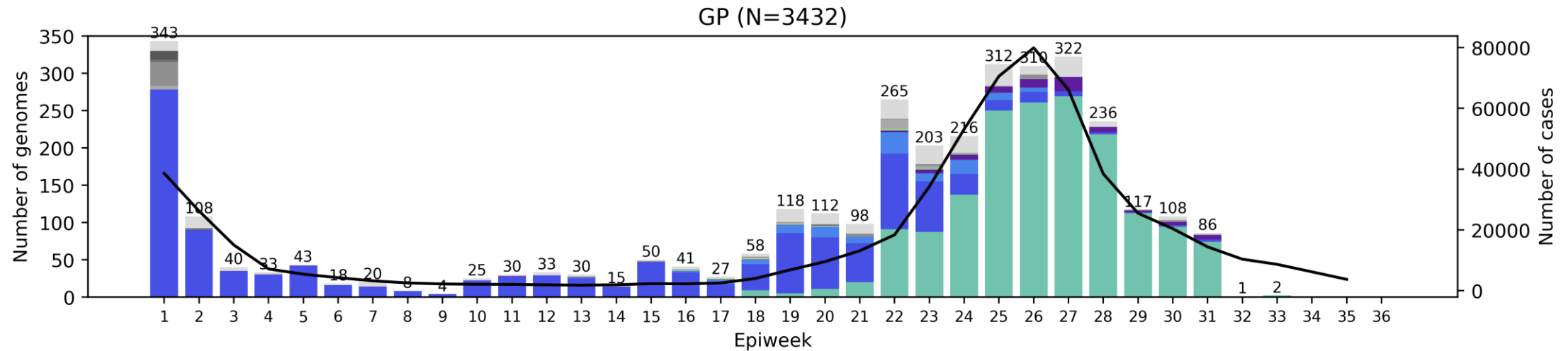
Delta variant dominating in Eastern Cape, Limpopo and North-West Provinces

Genomes sequenced from specimens collected in May to mid-August 2021 (epiweeks 18 – 36) from Free State, Mpumalanga and Northern Cape Provinces



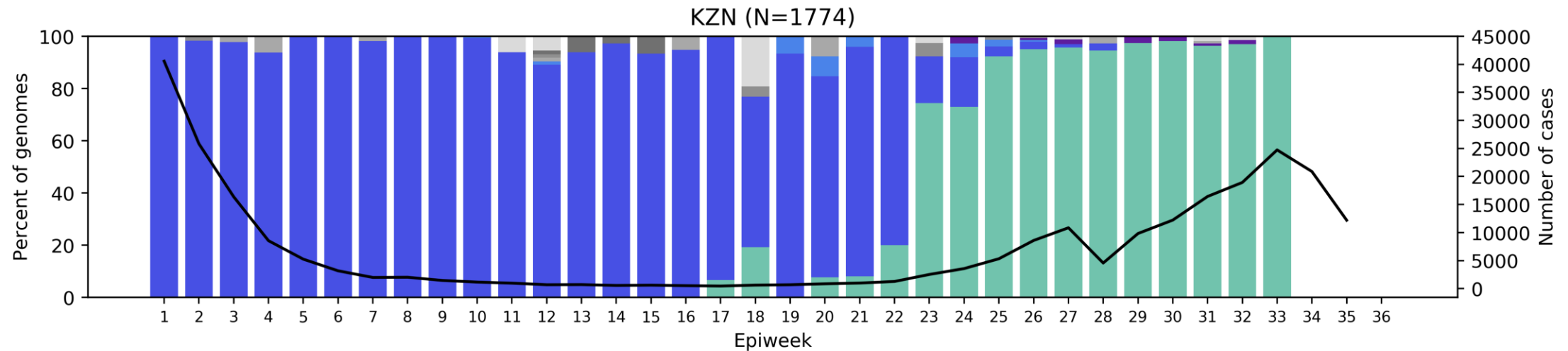
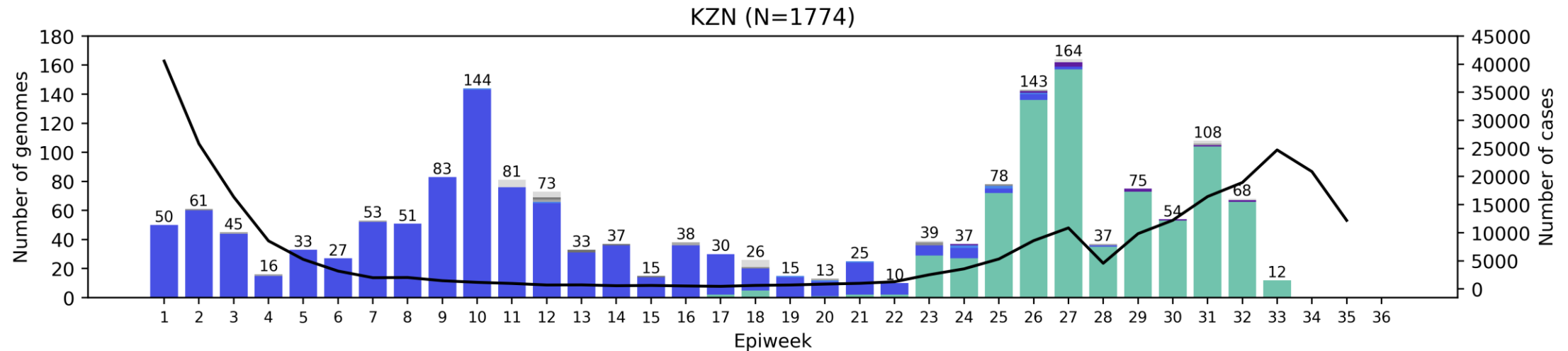
Despite small number of genomes, Delta dominating in FS, MP and NC provinces

Gauteng Province, 2021, n = 3432



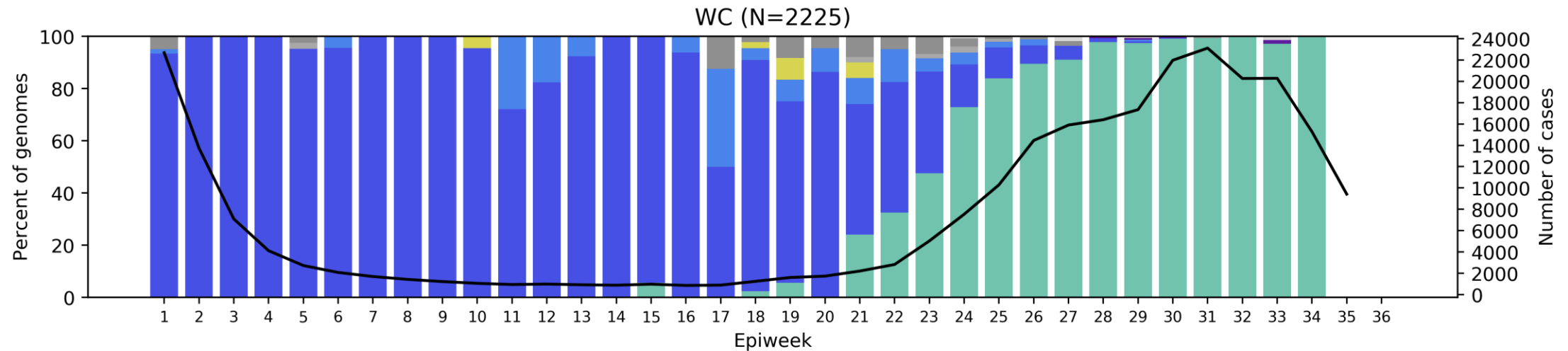
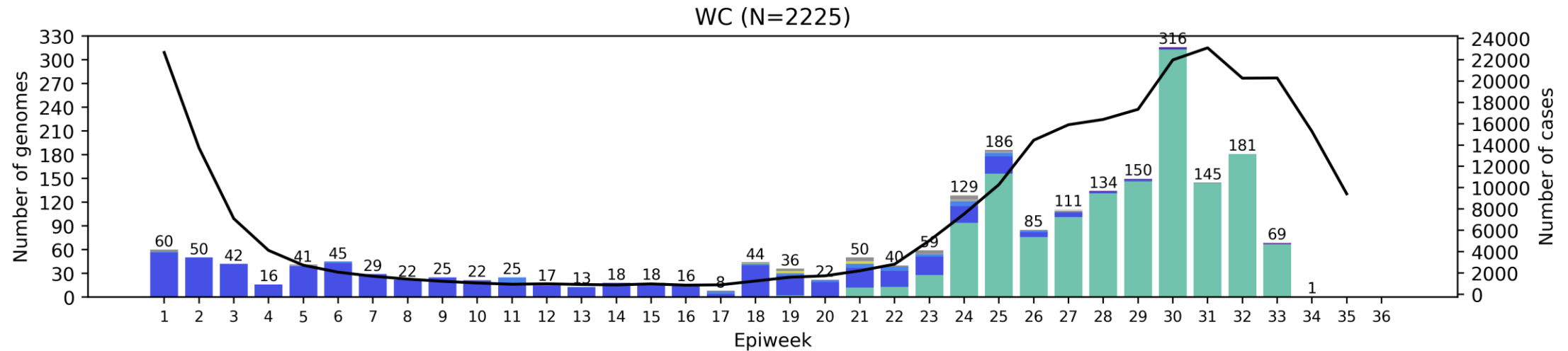
— cases Delta (21A) Beta (20H, V2) Alpha (20I, V1) C.1.2 (20D) Kappa (21B) Eta (21D) 20A 20B 20C 20D unassigned 19B

KwaZulu-Natal Province, 2021, n = 1774



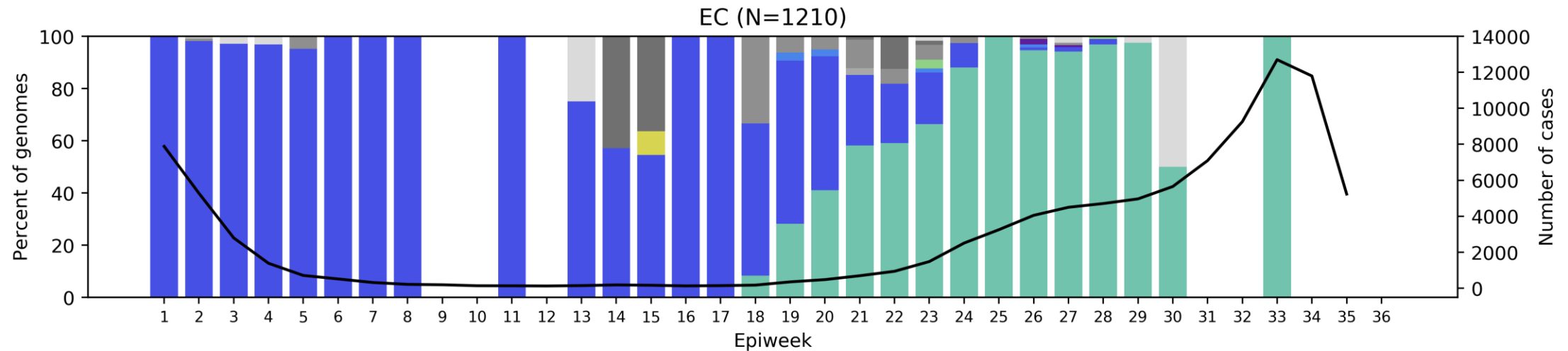
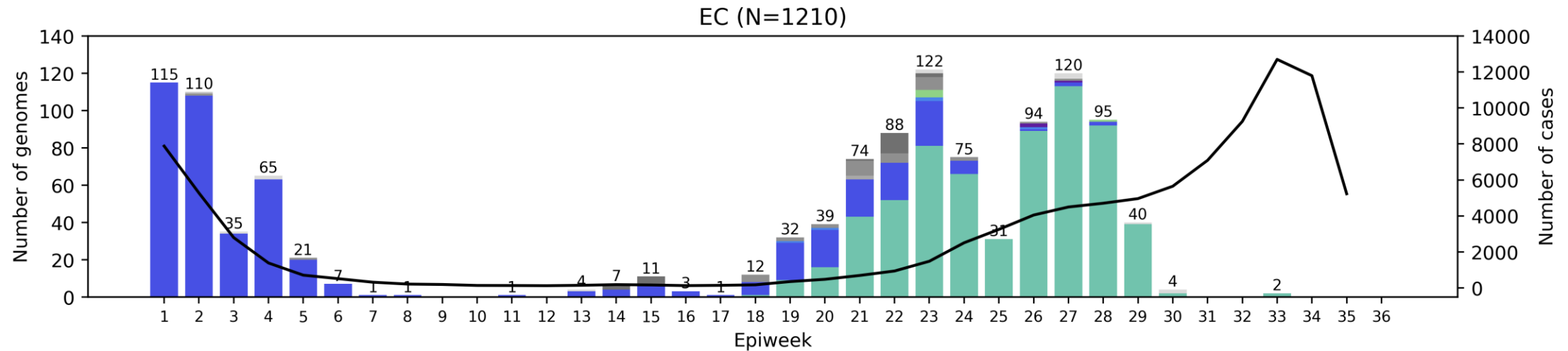
— cases Delta (21A) Beta (20H, V2) Alpha (20I, V1) C.1.2 (20D) Kappa (21B) Eta (21D) 20A 20B 20C 20D unassigned 19B

Western Cape Province, 2021, n = 2225



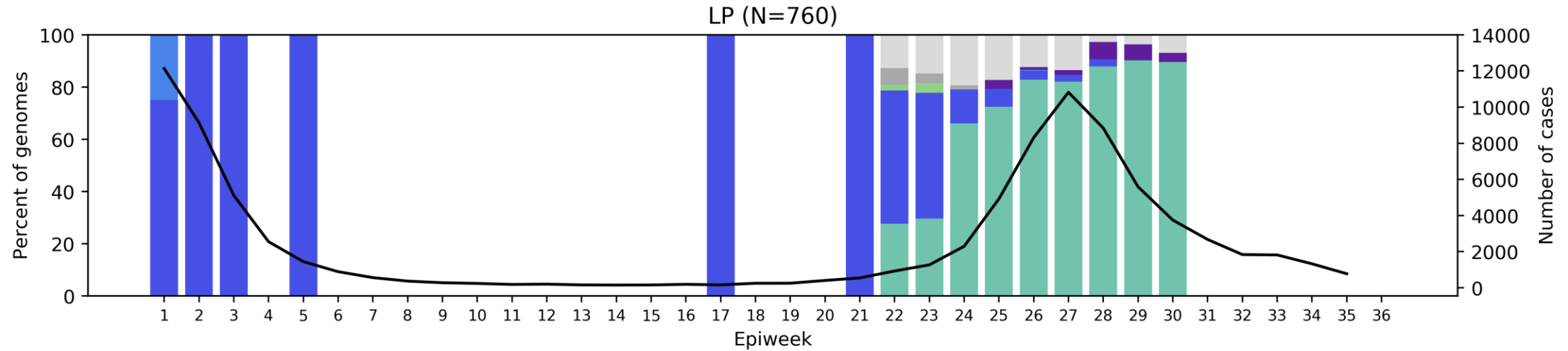
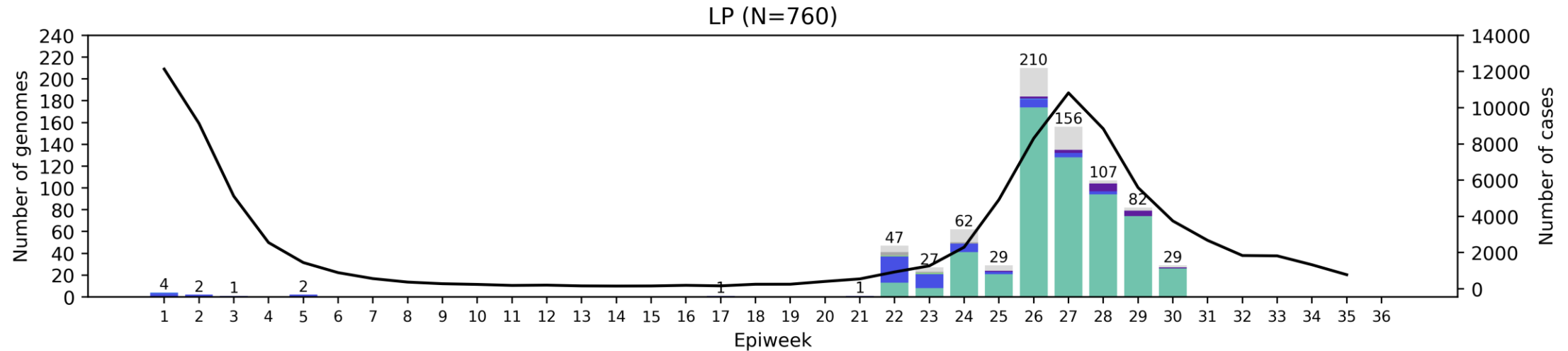
— cases Delta (21A) Beta (20H, V2) Alpha (20I, V1) C.1.2 (20D) Kappa (21B) Eta (21D) 20A 20B 20C 20D unassigned 19B

Eastern Cape Province, 2021, n = 1210



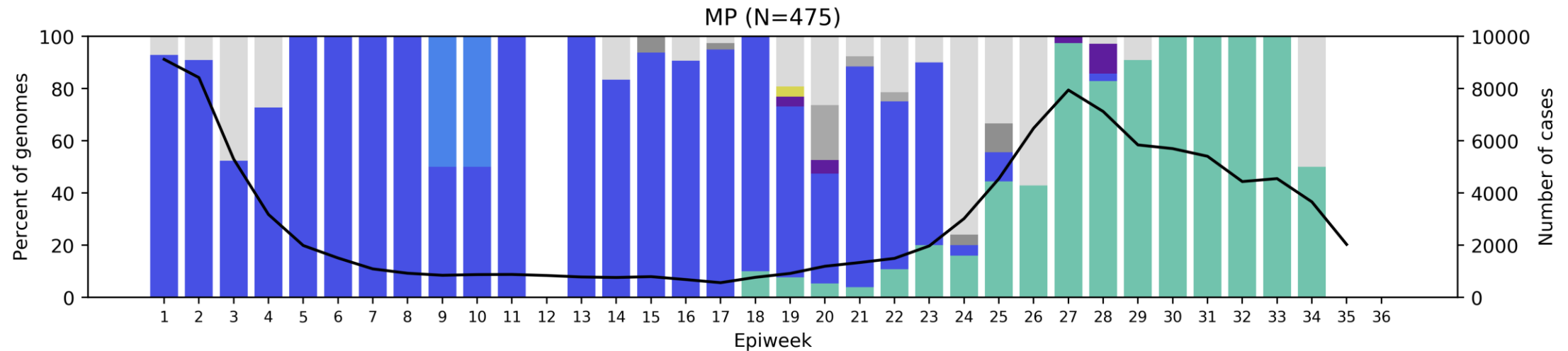
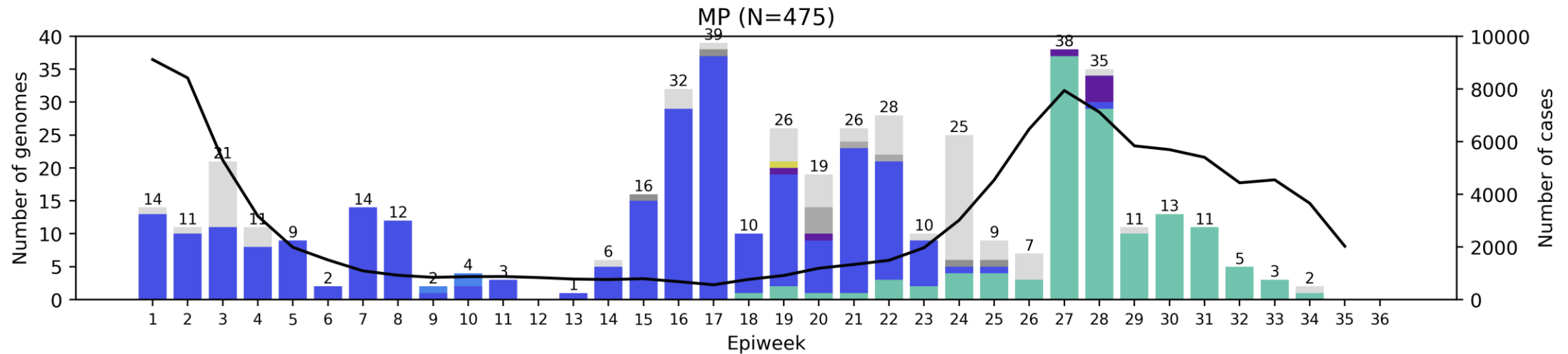
— cases Delta (21A) Beta (20H, V2) Alpha (20I, V1) C.1.2 (20D) Kappa (21B) Eta (21D) 20A 20B 20C 20D unassigned 19B

Limpopo Province, 2021, n = 760



— cases Delta (21A) Beta (20H, V2) Alpha (20I, V1) C.1.2 (20D) Kappa (21B) Eta (21D) 20A 20B 20C 20D unassigned 19B

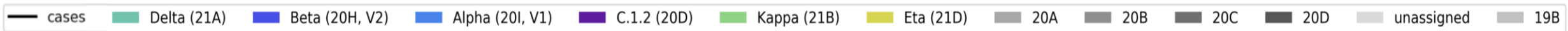
Mpumalanga Province, 2021, n = 475



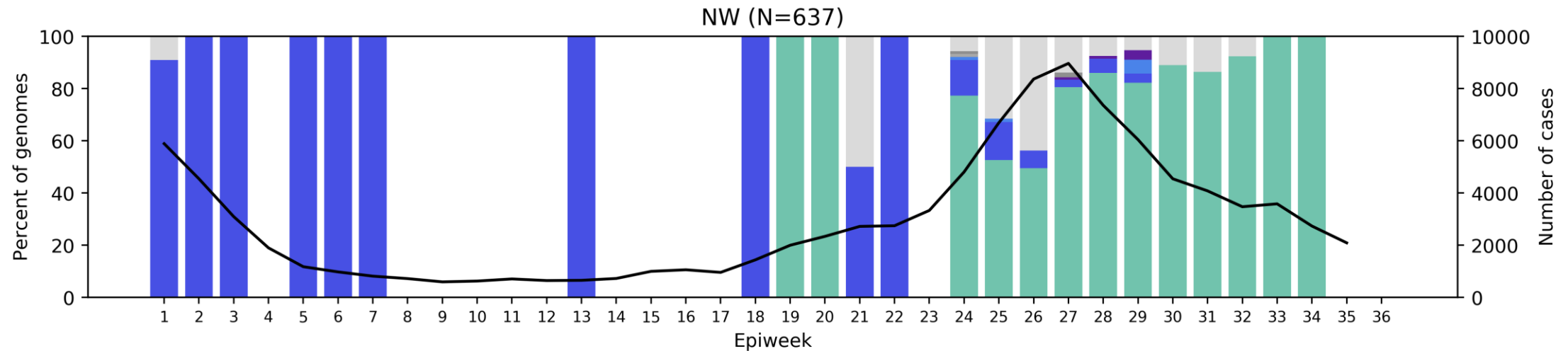
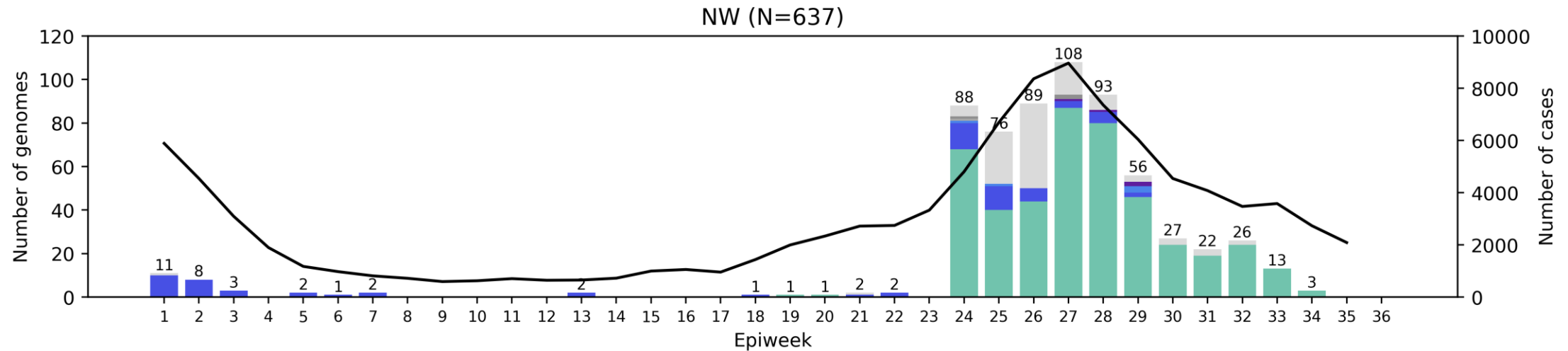
— cases Delta (21A) Beta (20H, V2) Alpha (20I, V1) C.1.2 (20D) Kappa (21B) Eta (21D) 20A 20B 20C 20D unassigned 19B

NC (N=675)

Epiweek	Number of genomes	Number of cases
1	8	2500
2	19	2200
3	27	1800
4	29	1500
5	22	1200
6	11	1000
7	9	900
8	2	800
9	15	700
10	41	1000
11	58	1200
12	48	1000
13	54	1100
14	14	1000
15	17	1500
16	54	1800
17	20	1700
18	61	2500
19	0	3000
20	0	3100
21	2	3000
22	13	2000
23	0	1800
24	1	1600
25	2	1500
26	9	1400
27	26	1500
28	23	1800
29	35	2200
30	50	2800
31	0	3500
32	0	3800
33	5	4200
34	0	3800
35	0	3200

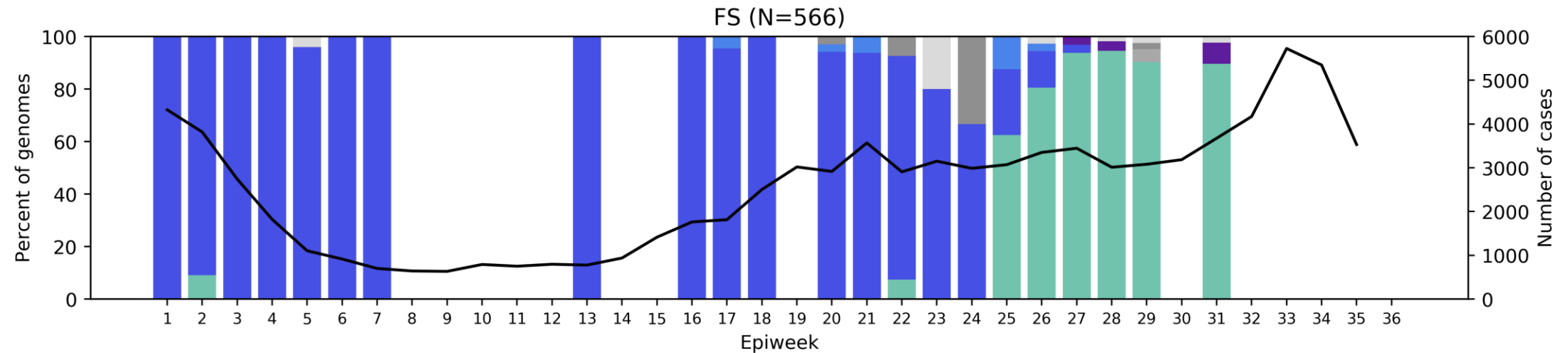
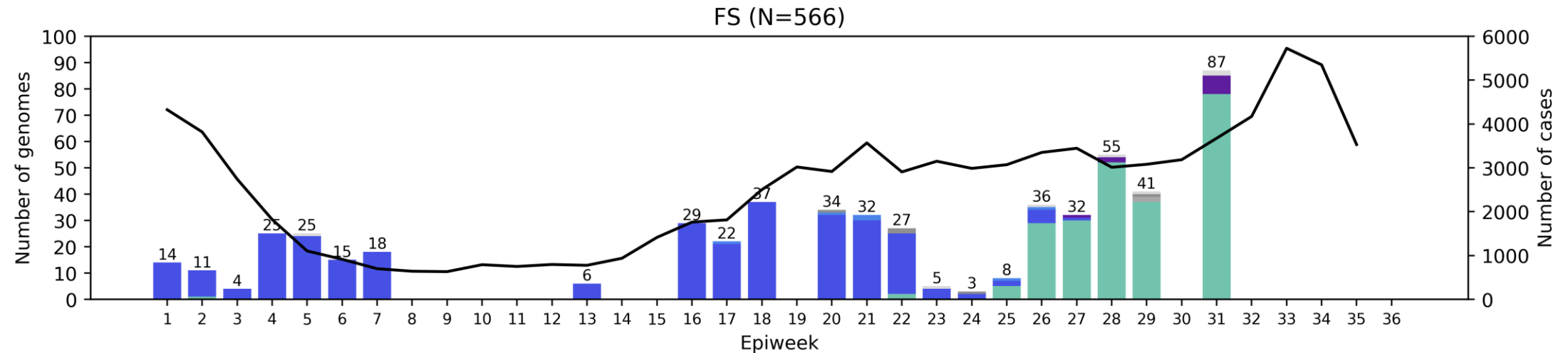


North West Province, 2021, n = 637



— cases Delta (21A) Beta (20H, V2) Alpha (20I, V1) C.1.2 (20D) Kappa (21B) Eta (21D) 20A 20B 20C 20D unassigned 19B

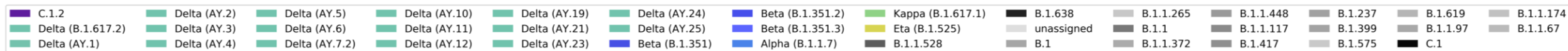
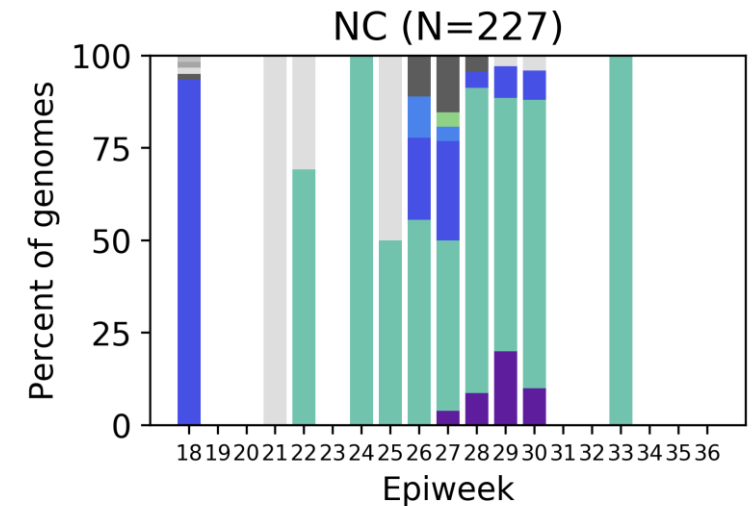
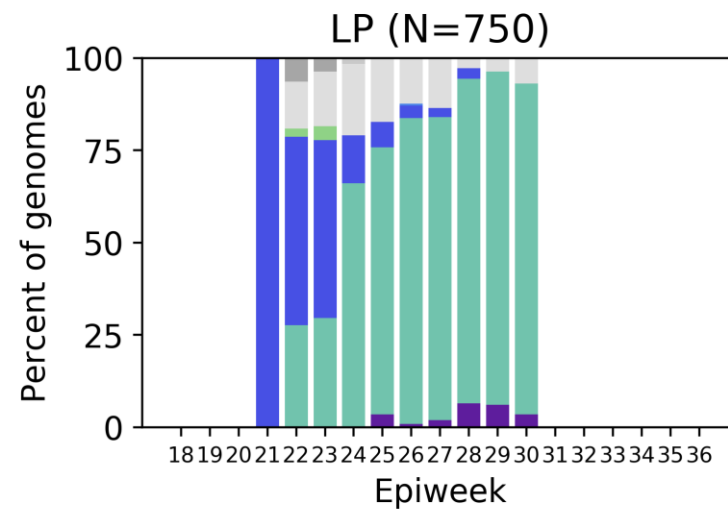
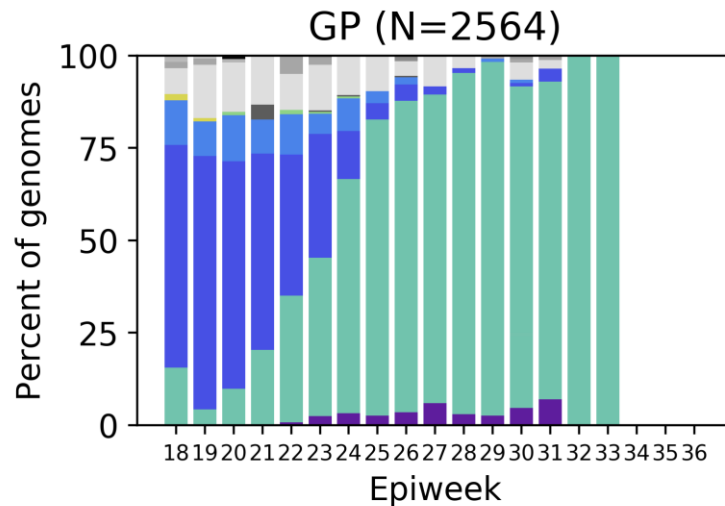
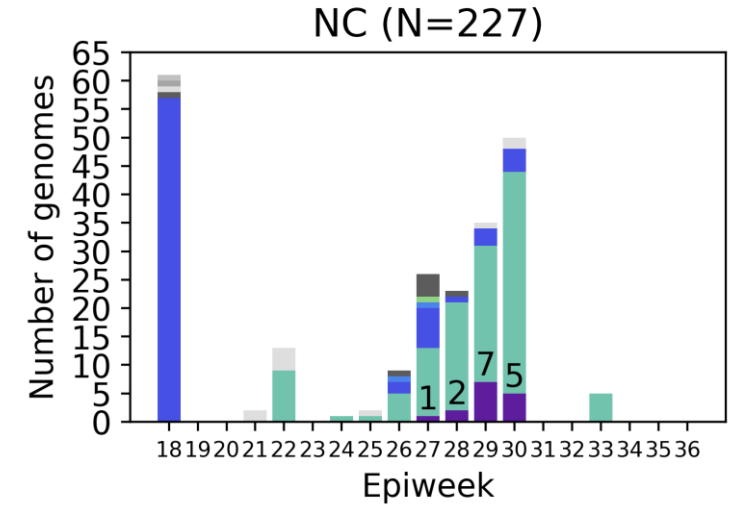
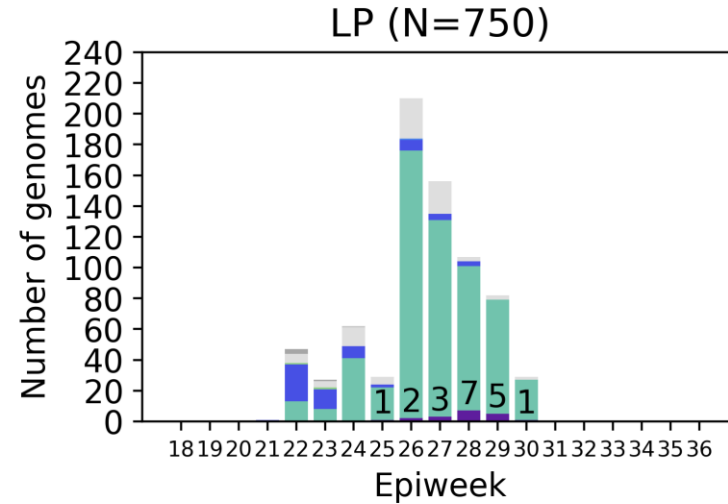
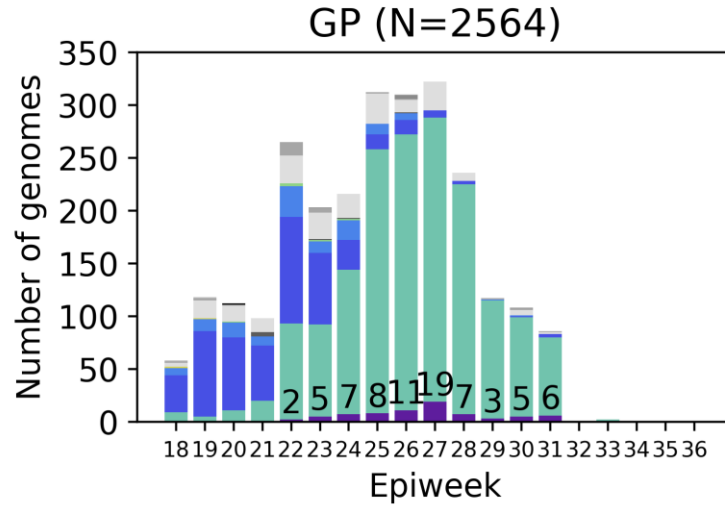
Free State Province, 2021, n = 566



— cases Delta (21A) Beta (20H, V2) Alpha (20I, V1) C.1.2 (20D) Kappa (21B) Eta (21D) 20A 20B 20C 20D unassigned 19B

C.1.2 (n=147 in SA) in May – August 2021 by epiweek

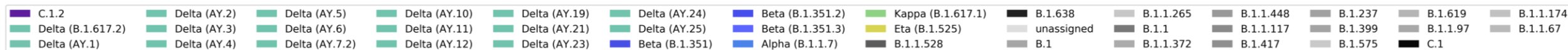
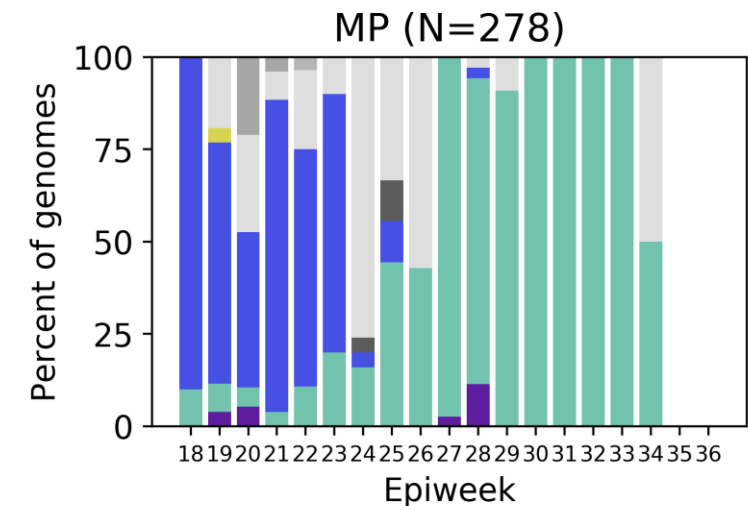
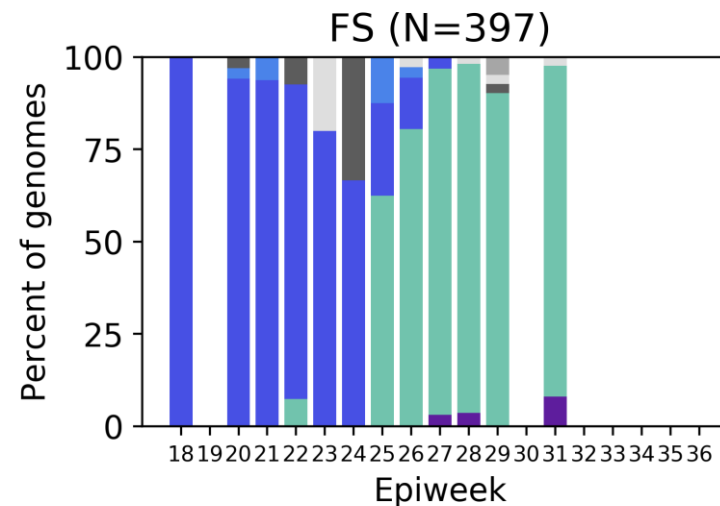
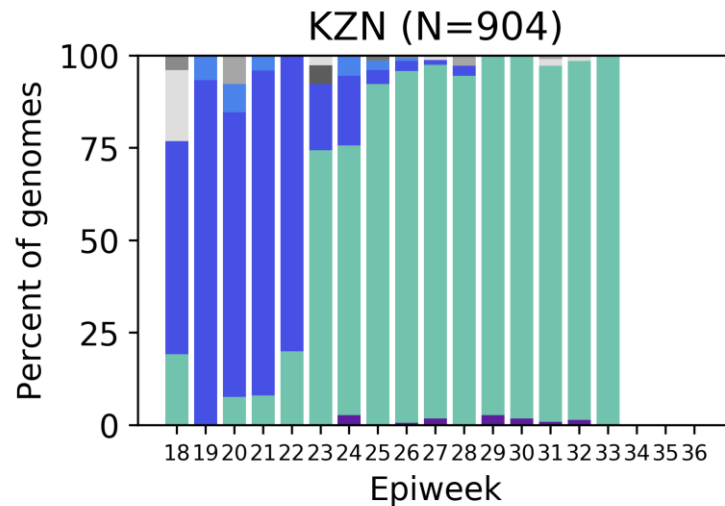
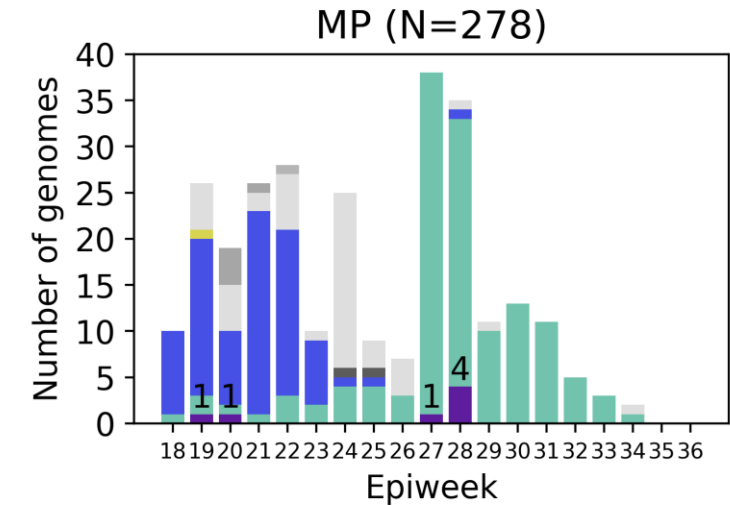
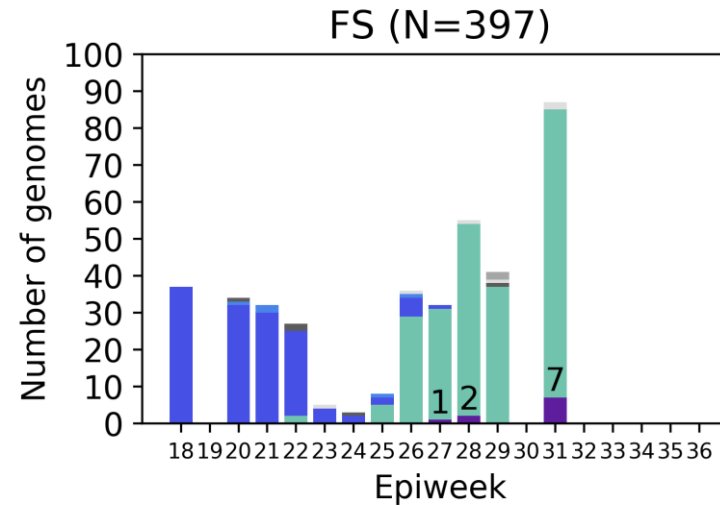
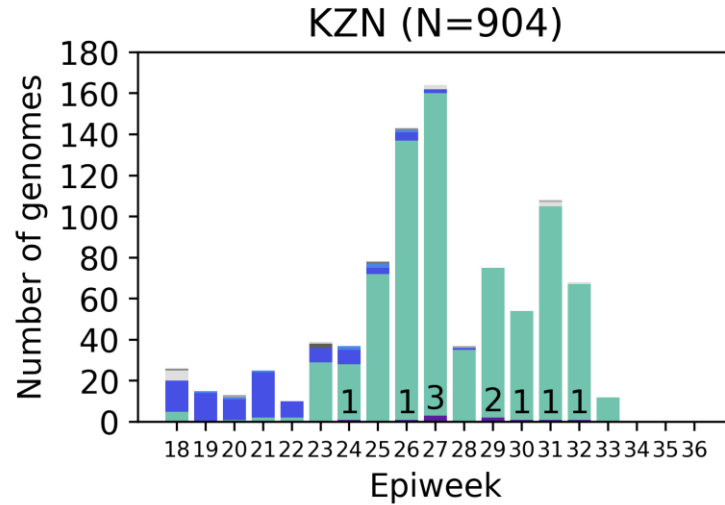
Number of C.1.2 samples indicated above bar



C.1.2 has now been detected in all provinces. The majority of samples have been detected in Gauteng (n=73), followed by Limpopo (n=19) and the Northern Cape (n=15).

C.1.2 (n=147 in SA) in May – August 2021 by epiweek

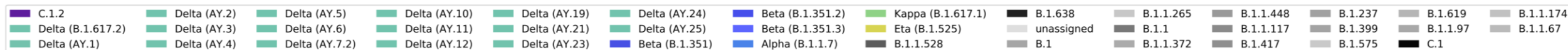
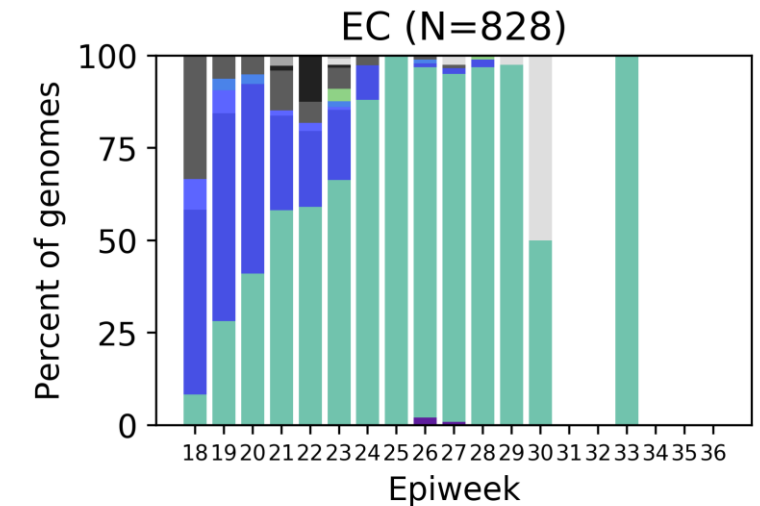
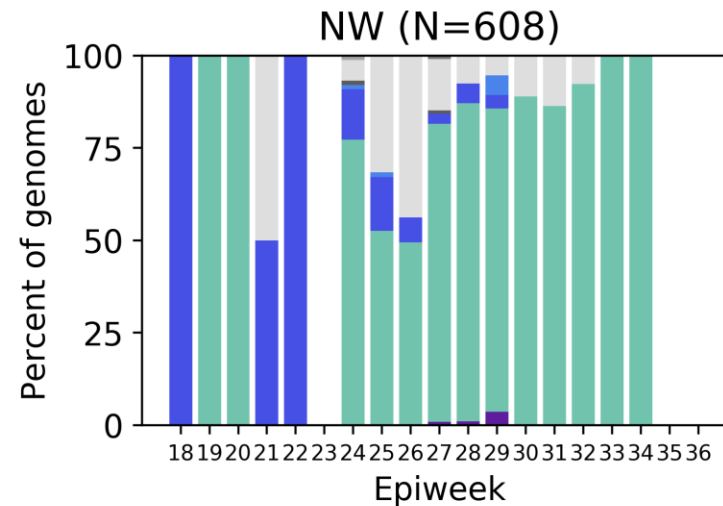
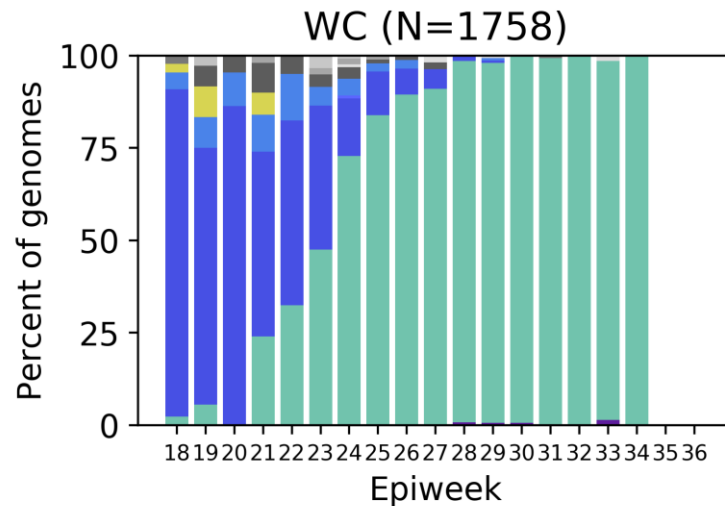
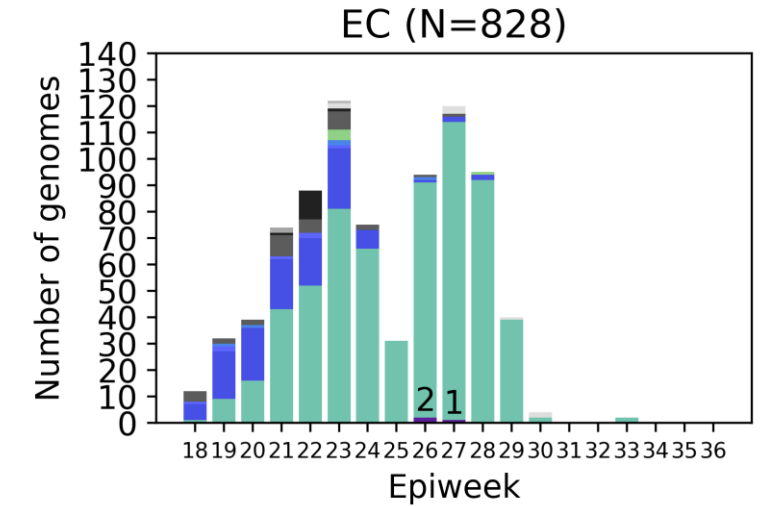
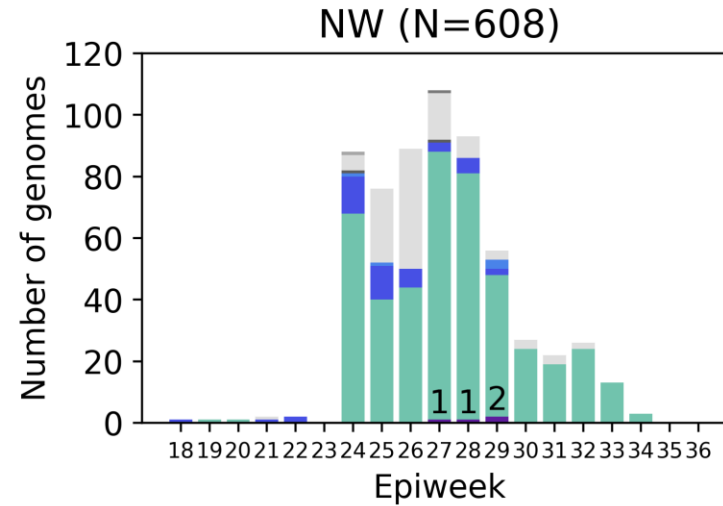
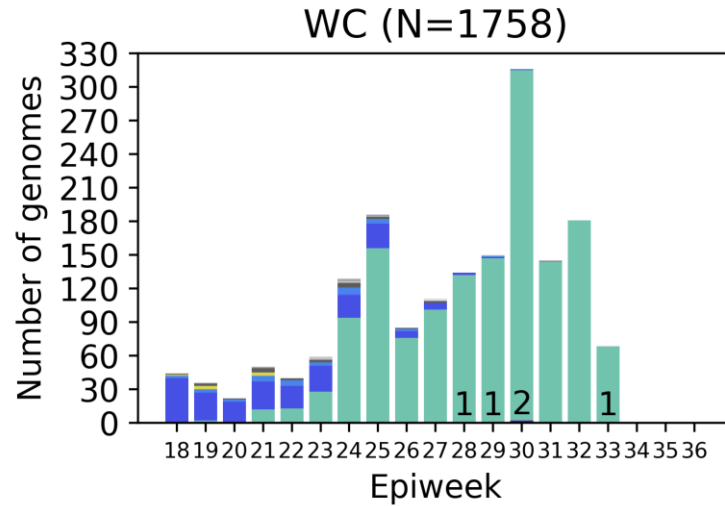
Number of C.1.2 samples indicated above bar



C.1.2 has now been detected in all provinces. 10 sequences have been detected in KwaZulu-Natal, 10 in the Free State, and 7 in Mpumalanga.

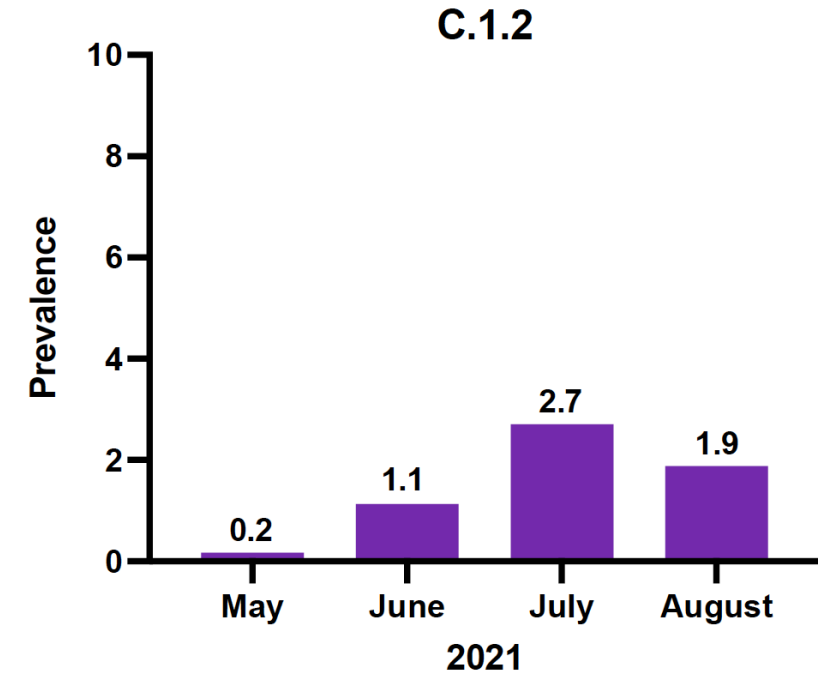
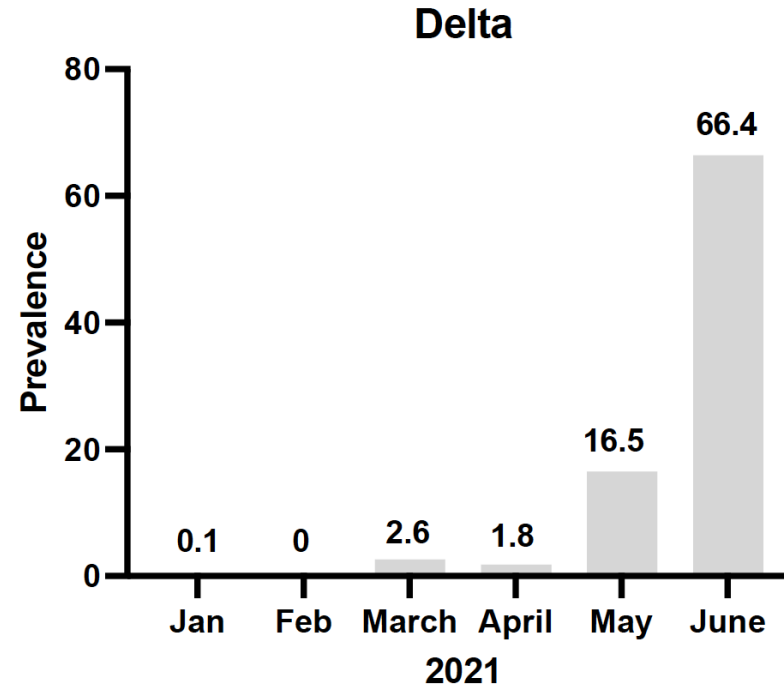
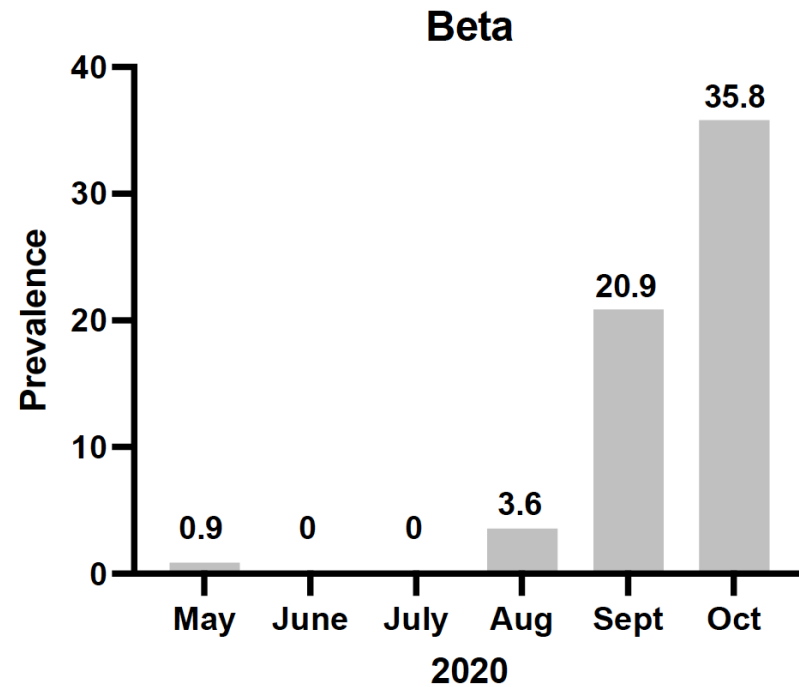
C.1.2 (n=147 in SA) in May – August 2021 by epiweek

Number of C.1.2 samples indicated above bar



The Western Cape has 5 sequences, the North West has 4, and the Eastern Cape has 3 detections of C.1.2.

C.1.2 growth compared to Beta and Delta



C.1.2 is being continually monitored and is currently only detected at low levels

Summary

- Delta continues to dominate in all provinces from specimens collected in August
- Overall diversity of lineages decreased as Delta became dominant
- Lambda and Mu variants not detected in South Africa
- Mutated C.1.2 lineage has now been detected in all provinces of South Africa
 - Frequency of C.1.2 is at less than 3% of genomes



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Dr Pravi Moodley
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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 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

University of Cape Town, NHLS & WCG



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



EDCTP



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



University of the Free State



UFS

Dominique Goedhals
Armand Bester
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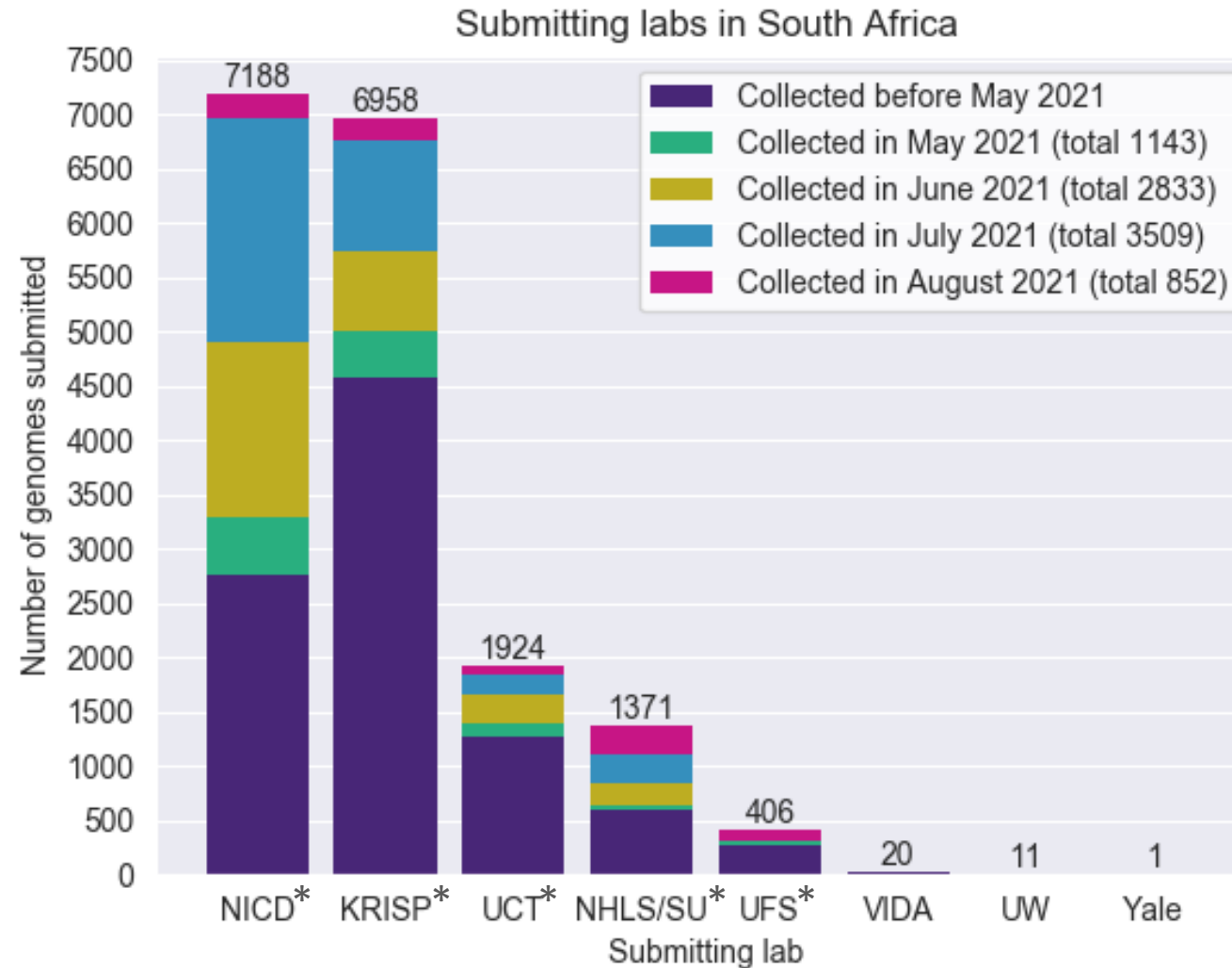
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South African genomes submitted per sequencing lab, 2020 and 2021 (N=17 879)



*NGS-SA laboratories

Multiple labs from NGS-SA are contributing to the sequencing effort.
Sequencing efforts have increased with the third wave.

Variants of Concern (VOC)

WHO label	Pango lineages ⁺	GISAID clade	Nextstrain clade	Additional amino acid changes monitored*	Earliest documented samples	Date of designation
Alpha	B.1.1.7 [#]	GRY	20I (V1)	+S:484K +S:452R	United Kingdom, Sep-2020	18-Dec-2020
Beta	B.1.351	GH/501Y.V2	20H (V2)	+S:L18F	South Africa, May-2020	18-Dec-2020
Gamma	P.1	GR/501Y.V3	20J (V3)	+S:681H	Brazil, Nov-2020	11-Jan-2021
Delta	B.1.617.2 [§]	G/478K.V1	21A	+S:417N	India, Oct-2020	VOI: 4-Apr-2021 VOC: 11-May-2021

<https://www.who.int/en/activities/tracking-SARS-CoV-2-variants/> accessed 1 September 2021

*Notable spike (S) amino acid changes under monitoring, which are currently reported in a minority of sequenced samples

⁺Includes all descendant lineages.

[#]Includes all Q.* lineages in the PANGO nomenclature system.

[§]Includes all AY.* lineages in the PANGO nomenclature system.

Currently designated Variants of Interest (VOI)

WHO label	Pango* lineages	GISAID clade	Nextstrain clade	Earliest documented samples	Date of designation
Eta	B.1.525	G/484K.V3	21D	Multiple countries, Dec-2020	17-Mar-2021
Iota	B.1.526	GH/253G.V1	21F	United States of America, Nov-2020	24-Mar-2021
Kappa	B.1.617.1	G/452R.V3	21B	India, Oct-2020	4-Apr-2021
Lambda	C.37	GR/452Q.V1	21G	Peru, Dec-2020	14-Jun-2021
Mu	B.1.631	GH	21H	Colombia, Jan-2021	30-Aug-2021

<https://www.who.int/en/activities/tracking-SARS-CoV-2-variants/> accessed 1 September 2021

* Includes all descendant lineages.

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.)