WEEKLY RESPIRATORY PATHOGENS SURVEILLANCE **REPORT**



SOUTH AFRICA WEEK 10 2022

CONTENTS

description Comments Systematic Influenza-like illness (ILI) surveillance

Influenza Respiratory syncytial virus SARS-CoV-2

Influenza-like illness (ILI) Viral 10-13 Watch

Influenza SARS-CoV-2

National syndromic 14-20 surveilance for pneumonia Influenza Respiratory syncytial virus SARS-CoV-2

Summary of laboratory 21-22 confirmed SARS-CoV-2 cases

SARS-CoV-2 Testing Methods

HIGHLIGHTS

- In 2022 to date, 17 sporadic influenza cases have been detected from Gauteng (n=1), Western Cape (n=1), Kwa-Zulu Natal (n=10) and Mpumalanga (n=5) sentinel surveillance sites.
- The 2022 RSV season started in week 7 (week starting 14 February 2022) when RSV detection rate among children aged < 5 years in pneumonia surveillance rose above the seasonal threshold and remained at low transmission, as determined by the Moving Epidemic Method.
- In 2022 to date, a total of 224 COVID-19 cases were detected from all surveillance programmes. In week 10, a decline in detection rate of COVID-19 cases has been noted in both ILI and pneumonia surveillance. Of the 137 hospitalised COVID-19 cases reported with available data on outcome, 9 (6%) died.
- · Of the 171/207 (83%) with variant data from SARI and ILI surveillance programmes, omicron variant predominated 49% (84/171), 1% (1/171) were alpha variant and for 50% (86/171) variant was not assigned.

CUMULATIVE DATA FROM





PROGRAMME DESCRIPTIONS

Programme	Influenza-like illness (ILI)	Viral Watch	National syndromic surveillance for pneumonia
Start year	2012	1984	2009
Provinces*	KZ NW WC** MP***	EC FS GP LP MP NC NW	GP KZ MP NW WC
Type of site	Primary health care clinics	General practitioners	Public hospitals
Case definition	ILI: An acute respiratory illness with a temperature (≥38°C) and cough, & onset ≤10 days Suspected pertussis Any person with an acute cough illness lasting ≥14 days (or cough illness of any duration for children <1 year), without a more likely diagnosis AND one or more of the following signs or symptoms: • paroxysms of coughing, • or inspiratory "whoop", • or post-tussive vomiting • or apnoea in children <1 year; OR Any person in whom a clinician suspects pertussis Suspected SARS-CoV-2 Any person presenting with an acute	ILI: An acute respiratory illness with a temperature (≥38°C) and cough, & onset ≤10 days Suspected SARS-CoV-2 Any person presenting with an	SRI: Acute (symptom onset≤10 days) or chronic (symptom onset >10) lower respiratory tract infection Suspected pertussis Any person with an acute cough illness lasting ≥14 days (or cough illness of any duration for children <1 year), without a more likely diagnosis AND one or more of the following signs or symptoms: • paroxysms of coughing, • or inspiratory "whoop", • or post-tussive vomiting • or apnoea in children <1 year; OR Any person in whom a clinician suspects pertussis. Suspected SARS-COV-2 Any person admitted with a physician-
Specimens	(≤14 days) respiratory tract infection or other clinical illness compatible with COVID-19 ^β Oropharyngeal & nasopharyngeal	acute (≤14 days) respiratory tract infection or other clinical illness compatible with COVID-19 [®] Throat and/or nasal swabs or	diagnosis of suspected COVID-19 and not meeting SRI case definition. Oropharyngeal & nasopharyngeal
collected	swabs	Nasopharyngeal swabs	swabs
Main pathogens tested****	INF RSV BP SARS-CoV-2	INF RSV BP SARS-CoV-2	INF RSV BP SARS-CoV-2
Testing Methods	INF and RSV - Fast-Track Diagnostics multiplex real-time reverse transcription polymerase chain reaction (until 31 March 2021) B. pertussis Multiplex real-time PCR (Tatti et al., J Clin Microbiol 2011) and culture (if PCR cycle threshold ≤25) SARS-CoV-2	INF and RSV - Fast-Track Diagnostics multiplex real-time reverse transcription polymerase chain reaction (until 31 March 2021) B. pertussis Multiplex real-time PCR (Tatti et al., J Clin Microbiol 2011) and culture (if PCR cycle threshold ≤25) SARS-CoV-2	INF and RSV - Fast Track Diagnostics multiplex real- time reverse transcription polymerase chain reaction (until 31 March 2021) B. pertussis Multiplex real-time PCR (Tatti et al., J Clin Microbiol 2011) and culture (if PCR cycle threshold ≤25) SARS-CoV-2 1 April 2020 – 31 March 2021: Roche E
	1 April 2020 – 31 March 2021: Roche E gene real-time PCR essay (Corman et al., Euro Surv 2020) 1 April 2021 to date: Allplex™ SARS- CoV-2/FluA/FluB/RSV PCR kit • positivity assigned if PCR cycle threshold is <40 for ≥1 gene targets (N, S, OR RdRp)	I April 2020 – 31 March 2021: Roche E gene real-time PCR essay Corman et al., Euro Surv 2020) 1 April 2021 to date: Allplex™ SARS- CoV-2/FluA/FluB/RSV PCR kit • positivity assigned if PCR cycle threshold is <40 for ≥1 gene targets (N, S, OR RdRp)	gene real-time PCR essay (Corman et al., Euro Surv 2020) 1 April 2021 to date: Allplex™ SARS-CoV-2/FluA/FluB/RSV PCR kit • positivity assigned if PCR cycle threshold is <40 for ≥1 gene targets (N, S, OR RdRp)

Epidemic Threshold

org/web/package=mem) designed to calculate the duration, start and end of the annual influenza epidemic. MEM uses the 40th, 90th and 97.5th $percentiles\ established\ from\ available\ years\ of\ historical\ data\ to\ calculate\ thresholds\ of\ activity.\ Thresholds\ of\ activity\ for\ influenza\ and\ RSV\ are\ defined\ percentiles\ and\ respectively.$ as follows: Below seasonal threshold, Low activity, Moderate activity, High activity, Very high activity. For influenza, thresholds from outpatient influenza like illness (ILI in primary health care clinics) are used as an indicator of disease transmission in the community and thresholds from pneumonia surveillance are used as an indicator of impact of disease. For RSV, thresholds from pneumonia surveillance, using data from children aged < 5 years

- * EC: Eastern Cape; FS: Free State; GP: Gauteng; KZ: KwaZulu-Natal; LP: Limpopo; MP: Mpumalanga: NC: Northern Cape; NW: North West; WC: Western Cape
- ***Started in November 2020
- symptoms (which may include fever, weakness, myalgia, or diarrhoea). Testing for SARS-CoV-2 was initiated in all three surveillance programmes in week 10 of 2020 (week starting 2 March 2020)

COMMENTS

Influenza

In 2022 to date, a total of 17 influenza cases have been reported. In week 10, transmission and impact were below threshold.

ILI programme: In 2022 to date, specimens from 313 patients meeting ILI case definition were tested from 4 ILI sites. Influenza was detected in eight (3%) patients, all were influenza A(H1N1)pdm09. (Fig1, Table1).

Viral Watch programme: In 2022 to date, specimens from 38 patients from three of the 8 provinces participating in Viral Watch surveillance were tested. Influenza was detected in one (3%) and influenza A subtype results are still pending. (Fig7, Table5)

Pneumonia surveillance: Since the beginning of 2022, specimens from 1 099 patients with severe respiratory illness (SRI) were tested from the 6 sentinel sites. Influenza was detected in seven (<1%) patients, of which six (86%) were influenza A(H1N1)pdm09 and influenza A subtype was pending for one (14%). (Fig12, Table9)

In addition, 30 specimens from patients who met suspected SARS-CoV-2 case definition but did not meet the pneumonia/ ILI surveillance case definitions were tested for influenza. One (3%) tested positive for influenza B(Victoria).

Respiratory syncytial virus

The 2022 RSV season started in week 7 (week starting 14 February 2022) when RSV detection rate among children aged < 5 years in pneumonia surveillance rose above the seasonal threshold and has remained at low transmission, as determined by the Moving Epidemic Method.

ILI programme: In 2022 to date, 313 specimens from patients meeting the ILI case definition were tested and RSV was detected in 25 (8%) patients. Of which, 19 (76%) were RSV subgroup A and six (24%) were pending RSV subgroup results. (Fig4, Table2)

Viral Watch programme: In 2022 to date, 38 specimens from Viral Watch patients were tested and RSV was not detected. (Fig9, Table6)

Pneumonia surveillance: Since the beginning of 2022, 1 099 specimens were tested and RSV was detected in specimens of 105 (10%) patients. Of which, 16 (15%) were RSV subgroup A, 44 (42%) were RSV subgroup B, and 45 (43%) were pending RSV subgroup results. (Fig14, Table10)

In addition, 30 specimens from patients who met suspected SARS-CoV-2 case definition but did not meet the pneumonia/ ILI surveillance case definitions were tested for RSV and none tested positive for RSV.

SARS-CoV-2 (Severe acute respiratory syndrome coronavirus 2)

ILI programme: From 3 January 2022 to date, 313 patients were tested and SARS-CoV-2 was detected in 50 (16%) patients. Of the 40 (40/50, 80%) with variant data, most common variant was omicron (11/40; 28%) and for 72% (29/40) variant was not assigned. (Fig6, Table4)

Viral Watch programme: From 3 January 2022 to date, 38 patients presenting with ILI were tested and SARS-CoV-2 was detected in 17 (45%). Of the 9 (9/17, 53%) with variant data, majority were omicron variant (5/9, 56%) and variant was not assigned for 44% (4/9). (Fig11, Table8)

Pneumonia surveillance: From 3 January 2022 to date, 1 099 patients with severe respiratory illness (SRI) were tested and SARS-CoV-2 was detected in 145 (14%) patients. Of the 119 (119/145, 82%) with variant data, majority were omicron variant 55% (65/119), 1% (1/119) was Alpha variant and the variant not assigned for was 45% (53/119). (Fig17, Table12)

In addition, SARS-CoV-2 was detected in 12 of 30 (40%) specimens from patients who met suspected SARS-CoV-2 case definition but did not meet the pneumonia/ILI surveillance case definitions. Of this 12 with variant data, 67% (8/12) were omicron variant and variant was not assigned for 33% (4/12).

WEEK 10 2022

INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE PRIMARY HEALTH CARE CLINICS

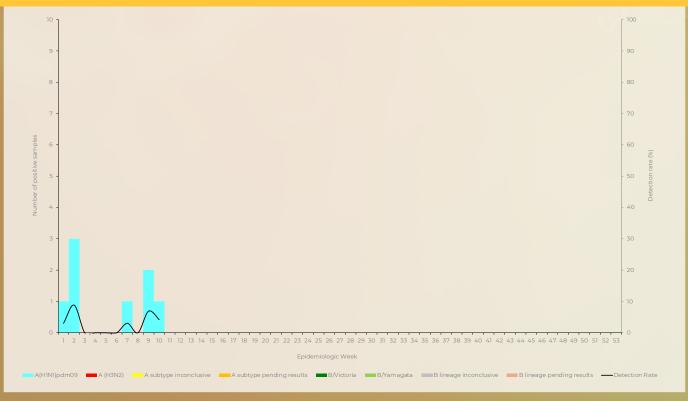


Figure 1. Number of influenza positive cases* by influenza subtype and lineage** and detection rate*** by week, Influenza-like illness (II I) surveillance in primary health care clinics 03/01/2022 – 13/03/2022

Table 1. Number of laboratory confirmed influenza cases by subtype and lineage** and total number of samples tested by clinic and province, Influenza-like illness (ILI) surveillance in primary health care clinics, 03/01/2022 - 13/03/2022

Clinic (Province)	A(H1N1) pdm09	A(H3N2)	A subtype inconclusive	A subtype pending results [§]	B/ Victoria	B/ Yamagata	B lineage inconclusive	B lineage pending results [§]	Total samples
Agincourt (MP)	3	0	0	0	0	0	0	0	49
Eastridge (WC)	О	О	0	0	0	0	0	0	42
Edendale Gateway (KZ)	5	0	О	0	0	0	0	0	103
Jouberton (NW)	0	О	0	0	0	0	0	0	101
Mitchell's Plain (WC)	0	О	0	0	0	0	0	0	18
Total:	8	0	0	0	0	0	0	0	313

KZ: KwaZulu-Natal; NW: North West; WC: Western Cape; MP: Mpumalanga

^{*}Specimens from patients with influenza-like illnesses at 5 sentinel sites in 4 provinces

^{***}Only reported for weeks with >10 specimens submitted

Inconclusive: insufficient viral load in sample and unable to characterise further

^{**}Influenza B(Victoria) was detected in one (6%) of 18 specimens from patients who met suspected SARS-CoV-2 case definition but did not meet Influenza-like illness (ILI

Inconclusive: insufficient viral load in sample and unable to characterise further

finfluenza A subtype or B lineage results are pending

^{**}Influenza B(Victoria) was detected in one (6%) of 18 specimens from patients who met suspected SARS-CoV-2 case definition but did not meet Influenza-like illness (ILI) case definition and it was influenza B(Victoria). These are not included in the table.

WEEK 10 2022

INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE PRIMARY HEALTH CARE CLINICS



Figure 2. Influenza percentage detections and epidemic thresholds* among cases of all ages, Influenza-like illness (ILI) surveillance in primary health care clinics, 03/01/2022 – 13/03/2022

*Thresholds based on 2012-2019 data

WEEK 10 2022

INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE PRIMARY HEALTH CARE CLINICS



Figure 3. Number of patients testing positive for respiratory syncytial virus* by province and detection rate by week, Influenza-like illness (ILI) surveillance in primary health care clinics, 03/01/2022 – 13/03/2022

*RSV was not detected from 18 specimens of patients who met suspected SARS-CoV-2 case definition but did not meet influenza-like illness (ILI) case definition.



WEEK 10 2022

INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE PRIMARY HEALTH CARE CLINICS

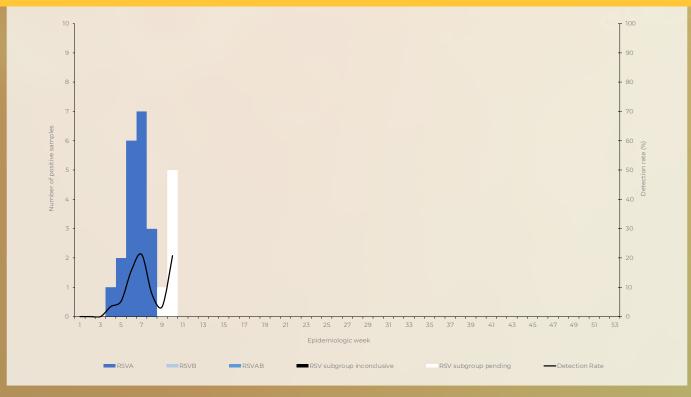


Figure 4. Number of patients testing positive for respiratory syncytial virus* by subgroup and detection rate by week, Influenza-

Table 2. Number of patients testing positive for respiratory syncytial virus (RSV)** by subgroups identified and total number of samples tested by clinic and province, Influenza-like illness (ILI) surveillance in primary health care clinics, 03/01/2022 –

Clinic (Province)	RSVA	RSVB	RSVAB	RSV subgroup inconclusive	RSV subgroup pending*	Total samples
Agincourt (MP)		0	0	0	0	49
Eastridge (WC)	0	0	0	0	0	42
Edendale Gateway (KZ)	18	0	0	0	6	103
Jouberton (NW)	0	0	0	0	0	101
Mitchell's Plain (WC)	0	0	0	0	0	18
Total	19	0	0	0	6	313

Inconclusive: insufficient viral load in sample and unable to characterise further RSV AB: Both RSV A and B subgroup identified

*RSV results for subgroups are performing
**RSV was not detected from 18 specimens of patients who met suspected SARS-CoV-2 case definition but did not meet influenza-like illness (ILI) case definition. These

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INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE PRIMARY HEALTH CARE CLINICS



Figure 5. Number of patients testing positive for SARS-CoV-2* by province and detection rate by week, Influenza-like illness (ILI)

Table 3. Number of patients positive for SARS-CoV-2* identified and total number of samples tested by clinic and province, Influenza-like illness (ILI) surveillance primary health care clinics, 03/01/2022 - 13/03/2022

Clinic (Province)	SARS-CoV-2 positive	Total samples tested
Agincourt (MP)	19	49
Eastridge (WC)	3	42
Edendale Gateway (KZ)	5	103
Jouberton (NW)	19	101
Mitchell's Plain (WC)	4	18
Total:	50	313

KZ: KwaZulu-Natal; NW: North West; WCP: Western Cape; MP: Mpumalanga *SARS-CoV-2 was detected in 5 of 18 (28%) specimens from patients who met suspected SARS-CoV-2 case definition but did not meet influenza-like illness (ILI) case definition. These are not included in the table.

WEEK **10** 2022

INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE PRIMARY HEALTH CARE CLINICS



Figure 6. Number and detection rate of laboratory confirmed SARS-CoV-2* cases by variant type (variant PCR/sequencing) and

Unable to assign: no lineage assigned due to poor- sequence quality OR low viral load (ct=>35) OR variant PCR could not assign variant and no sequencing result Pending: outstanding variant results

Table 4. Number of SARS-CoV-2* positive cases by variant (variant PCR and/or sequencing) identified and total number of samples tested by clinic and province, Influenza-like illness (ILI) surveillance primary health care clinics, 03/01/2022 - 13/03/2022

Clinic (Province)	Non- Alpha/ Beta/ Delta	Alpha (201)	Beta (20H)	Delta (21A, 21I, 21J)	C.1.2 (20D)	Omicron (21K, 21L, 21M)	Pending	Unable to assign	Total SARS- CoV-2 positive
Agincourt (MP)	0	0	0	0	0	5	2	14	21
Eastridge (WC)	0	0	0	О	0	2	0		3
Edendale Gateway (KZ)	0	О	О	0	0	2		4	7
Jouberton (NW)	0	О	0	О	0	3	7	10	20
Mitchell's Plain (WC)	0	0	0	0	0	2	0	2	4
Total:	0	0	0	0	0	14	10	31	55

Unable to assign: no lineage assigned due to poor- sequence quality OR low viral load (ct=>35) OR variant PCR could not assign variant and no sequencing result Pending: outstanding variant results

KZ: KwaZulu-Natal; NW: North West; WCP: Western Cape; MP: Mpumalanga
*Specimens are from patients with influenza-like illness at 5 sentinel sites in 4 provinces who met suspected SARS-CoV-2 case definition or met ILI case definition

WEEK 10 2022

INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE VIRAL WATCH

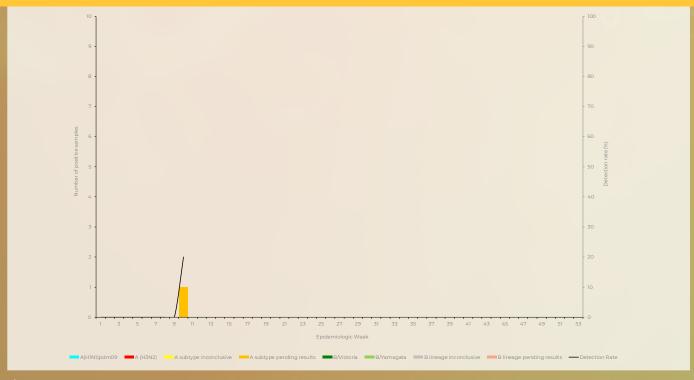


Figure 7. Number of positive patients* by influenza subtype and lineage and detection rate** by week, ILI surveillance - Viral Watch, 03/01/2022 – 13/03/2022

Inconclusive: insufficient viral load in sample and unable to characterise further

Table 5. Number of laboratory confirmed influenza cases by influenza subtype and lineage and total number of samples tested by province, ILI surveillance - Viral Watch, 03/01/2022 – 13/03/2022

Province	A(H1N1) pdm09	A(H3N2)	A subtype inconclusive	A subtype pending results*	B/ Victoria	B/ Yamagata	B lineage inconclusive	B lineage pending results*	Total samples
Eastern Cape	0	0	0	0	0	О	0	0	
Free State	Ο	О	0	O	0	O	0	О	0
Gauteng	Ο	О	0		0	О	0	О	26
Limpopo	Ο	О	0	О	0	O	0	0	0
Mpumalanga	0	О	0	0	0	О	0	О	0
North West	0	0	0	0	0	0	0	0	0
Northern Cape	0	О	0	0	0	0	0	0	О
Western Cape	0	0	0	0	0	0	0	0	11
Total:	0	0	0	1	0	0	0	0	38

Inconclusive: insufficient viral load in sample and unable to characterise further *Influenza A subtype or B lineage results are pending

^{*}Specimens from patients with Influenza-like illnesses at 90 sentinel sites in 8 provinces

^{**} Only reported for weeks with >10 specimens submitted.

WEEK 10 2022

INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE VIRAL WATCH



Figure 8. Number of RSV positive cases testing positive for respiratory syncytial virus (RSV)* by subgroup and detection rate** by week, ILI surveillance - Viral Watch, 03/01/2022 – 13/03/2022

Table 6. Number of RSV positive cases identified and total number of samples tested by province, ILI surveillance - Viral Watch, 03/01/2022 – 13/03/2022

Province	RSV A	RSV B	RSV AB	RSV subgroup inconclusive**	RSV subgroup pending results*	Total samples tested
Eastern Cape	0	0	0	0	0	
Free State	О	О	0	0	0	0
Gauteng	О	О	0	0	0	26
Limpopo	0	0	0	0	0	0
Mpumalanga	О	0	0	0	0	0
North West	0	0	0	0	0	0
Northern Cape	0	О	0	0	0	0
Western Cape	0	0	0	0	0	11
Total:	0	0	0	0	0	38

^{*}RSV results for subgroups are pending

^{*}Specimens from patients with Influenza-like illnesses at 92 sentinel sites in 8 provinces

^{**} Only reported for weeks with >10 specimens submitted

^{**}Inconclusive: insufficient viral load in sample and unable to characterise further

WEEK 10 2022

INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE VIRAL WATCH

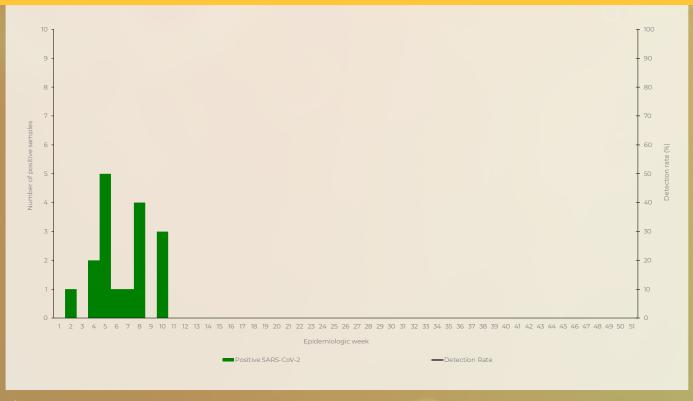


Figure 9. Number of patients testing positive for SARS-CoV-2*, by site and detection rate** by week, ILI surveillance - Viral Watch, 03/01/2022 – 13/03/2022

Table 7. Number of SARS-CoV-2 positive cases identified and total number tested by province, ILI surveillance - Viral Watch, 03/01/2022 – 13/03/2022

Province	SARS-CoV-2 positive	Total samples tested
Eastern Cape	1 1 10 7	1
Free State	0	0
Gauteng	12	26
Limpopo	0	0
Mpumalanga	0	0
North West	0	0
Northern Cape	0	0
Western Cape	4	11
Total:	17	38

^{*}Specimens from patients with Influenza-like illnesses at 92 sentinel sites in 8 provinces

^{**} Only reported for weeks with >10 specimens submitted

WEEK 10 2022

INFLUENZA-LIKE ILLNESS (ILI) SURVEILLANCE VIRAL WATCH



Figure 10. Number and detection rate of laboratory confirmed SARS-CoV-2* cases by variant type (variant PCR/sequencing) and week, ILI surveillance - Viral Watch, 03/01/2022 – 13/03/2022

Unable to assign: no lineage assigned due to poor- sequence quality OR low viral load (ct=>35) OR variant PCR could not assign variant and no sequencing result Pending: outstanding variant results

Table 8. Number of SARS-CoV-2* positive cases by variant (variant PCR and/or sequencing) identified and total number of samples tested by province, ILI surveillance - Viral Watch, 03/01/2022 - 13/03/2022

Clinic (Province)	Non-Alpha/ Beta/Delta	Alpha (20I)	Beta (20H)	Delta (21A, 21I, 21J)	C.1.2 (20D)	Omicron (21K, 21L, 21M)	Pending	Unable to assign	Total SARS-CoV-2 positive
Eastern Cape	0	0	0	0	0	0		0	
Free State	О	О	0	О	0	0	Ο	О	0
Gauteng	О	О	Ο	О	0	3	5	4	12
Limpopo	О	О	Ο	О	0	0	Ο	О	0
Mpumalanga	О	0	Ο	0	0	0	Ο	О	0
North West	О	Ο	0	О	0	0	Ο	О	0
Northern Cape	0	Ο	0	О	0	0	Ο	О	0
Western Cape	О	О	0	О	0	2	2	0	4
Total:	0	О	0	0	0	5	8	4	17

^{*}Specimens from patients with Influenza-like illnesses at 92 sentinel sites in 8 provinces

Unable to assign: no lineage assigned due to poor- sequence quality OR low viral load (ct=>35) OR variant PCR could not assign variant and no sequencing result **Pending:** outstanding variant results

^{*}Specimens from patients with Influenza-like illnesses at 92 sentinel sites in 8 provinces

NATIONAL SYNDROMIC SURVEILLANCE FOR PNEUMONIA

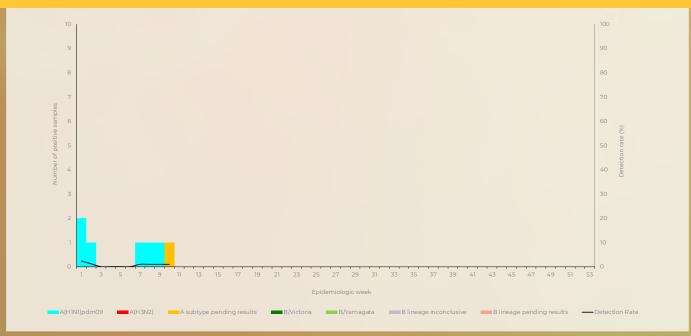


Figure 11. Number of positive influenza positive cases* by influenza subtype and lineage** and detection rate*** by week,

Table 9. Number of laboratory confirmed influenza cases by subtype and lineage* and total number of samples tested by hospital, pneumonia surveillance public hospitals, 03/01/2022 – 13/03/2022

Hospital (Province)	A(H1N1) pdm09	A(H3N2)	A subtype inconclusive	A subtype pending results***	B/ Victoria	B/ Yamagata	B lineage inconclusive	B lineage pending results***	Total samples
Edendale (KZ)	3	0	0	- 1	0	0	0	0	216
Helen Joseph- Rahima Moosa (GP)	0	0	0	0	0	0	0	0	304
Klerksdorp- Tshepong (NW)	0	0	0	0	0	0	О	0	106
Mapulaneng- Matikwana (MP)	0	0	0	0	0	0	О	0	97
Red Cross (WC)	0	0	0	0	0	0	0	0	183
Mitchell's Plain (WC)		0	0	0	0	0	Ο	0	138
Tintswalo (MP)	2	0	0	0	О	0	0	0	55
Total:	6	0	0	1	0	0	0	0	1 099

CP: Gauteng; KZ: KwaZulu-Natal; NW: North West; MP: Mpumalanga; WC: Western Cape Inconclusive: insufficient viral load in sample and unable to characterise further

***influenza A subtype or B lineage results are pending

*Influenza was not detected in 12 specimens from patients who met suspected SARS-CoV-2 case definition but did not meet pneumonia (SRI) case definition. These are not included in the table.

WEEK 10 2022

NATIONAL SYNDROMIC SURVEILLANCE FOR PNEUMONIA

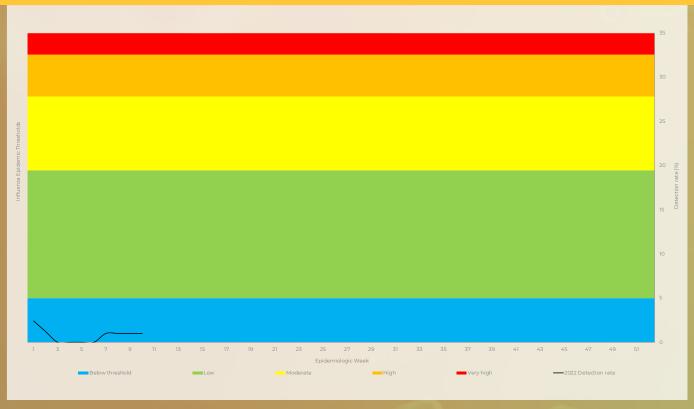


Figure 12. Influenza percentage detections and epidemic thresholds* among cases of all ages, pneumonia surveillance public hospitals, 03/01/2022 – 13/03/2022

*Thresholds based on 2010-2019 data

WEEK 10 2022

NATIONAL SYNDROMIC SURVEILLANCE FOR PNEUMONIA



Figure 13. Number of patients testing positive for respiratory syncytial virus* by province and detection rate by week, pneumonia surveillance public hospitals. 03/01/2022 – 13/03/2022

*RSV was not detected in 12 specimens from patients who met suspected SARS-CoV-2 case definition but did not meet pneumonia (SRI) case definition.



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PAGE 16

NATIONAL SYNDROMIC SURVEILLANCE FOR PNEUMONIA

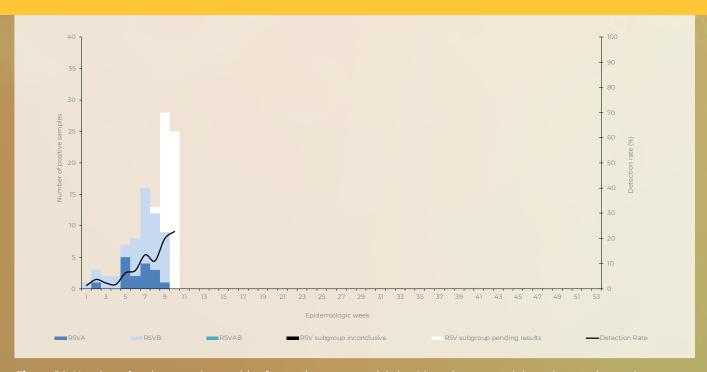


Figure 14. Number of patients testing positive for respiratory syncytial virus* by subgroup and detection rate by week,

Table 10: Number of patients positive for respiratory syncytial virus subgroups** by subgroups identified and total number of samples tested by hospital, pneumonia surveillance public hospitals, 03/01/2022 - 13/03/2022

Hospital (Province)	RSVA	RSVB	RSVAB	RSV subgroup inconclusive	RSV subgroup pending*	Total samples
Edendale (KZ)	14	0	0	0	21	216
Helen Joseph-Rahima Moosa (GP)	2	41	0	0	19	304
Klerksdorp-Tshepong (NW)	0	О	О	0		106
Mapulaneng-Matikwana (MP)	0	О	0	0	0	97
Red Cross (WC)	0	2	0	0	2	183
Mitchell's Plain (WC)	0		0	0		138
Tintswalo (MP)	0	О	0	0	1	55
Total:	16	44	0	0	45	1 099

GP: Gauteng; KZ: KwaZulu-Natal; NW: North West; MP: Mpumalanga; WC: Western Cape Inconclusive: insufficient viral load in sample and unable to characterise further RSV AB: Both RSV A and B subgroup identified *RSV results for subgroups are pending

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WEEK 10 2022

NATIONAL SYNDROMIC SURVEILLANCE FOR PNEUMONIA



Figure 15. RSV percentage detections and epidemic thresholds* among children aged < 5 years, pneumonia surveillance public hospitals, 03/01/2022 – 13/03/2022

*Thresholds based on 2010-2019 data

WEEK 10 2022

NATIONAL SYNDROMIC SURVEILLANCE FOR PNEUMONIA

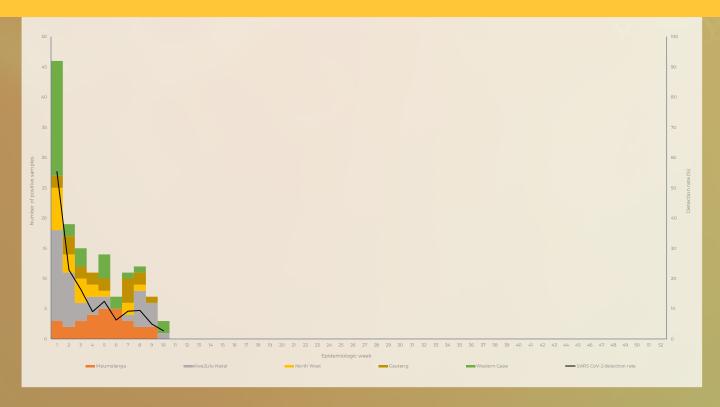


Figure 16. Number of patients testing positive for SARS-CoV- 2^* by province and detection rate by week, pneumonia surveillance public hospitals, 03/01/2022 - 13/03/2022

*Specimens from patients hospitalized with pneumonia at 6 sentinel sites in 5 provinces

Table 11. Number of patients positive for SARS-CoV- 2^* and total number of samples tested by hospital, pneumonia surveillance public hospitals, 03/01/2022 - 13/03/2022

Hospital (Province)	SARS-CoV-2 positive	Total samples tested
Edendale (KZ)	44	216
Helen Joseph-Rahima Moosa (GP)	18	304
Klerksdorp-Tshepong (NW)	20	106
Mapulaneng-Matikwana (MP)	19	97
Red Cross (WC)	13	183
Mitchell's Plain (WC)	21	138
Tintswalo (MP)	10	55
Total:	145	1 099

GP: Gauteng; KZ: KwaZulu-Natal; NW: North West; MP: Mpumalanga; WC: Western Cape
*SARS-CoV-2 was detected in 7 of 12 (58%) specimens from patients who met suspected SARS-CoV-2 case definition but did not meet pneumonia (SRI) case definition. These are not included in the table.

^{*}SARS-CoV-2 was detected in 7 of 12 (58%) specimens from patients who met suspected SARS-CoV-2 case definition but did not meet pneumonia (SRI) case definition These are not included in the epidemiological curve.

NATIONAL SYNDROMIC SURVEILLANCE FOR PNEUMONIA



Figure 17. Number and detection rate of laboratory confirmed SARS-CoV-2 cases* by variant type (variant PCR/sequencing),

Unable to assign: no lineage assigned due to poor- sequence quality OR low viral load (ct=>35) OR variant PCR could not assign variant and no sequencing result Pending: outstanding variant results

Table 12. Number of SARS-CoV-2 positive cases* by variant (variant PCR and/or sequencing) identified and total number of samples tested by hospital, pneumonia surveillance public hospitals, 03/01/2022 - 13/03/2022

Hospital (Province)	Non-Alpha/ Beta/Delta	20I (Alpha (20I)	Beta (20H)	Delta (21A, 21I, 21J)	C.1.2 (20D)	Omicron (21K, 21L, 21M)	Pending	Unable to assign	SARS- CoV-2 positive
Edendale (KZ)	0	0	0	О	0	23	12	14	49
Helen Joseph-Rahima Moosa (GP)	O		0	0	0	9	3	5	18
Klerksdorp-Tshepong (NW)	O	0	0	0	0	9	2	9	20
Mapulaneng- Matikwana (MP)	0	0	Ο	0	0	8	4	9	21
Red Cross (WC)	0	0	0	О	0	6	2	5	13
Mitchell's Plain (WC)	0	0	О	О	О	10		10	21
Tintswalo (MP)	0	0	0	О	0	5	2	3	10
Total:	0	1	0	0	0	70	26	55	152

GP: Gauteng; KZ: KwaZulu-Natal; NW: North West; MP: Mpumalanga; WC: Western Cape
*Specimens are from hospitalized patients at 7 sentinel sites in 5 provinces who met suspected SARS-CoV-2 case definition and met pneumonia (SRI) case definition

Unable to assign: no lineage assigned due to poor- sequence quality OR low viral load (ct=>35) OR variant PCR could not assign variant and no sequencing result Pending: outstanding variant results

WEEK 10 2022

SUMMARY OF LABORATORY CONFIRMED SARS-COV-2 CASES

Table 13. Characteristics of individuals with laboratory-confirmed SARS-CoV-2, enrolled in influenza-like illness

Characteristic	Influenza-like illness (ILI), public-sector, n=55 (%)	Pneumonia, n=152 (%)		
Age group (years)				
0-9	8/55 (15)	35/152 (23)		
10-19	9/55 (16)	5/152 (3)		
20-39	12/55 (22)	43/152 (28)		
40-59	20/55 (36)	40/152 (26)		
60-79	5/55 (9)	23/152 (15)		
≥80	1/55 (2)	6/152 (4)		
Sex-female	33/55 (60)	82/152 (54)		
Province*				
Gauteng	N/A	18/152 (12)		
KwaZulu-Natal	7/55 (13)	49/152 (32)		
Mpumalanga	21/55 (38)	31/152 (20)		
North West	20/55 (36)	20/152 (13)		
Western Cape	7/55 (13)	34/152 (22)		
Race				
Black	38/55 (69)	123/149 (83)		
Coloured	7/55 (13)	24/149 (16)		
Asian/Indian	0/55 (0)	0/149 (0)		
White	9/55 (16)	0/149 (0)		
Other	1/55 (2)	2/149 (1)		
Variant				
Non-Alpha/Beta/Delta	0/55 (0)	0/152 (0)		
Alpha(201)	0/55 (0)	1/152 (1)		
Beta(20H)	0/55 (0)	0/152 (0)		
Delta(21A, 21I, 21J)	0/55 (0)	0/152 (0)		
C.1.2(20D)	0/55 (0)	0/152 (0)		
Omicron(21K,21L,21M)	14/55 (25)	70/152 (46)		
Pending results ^{\$}	10/55 (18)	26/152 (17)		
Unable to assign ^{\$\$}	31/55 (56)	55/152 (36)		

WEEK 10 2022

Characteristic	Influenza-like illness (ILI), public-sector, n=55 (%)	Pneumonia, n=152 (%)		
Presentation				
Fever	38/55 (69)	59/149 (40)		
Cough	54/55 (98)	133/149 (89)		
Shortness of breath	24/55 (44)	87/149 (58)		
Chest pain	23/55 (42)	62/149 (42)		
Diarrhoea	6/55 (11)	17/149 (11)		
Underlying conditions				
Hypertension	11/55 (20)	25/149 (17)		
Cardiac	1/55 (2)	1/149 (1)		
Lung disease	0/55 (0)	1/149 (1)		
Diabetes	2/55 (4)	15/149 (10)		
Cancer	0/55 (0)	3/149 (2)		
Tuberculosis	0/55 (0)	15/149 (10)		
HIV-infection	8/55 (17)	57/149 (39)		
Other **	1/55 (2)	2/149 (1)		
SARS-CoV-2 Vaccine				
Pfizer-BioNTech (1st dose)	8/53 (15)	15/145 (10)		
Pfizer-BioNTech (2nd dose)	8/53 (15)	10/145 (7)		
Johnson & Johnson	8/53 (15)	14/145 (10)		
Booster	0/53 (0)	0/145 (0)		
Management				
Oxygen therapy	0/55 (0)	72/149 (48)		
ICU admission	N/A	0/149 (0)		
Ventilation	N/A	2/149 (1)		
Outcome***				
Died	0/55 (0)	9/137 (7)		

WEEK 10 2022

METHODS

SARS-CoV-2 Testing

March 2020 - March 2021: SARS-CoV-2 was detected using the Roche E gene real-time PCR assay (Corman et al. Euro Surveillance 2020) with cycle threshold (Ct) <40 interpreted as positive for SARS-CoV-2. From April 2021 to date the laboratory changed to the Allplex™ SARS-CoV-2/FluA/FluB/RSV kit (Seegene Inc., Seoul, South Korea), with positivity assigned if the PCR cycle threshold (Ct) was <40 for ≥1 gene targets (N, S or RdRp).

A confirmed SARS-CoV-2 case is a person of any age enrolled in surveillance with laboratory confirmation of SARS-CoV-2 infection by PCR. Only positive SARS-CoV-2 specimens on PCR are further tested to determine variant/lineage type by variant PCR or genomic sequencing.

Variant PCR

Allplex™ SARS-CoV-2 Variants I PCR detects Alpha and Beta/Gamma variants. The assay was conducted on all SARS-CoV-2-positive samples from 1 March 2020 – 30 June 2021.

Allplex™ SARS-CoV-2 Variants II PCR detects Delta variant and distinguishes Beta from Gamma. The assay was conducted on SARS-CoV-2-positive samples from 1 Jan to 30 June 2021.

Extraction: Total nucleic acids were extracted from 200µl NP/OP samples in universal or viral transport medium using a MagNA Pure 96 automated extractor and DNA/Viral NA Small Volume v2.0 extraction kit (Roche Diagnostics, Mannheim, Germany).

SARS-CoV-2 genomic surveillance

SARS-CoV-2 Whole-Genome Sequencing and Genome Assembly

RNA Extraction

RNA was extracted either manually or automatically in batches, using the QIAamp viral RNA mini kit (QIAGEN, CA, USA) or the Chemagic 360 using the CMG-1049 kit (PerkinElmer, MA, USA). A modification was done on the manual extractions by adding 280 µl per sample, in order to increase yields. 300 µl of each sample was used for automated magnetic bead-based extraction using the Chemagic 360. RNA was eluted in 60 µl of the elution buffer. Isolated RNA was stored at -80 °C prior to use

PCR and Library Preparation

Sequencing was performed using the Illumina COVIDSeq protocol (Illumina Inc., CA, USA) or nCoV-2019 ARTIC network sequencing protocol v3 (https://artic.network/ncov-2019). These are amplicon-based next-generation sequencing approaches. Briefly, for the nCoV-2019 ARTIC network sequencing protocol, the first strand synthesis was carried out on extracted RNA samples using random hexamer primers from the SuperScript IV reverse transcriptase synthesis kit (Life Technologies, CA, USA) or LunaScript RT SuperMix Kit (New England Biolabs (NEB), MA, USA). The synthesized cDNA was amplified using multiplex polymerase chain reactions (PCRs) using ARTIC nCoV-2019 v3 primers. For the COVIDSeq

protocol, the first strand synthesis was carried out using random hexamer primers from Illumina and the synthesized cDNA underwent two separate multiplex PCR reactions.

For Illumina sequencing using the nCoV-2019 ARTIC network sequencing protocol, the pooled PCR products underwent bead-based tagmentation using the Nextera Flex DNA library preparation kit (Illumina Inc., CA, USA). The adapter-tagged amplicons were cleaned up using AmpureXP purification beads (Beckman Coulter, High Wycombe, UK) and amplified using one round of PCR. The PCRs were indexed using the Nextera CD indexes (Illumina Inc., CA, USA) according to the manufacturer's instructions. For COVIDSeq sequencing protocol, pooled PCR amplified products were processed for tagmentation and adapter ligation using IDT for Illumina Nextera UD Indexes. Further enrichment and cleanup was performed as per protocols provided by the manufacturer (Illumina Inc., CA, USA). Pooled samples from both COVIDSeq protocol and nCoV-2019 ARTIC network protocol were quantified using Qubit 3.0 or 4.0 fluorometer (Invitrogen Inc., MA, USA) using the Qubit dsDNA High Sensitivity assay according to manufacturer's instructions. The fragment sizes were analyzed using TapeStation 4200 (Invitrogen Inc., MA, USA). The pooled libraries were further normalized to 4nM concentration and 25 µl of each normalized pool containing unique index adapter sets were combined in a new tube. The final library pool was denatured and neutralized with 0.2 N sodium hydroxide and 200 mM Tris-HCL (pH7), respectively. 1.5 pM sample library was spiked with 2% PhiX. Libraries were loaded onto a 300-cycle NextSeq 500/550 HighOutput Kit v2 and run on the Illumina NextSeq 550 instrument (Illumina Inc., CA, USA).

Assembly, Processing and Quality Control of Genomic Sequences

Raw reads from Illumina sequencing were assembled using the Exatype NGS SARS-CoV-2 pipeline v1.6.1, (https://sarscov-2.exatype.com/). The resulting consensus sequence was further manually polished by considering and correcting indels in homopolymer regions that break the open reading frame (probably sequencing errors) using Aliview v1.27, (http://ormbunkar.se/aliview/) (Larsson, 2014). Mutations resulting in mid-gene stop codons and frameshifts were reverted to wild type. All assemblies determined to have acceptable quality (defined as having at least 1 000 000 reads and at least 40 % 10 X coverage) were deposited on GISAID (https://www.gisaid.org/) (Elbe & Buckland-Merrett, 2017; Shu & McCauley, 2017).

Classification of Lineage, Clade and Associated Mutations

Assembled genomes were assigned lineages using the 'Phylogenetic Assignment of Named Global Outbreak Lineages' (PANGOLIN) software suite (https://github.com/hCoV-2019/pangolin) (Rambaut et al., 2020), a tool used for dynamic SARS-CoV-2 lineage classification. The SARS-CoV-2 genomes in our dataset were also classified using the clade classification proposed by NextStrain (https://nextstrain.org/), a tool built for real-time tracking of the pathogen evolution (Hadfield et al., 2018).