

# Australian Respiratory Surveillance Report

# Key messages

This report presents a national epidemiological update for acute respiratory infections, including coronavirus disease 2019 (COVID-19), influenza and respiratory syncytial virus (RSV), with a focus on the current reporting period (16 June to 29 June 2025) and earlier severity reporting periods (up to 15 June 2025).

**In the community:** Respiratory illness activity (self-reported new fever and cough symptoms) remains lower than the trends observed at the same time in previous years. While slightly more people reported new fever and cough symptoms compared to the previous fortnight, fewer people reported taking time off work due to respiratory illness (self-reported new fever and cough symptoms). The number of COVID-19 cases was moderate in the last fortnight with cases increasing in most jurisdictions but remaining lower than at the same time in previous years. The number of influenza cases was moderate in the last fortnight and while case numbers are above the five-year average, cases remain lower than at the same time last year. The number of RSV cases in the last fortnight was high; however, trends varied across jurisdictions. In most jurisdictions, expected seasonal increases in RSV cases were observed in the last fortnight.

**In general practice:** There were slightly more influenza-like-illness (new fever and cough symptoms) consultations at sentinel surveillance sites in the last fortnight compared with the previous fortnight. Influenza-like-illness rates in the last fortnight were slightly higher than the rates observed at the same time in 2023 and the five-year average but remain lower than rates observed in 2022 and 2024.

**In hospitals:** Sentinel hospital-based surveillance shows the number of patients admitted with severe acute respiratory infections this year has continued to increase since late March. The length of hospital stay continues to vary only slightly between illnesses and the proportion of those patients who were admitted directly to intensive care at a sentinel hospital site has remained low. At sentinel hospitals, more children (those aged 16 years and younger) were admitted with RSV than with influenza or COVID-19, while more adults were admitted with COVID-19 compared to influenza or RSV. Sentinel intensive care surveillance shows the overall number of patients with severe acute respiratory infections has remained low and stable this year to date, though now appears to be increasing. In 2025 to date, most patients were admitted to sentinel intensive care with rhinovirus / enterovirus, followed by influenza. The duration of intensive care stay varies slightly between illnesses. In the last fortnight, the average number of COVID-19 cases in intensive care decreased, while the average number of intensive care staff unavailable due to COVID-19 illness or exposure increased.

**Deaths:** COVID-19 has been the leading cause of acute respiratory infection mortality across 2023–2025. All three acute respiratory infections (COVID-19, influenza and RSV) are more likely to cause death in older age groups than younger age groups.

**In laboratories:** Test positivity for SARS-CoV-2, influenza and RSV increased in the last fortnight. One of the most recently designated variants under monitoring, NB.1.8.1, is the dominant SARS-CoV-2 variant in the last 28 days in Australia accounting for 48.6% of sequences. Small numbers of sequences of other variants under monitoring, including LP.8.1, LB.1, KP.3.1.1, XEC and XFG continue to be observed in Australia.

**Vaccine coverage, effectiveness and match:** Nationally, fewer adults have received a COVID-19 vaccine in the past 12 months compared to the 12 months prior. Influenza vaccine coverage this year to date is increasing but remains lower than the same time in previous years. For RSV, nirsevimab uptake is increasing; however, there is substantial variation in nirsevimab uptake in infants across jurisdictions. Of influenza isolates characterised in 2025 thus far, over 99% have been a good match to the corresponding 2025 vaccine components.

# Australian Respiratory Surveillance Report

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This report was prepared by Lauren Kutzner, Lauren Welsh, Ash Donovan, and Siobhan St George on behalf of the interim Australian Centre for Disease Control. We thank the staff and participants from the surveillance systems who contribute data for acute respiratory illness surveillance across Australia.

The report presents a national overview of acute respiratory infections in Australia, drawing information from several different surveillance systems. These surveillance systems help us to understand the distribution of acute respiratory illnesses in the community, the severity of infections including which populations might be at risk, and the impact of acute respiratory illnesses on the community and health system in Australia.

Surveillance indicators presented in this report are based on the [Australian National Surveillance Plan for COVID-19, Influenza, and RSV](#). Please refer to the [Technical Supplement – Australian Respiratory Surveillance Report](#) for information on our surveillance sources and data considerations, including the considerable impact of the COVID-19 pandemic on acute respiratory infection surveillance in Australia. A summary of data considerations for this report are provided below:

- Due to the dynamic nature of the surveillance systems used in this report, surveillance data are considered preliminary and subject to change as updates are received, with the most recent weeks considered particularly incomplete. Data in this report may vary from data reported in other national reports and reports by states and territories.
- Data in this report are presented by date of event (diagnosis, admission or death) or by the International Organization for Standardization (ISO) week date system, with weeks defined as seven-day periods which begin on a Monday and end on a Sunday. The ISO week date system is used to support trends comparisons over time more effectively. The current reporting period includes 16 June to 29 June 2025 and where comparisons to the previous fortnight are made this includes 2 June to 15 June 2025.
- In Australia, states and territories (the Australian Capital Territory [ACT], New South Wales [NSW], the Northern Territory [NT], Queensland [Qld], South Australia [SA], Tasmania [Tas], Victoria [Vic], and Western Australia [WA]) report notified cases to the **National Notifiable Diseases Surveillance System (NNDSS)** based on the [Australian national surveillance case definitions](#). NNDSS data are analysed and reported based on diagnosis date, which is the true onset date of a case if known, otherwise it is the earliest of the specimen date, the notification date, or the notification received date. The NNDSS data for this report were extracted on 2 July 2025.
- To account for the lag in collection and provision of severity data from some surveillance systems, and for the time delay between illness onset and the development of severe disease outcomes, cases with an admission date or a diagnosis date in the last two weeks are excluded from severity analyses for hospitalisations and intensive care admissions. As such, the severity reporting periods are two weeks behind the end of the current reporting period. For this report, severity reporting includes data from 2 June to 15 June 2025 unless specified otherwise. Where comparisons to the previous severity fortnight are made this includes 19 May to 1 June 2025.
- Death registrations from the Australian Bureau of Statistics (ABS) Provisional Mortality Statistics are now used as the primary data source for measuring acute respiratory infection associated deaths. The ABS mortality data is sourced from the Registry of Births, Deaths and Marriages and is separate from the NNDSS. Registration-based mortality data needs time to be received and processed. For this reason, mortality statistics in this report may lag by at least two months.
- Analysis and reporting outputs were produced using R Statistical Software v4.3.1. While every care has been taken in preparing this report, the Australian Government Department of Health, Disability and Ageing does not accept liability for any injury or loss or damage arising from the use of, or reliance upon, the content of the report or Technical Supplement. For further information about this report refer to the [Technical Supplement – Australian Respiratory Surveillance Report](#) or contact [respiratory.surveillance@health.gov.au](mailto:respiratory.surveillance@health.gov.au).

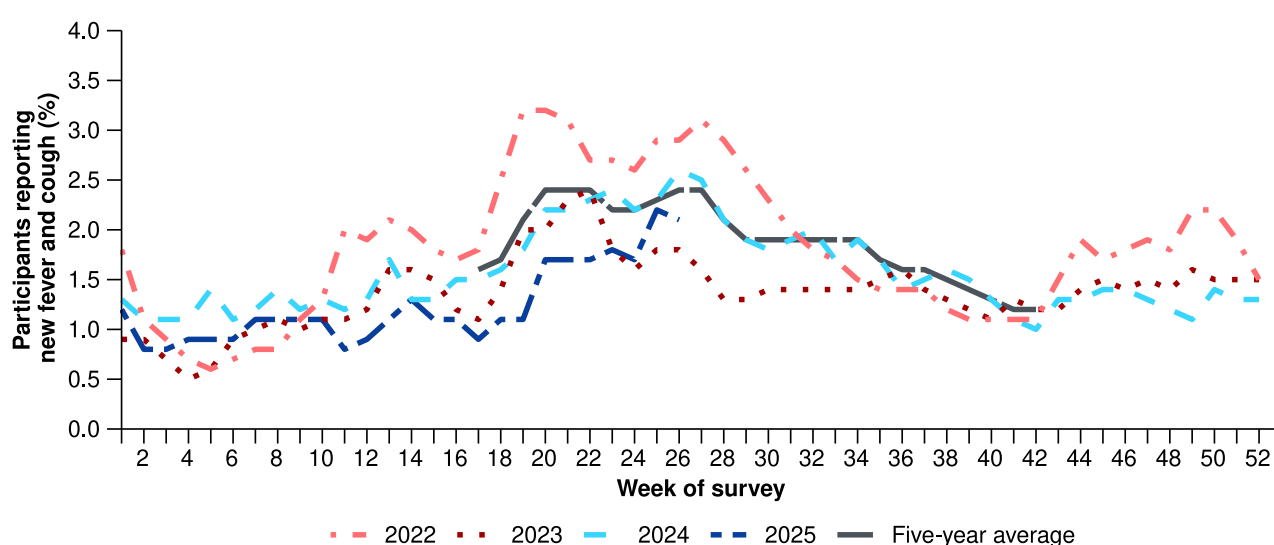
# Community surveillance

Community surveillance monitors respiratory illnesses in the community, providing information on the number of people reporting respiratory symptoms, testing practices, and the impact of respiratory illnesses.

Community surveillance includes notification data obtained from laboratory tests for infections. Infections that are diagnosed and notified are only a subset of the total number of infections occurring in the community.

- Community surveys via FluTracking indicate current respiratory illness symptoms and test positivity remain lower than the five-year average.
- In the last fortnight (16 June to 29 June 2025), slightly more survey participants reported new fever and cough symptoms (2.2%), than in the previous fortnight (1.8%) (Figure 1).
- In the last fortnight, more survey participants with new fever and cough symptoms used a rapid antigen test (RAT) (56.1%; 728/1,298) than a polymerase chain reaction (PCR) test (9.2%; 120/1,298) to test for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2).
- Self-reported SARS-CoV-2 RAT positivity was slightly higher in the last fortnight (31.5%; 229/728) than in the previous fortnight (30.9%; 226/732). Self-reported SARS-CoV-2 PCR positivity was also higher in the last fortnight (16.7%; 20/120) than in the previous fortnight (11.6%; 17/147).
- In the last fortnight, 8.6% (112/1,298) of survey participants with new fever and cough symptoms used a PCR test to test for influenza. Self-reported influenza PCR positivity was slightly higher this fortnight (23.2%; 26/112), than in the previous fortnight (23.0%; 31/135).
- In the last fortnight, less survey participants reported taking three or more days off work or normal duties due to fever and cough symptoms (47.6%; 618/1,298), than in the previous fortnight (50.7%; 659/1,300).
- From January to early March 2025, the weekly proportion of survey participants with new fever and cough symptoms was relatively consistent with the proportions observed at the same time in 2022–2024. Since mid-March, the weekly proportion has been lower than observed at the same time in 2022–2024, and from late April has been lower than the five-year average (Figure 1).

**Figure 1: Age standardised proportion of survey participants reporting new fever and cough symptoms compared with the five-year average\* by year and week of survey, Australia, 2022 to 29 June 2025**



Source: FluTracking

\* From 2020, FluTracking expanded their data capture period to year-round. Data before May and after October for any year before 2020 are not available for historical comparisons. The years 2020 and 2021 are excluded when comparing the current season to historical periods when influenza virus has circulated without public health restrictions. As such, the five-year average includes the years 2018 to 2019 and 2022 to 2024. Please refer to the [Technical Supplement](#) for interpretation of the five-year average.

- In the last fortnight (16 June to 29 June 2025), there was a 12.7% increase in COVID-19 cases, a 55.4% increase in influenza cases, and a 7.3% increase in RSV cases.

**Table 1: Notified cases and notification rate per 100,000 population by disease, five-year age group, and jurisdiction\*†, Australia, 1 January to 29 June 2025**

	COVID-19			Influenza			RSV		
	Reporting period (n)	Year to date (n)	Year to date (rate)	Reporting period (n)	Year to date (n)	Year to date (rate)	Reporting period (n)	Year to date (n)	Year to date (rate)
<b>Age group (years)</b>									
0–4	1,305	10,091	669	3,933	17,363	1,151	5,773	43,862	2,907
5–9	448	2,727	169	5,713	21,837	1,356	1,262	6,203	385
10–14	483	3,126	187	3,957	14,736	880	502	2,561	153
15–19	475	3,635	219	2,143	8,844	532	302	1,752	105
20–24	508	3,611	202	846	4,862	272	191	1,427	80
25–29	663	4,549	228	898	4,915	246	285	1,759	88
30–34	800	5,471	268	1,395	6,905	339	368	2,247	110
35–39	912	6,267	316	1,956	9,280	468	330	2,270	114
40–44	899	5,986	323	2,061	9,714	525	330	1,912	103
45–49	809	5,223	321	1,567	7,625	468	307	1,786	110
50–54	743	5,321	315	1,337	7,178	425	345	2,188	129
55–59	765	5,149	336	1,228	6,558	428	348	2,340	153
60–64	831	5,522	360	1,136	6,445	420	411	2,647	173
65–69	840	5,686	418	967	5,725	421	420	2,774	204
70+	5,645	35,245	1,055	3,342	18,250	546	1,816	11,344	340
<b>Jurisdiction</b>									
ACT	314	1,698	358	937	2,881	608	351	1,266	267
NSW	7,802	49,083	579	11,997	55,281	652	5,390	45,404	535
NT	60	817	320	124	1,923	754	32	524	205
Qld	2,627	21,318	382	5,437	29,548	529	1,806	17,305	310
SA	889	6,789	361	3,015	10,790	575	692	3,026	161
Tas	197	1,423	247	430	2,201	383	117	718	125
Vic	3,116	19,362	277	8,798	37,125	532	3,941	15,973	229
WA	1,132	7,178	242	1,747	10,522	355	661	2,866	97
<b>Total</b>	<b>16,137</b>	<b>107,668</b>	<b>396</b>	<b>32,485</b>	<b>150,271</b>	<b>552</b>	<b>12,990</b>	<b>87,082</b>	<b>320</b>

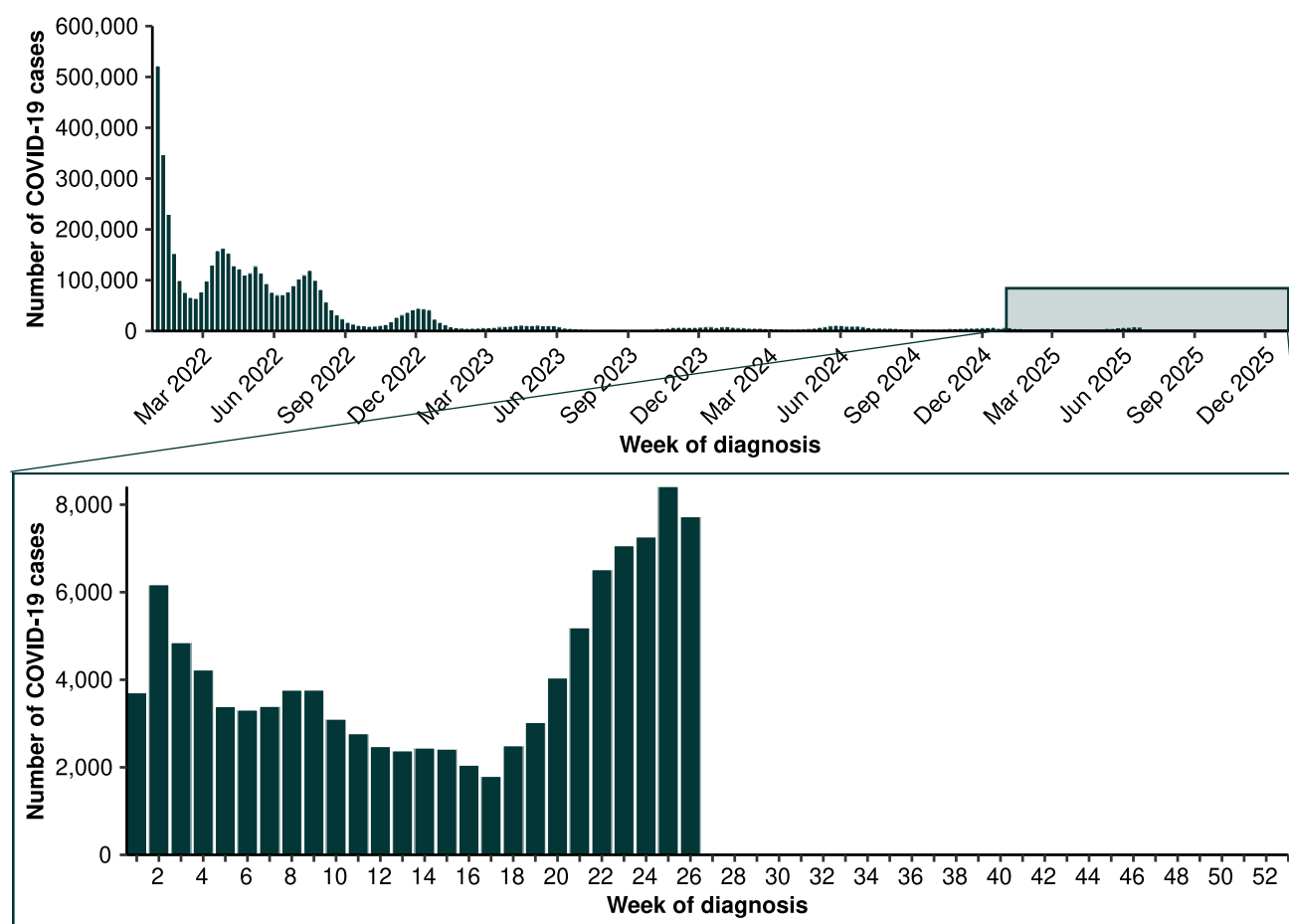
Source: National Notifiable Diseases Surveillance System (NNDSS)

\* Rate per 100,000 population for the given time period. Population data are based on the Australian Bureau of Statistics (ABS) [Estimated Resident Population \(ERP\) for the reference period June 2024, released 12 December 2024](#).

† Total includes cases with missing age.

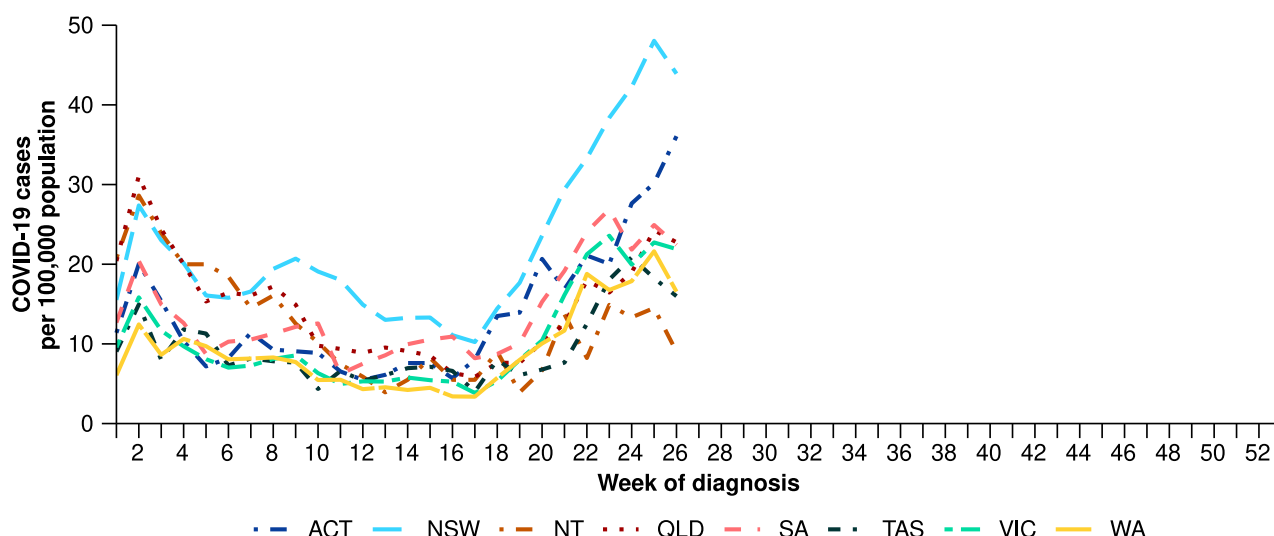
- In the last fortnight, the number of COVID-19 cases was moderate and increasing; however, case notifications remain lower than the number of cases reported at the same time last year (16,137 in the last fortnight, compared to 19,860 COVID-19 cases in the fortnight ending 30 June 2024).
- Following a decreasing trend from mid-January to late April 2025, COVID-19 cases have increased week-on-week from late April 2025; however, the overall number of cases this year to date (n=107,668) is 35.9% less than the number of cases observed in the same time period last year (n=167,984) (Figure 2).
- The current increase in COVID-19 case numbers is not unexpected based on previous transmission patterns, with COVID-19 case numbers peaking in May 2023 and in June 2024 (Figure 2). The current increase in COVID-19 cases is likely driven by the new SARS-CoV-2 variant NB.1.8.1.
- In the last fortnight, trends in COVID-19 notification rates have varied across jurisdictions (Figure 3). COVID-19 notification rates in the ACT increased steeply this fortnight, with slower increases in Qld and Vic. COVID-19 notification rates in NSW, NT, SA, Tas and WA decreased slightly.
- In the year to date, COVID-19 notification rates remain highest in people aged 70 years or over, likely due to higher case ascertainment from targeted testing strategies for populations at-risk of severe disease or who live in a high-risk setting such as a residential aged care home (Table 1).
- In the year to date, COVID-19 notification rates are highest in NSW and lowest in WA (Table 1).

**Figure 2: Notified COVID-19 cases (laboratory-confirmed only) by year and week of diagnosis, Australia, 2022 to 29 June 2025**



Source: National Notifiable Diseases Surveillance System (NNDSS)

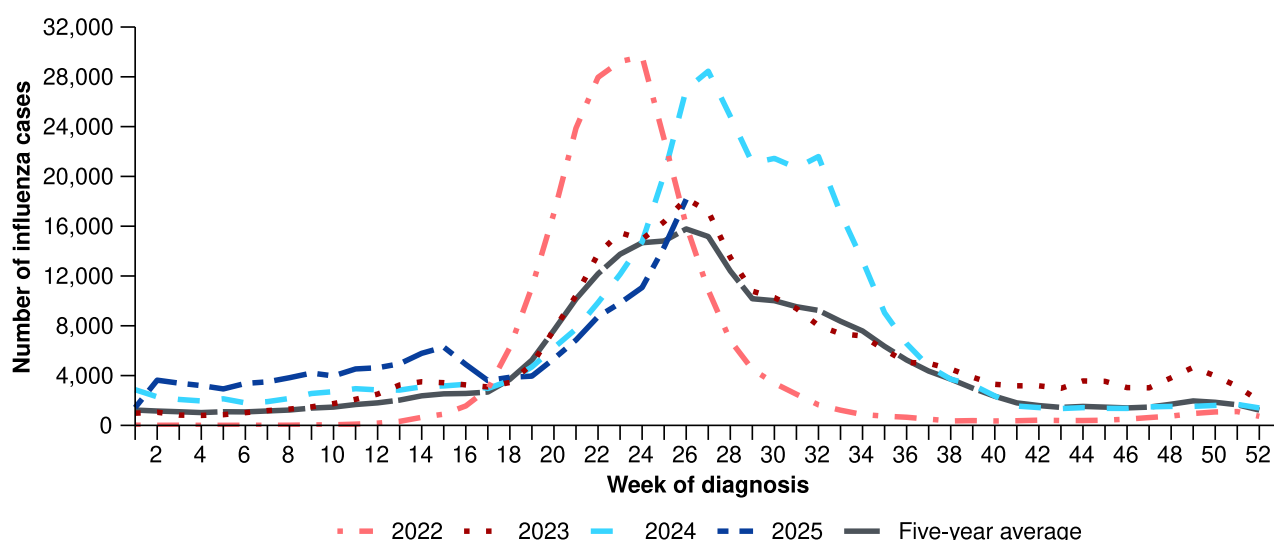
**Figure 3: Notification rates\* per 100,000 population for COVID-19 cases by state or territory and week of diagnosis, Australia, 1 January to 29 June 2025**



Source: National Notifiable Diseases Surveillance System (NNDSS)

\* Rate per 100,000 population for the given time period. Population data are based on the Australian Bureau of Statistics (ABS) [Estimated Resident Population \(ERP\)](#) for the reference period June 2024, released 12 December 2024

**Figure 4: Notified influenza cases and five-year average\* by year and week of diagnosis, Australia, 2022 to 29 June 2025**



Source: National Notifiable Diseases Surveillance System (NNDSS)

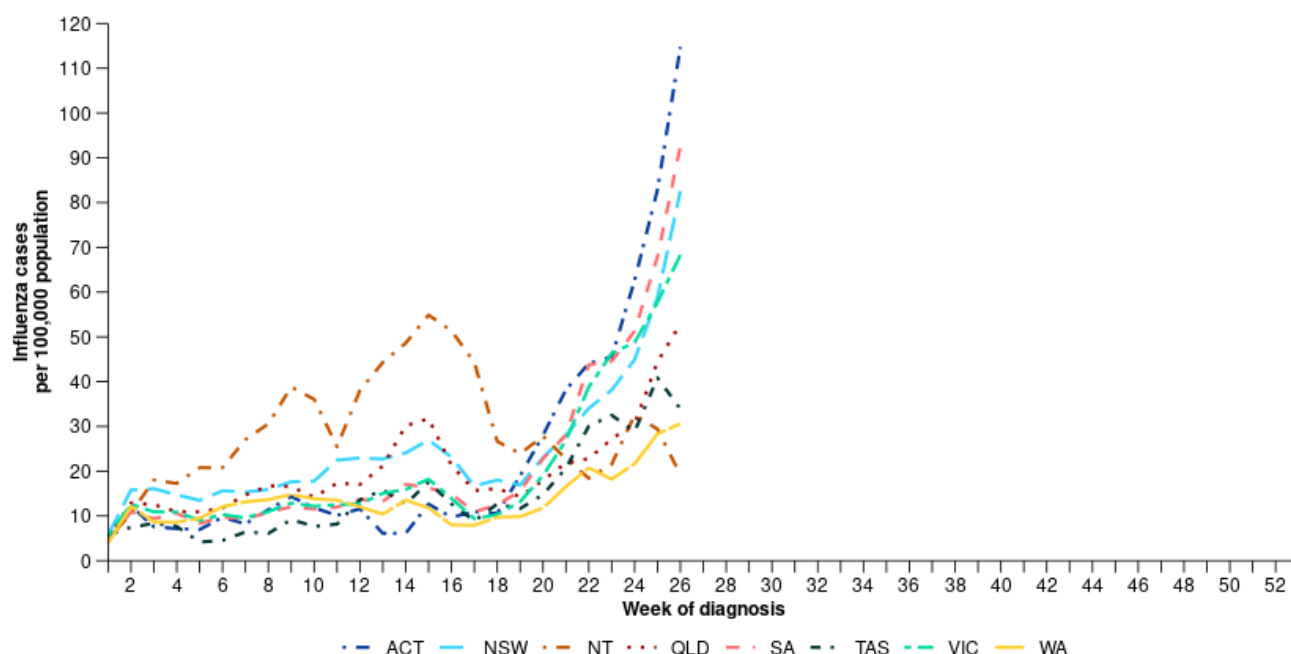
\* The years 2020 and 2021 are excluded when comparing the current season to historical periods when influenza virus has circulated without public health restrictions. As such, the five-year average includes the years 2018 to 2019 and 2022 to 2024. Please refer to the [Technical Supplement](#) for interpretation of the five-year average.

- In the last fortnight, the number of influenza cases was moderate and increased above the five-year average.
- Following higher than previously observed influenza case numbers in January to mid-April 2025, influenza cases remained relatively stable throughout April. From early May, influenza case numbers increased in line with trends observed in 2023–2024 and the five-year average. In the last fortnight, influenza case numbers have increased above the five-year average; however, remain lower than the same time period last year. The current increase in influenza cases case numbers is not unexpected based on previous transmission patterns, with influenza case numbers in 2024 peaking in July.
- The overall number of influenza cases this year to date (n=150,271) is 1.9% more than the number of cases observed in the same time period last year (n=147,435) (Figure 4).



- In the last fortnight, most jurisdictions have observed a steeply increasing trend in influenza notification rates, although the rate of increase was slower in Tas and WA. A decreasing trend in influenza notification rates was observed in the NT (Figure 5).
- In the year to date, influenza notification rates remain highest in children aged 5–9 years and children aged 0–4 years (Table 1).
- In the year to date, influenza notification rates are highest in the NT and lowest in WA (Table 1).

**Figure 5: Notification rates\* per 100,000 population for influenza cases by state or territory and week of diagnosis, Australia, 1 January to 29 June 2025**



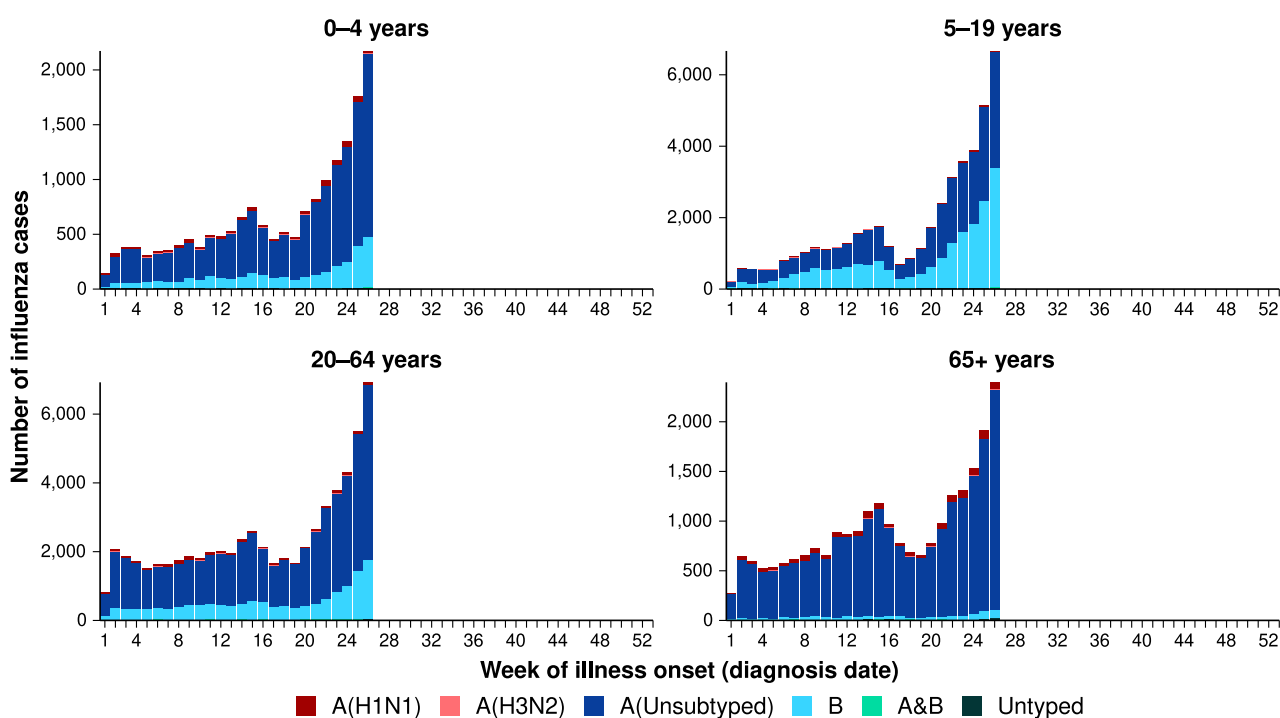
Source: National Notifiable Diseases Surveillance System (NNDSS)

\* Rate per 100,000 population for the given time period. Population data are based on the Australian Bureau of Statistics (ABS) Estimated Resident Population (ERP) for the reference period June 2024, released 12 December 2024.

- In the last fortnight, most influenza notifications were influenza A(Unsubtyped) (67.3%; 21,869/32,485), followed by influenza B (30.5%; 9,920/32,485), then influenza A(H1N1) (1.6%; 507/32,485), and influenza untyped (0.4%; 140/32,485). In the last fortnight, there have been 35 influenza A&B co-detections (Figure 6).
- In the year to date, influenza A (Unsubtyped) has accounted for most cases across all age groups, followed by influenza B. The proportion of influenza B cases is the highest in the 5–19 years age group. There has been a small number of influenza A(H1N1) and influenza A(H3N2) cases across all age groups (Figure 6).
  - There is a comparatively higher proportion of influenza B cases this season than observed in 2024. While influenza B is often a good match with the seasonal influenza vaccine strain, influenza B can result in more severe infections in children.
- In the year to date, influenza A(Unsubtyped) has accounted for the majority of influenza cases across all jurisdictions. Several jurisdictions have been experiencing increasing numbers of influenza B cases in the year to date; however, the proportion of influenza B and influenza A varies week-on-week (Figure 7).
- Influenza A(H1N1) and influenza A(H3N2) cases were most commonly observed in the NT, Qld, Tas and WA (Figure 7); however, trends in influenza subtypes should be interpreted with care as there are jurisdictional differences in the number and selection of influenza samples that undergo typing.



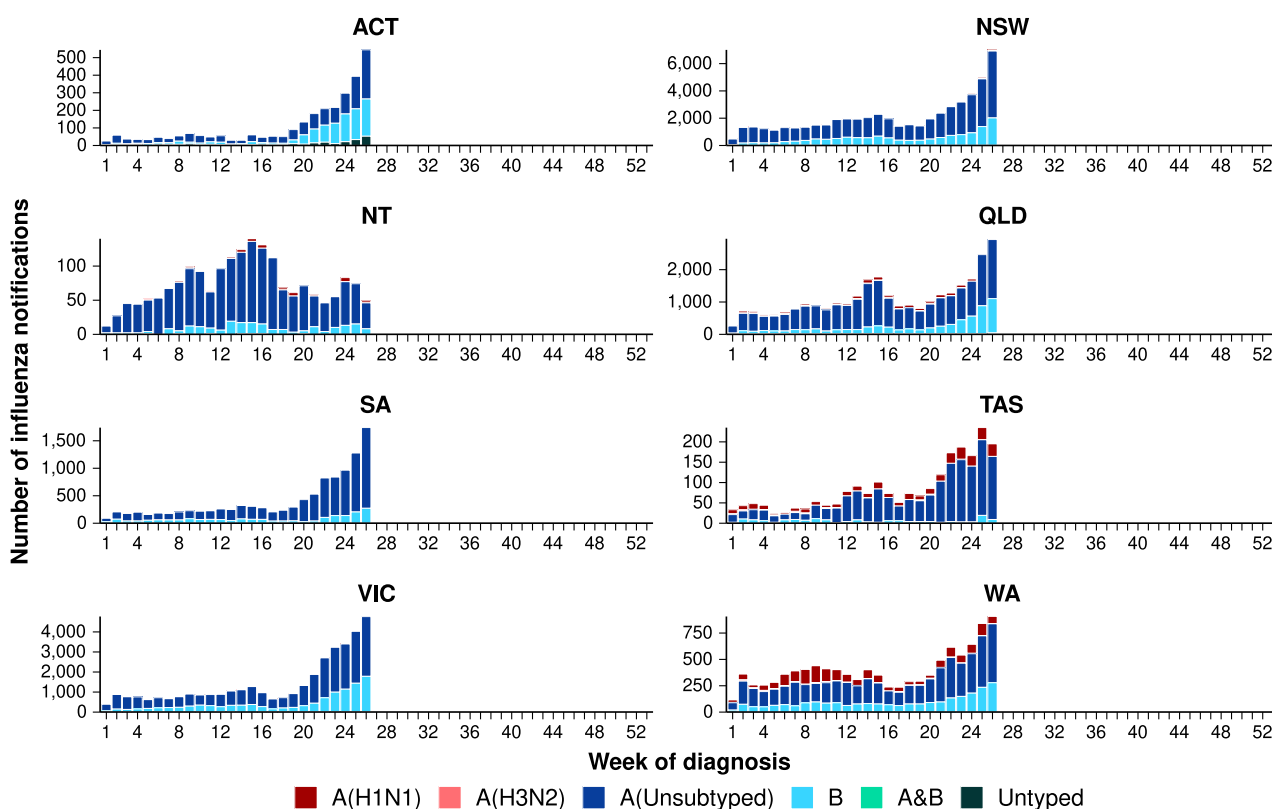
**Figure 6: Notified influenza cases by influenza subtype, age group\*, and week of diagnosis, Australia, 1 January to 29 June 2025**



Source: National Notifiable Diseases Surveillance System (NNDSS)

\* Axis varies between age groups.

**Figure 7: Notified influenza cases by influenza subtype, jurisdiction\*, and week of diagnosis, Australia, 1 January to 29 June 2025**

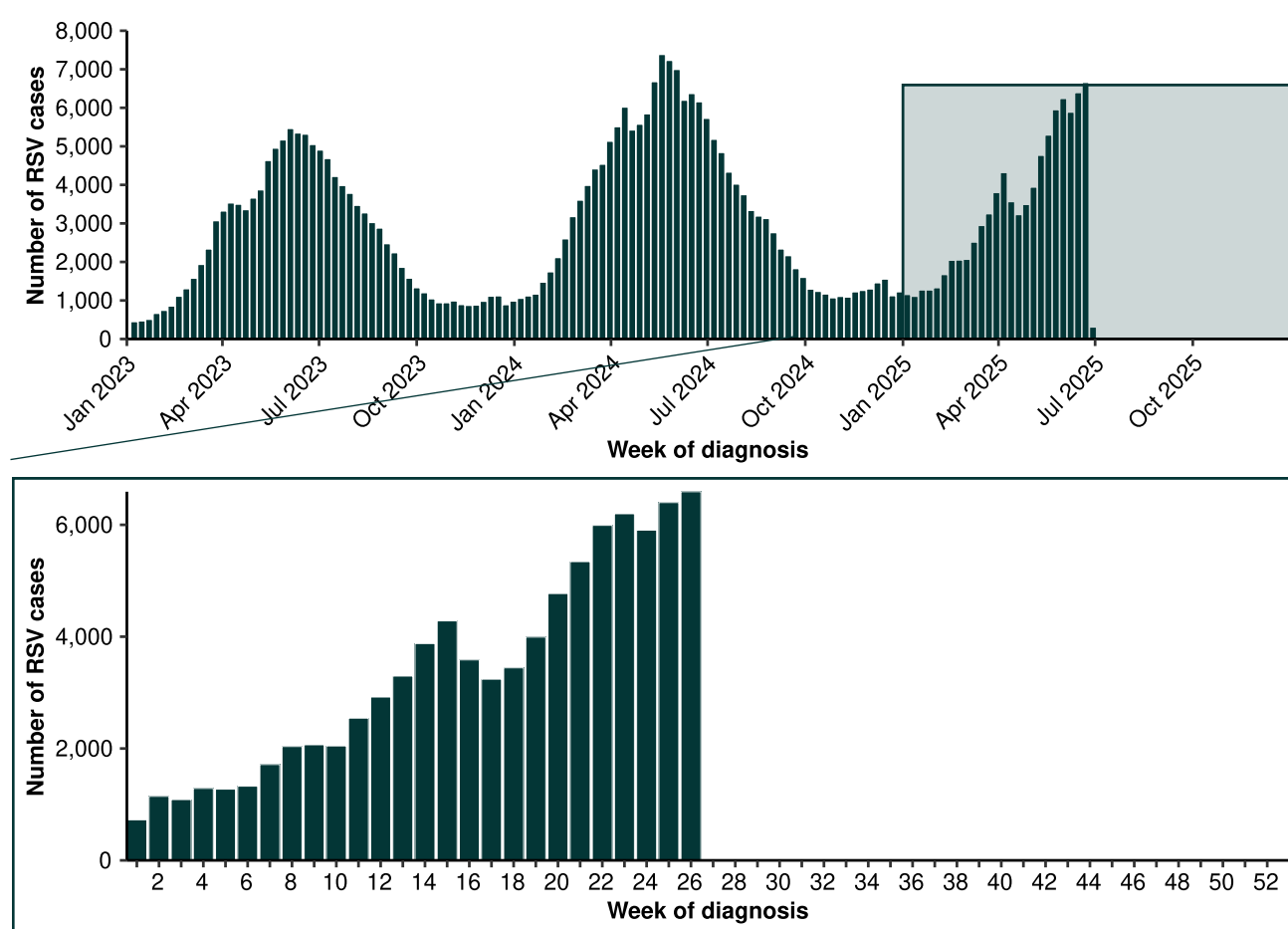


Source: National Notifiable Diseases Surveillance System (NNDSS)

\* Axis varies between jurisdictions.

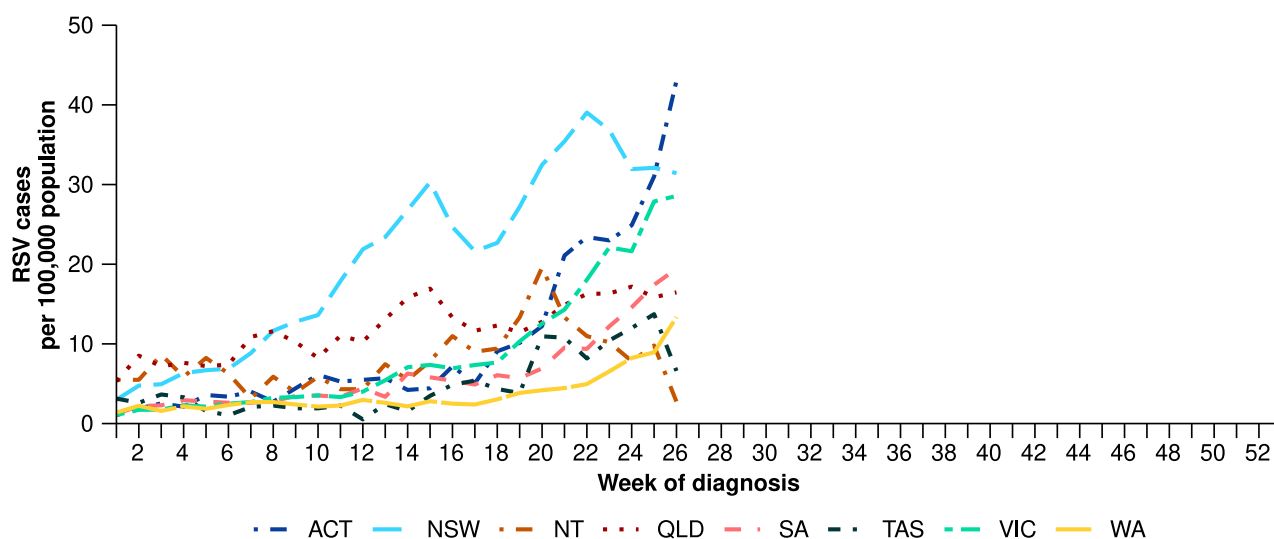
- In the last fortnight, the number of RSV cases was high and slightly above the number of cases reported at the same time last year (12,990 in the last fortnight, compared to 12,107 in the fortnight ending 30 June 2024).
- The number of RSV cases has been steadily increasing since the start of 2025, with slight week-on-week decreases observed across mid- to late April. Overall, the trend in case numbers this year to date has been similar to trends observed in previous years; however, the number of cases this year to date (n=87,082) is 22.5% less than the number of cases observed in the same time period last year (n=112,328) (Figure 8).
- In the last fortnight, notification rates increased steeply in the ACT with slower increases observed in Qld, SA, Vic and WA. (Figure 9). RSV notification rates decreased slightly in NSW and the NT.
- In the year to date, RSV notification rates remain considerably higher in children aged 0–4 years than in other age groups (Table 1).
- In the year to date, RSV notification rates are highest in NSW and lowest in WA (Table 1).

**Figure 8: Notified RSV cases by year and week of diagnosis\*, Australia, 2023 to 29 June 2025**



Source: National Notifiable Diseases Surveillance System (NNDSS). Please note, RSV became notifiable in all states and territories on 1 September 2022 and comprehensive national notification data became available after this point. For this reason, RSV notification trends are only presented from 1 January 2023.

**Figure 9: Notification rates\* per 100,000 population for RSV cases by state or territory and week of diagnosis, Australia, 1 January to 29 June 2025**



Source: National Notifiable Diseases Surveillance System (NNDSS)

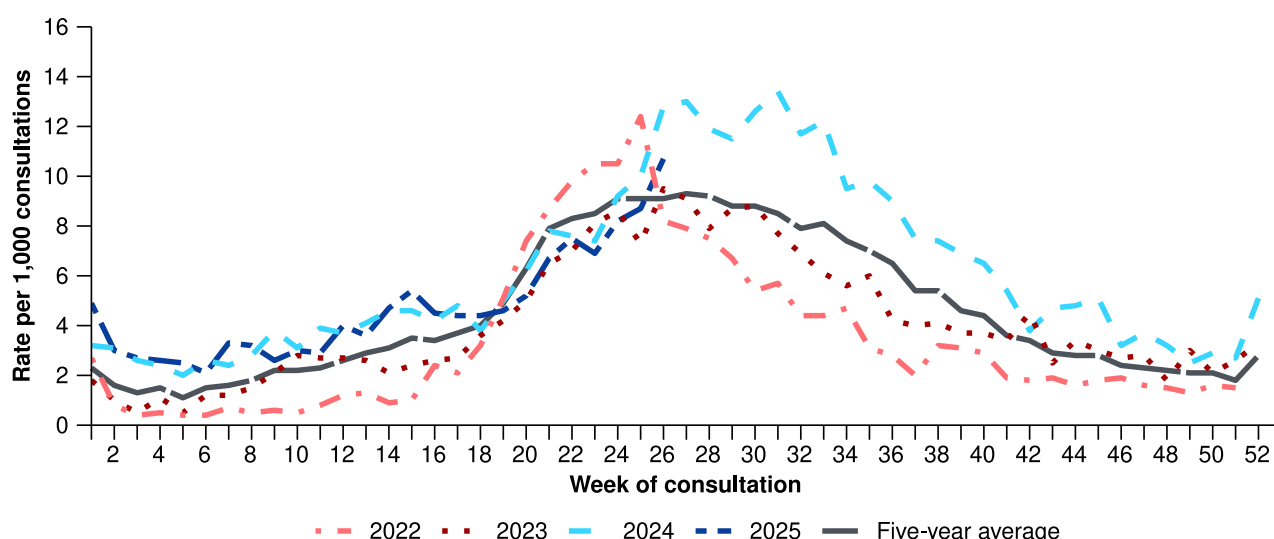
\* Rate per 100,000 population for the given time period. Population data are based on the Australian Bureau of Statistics (ABS) [Estimated Resident Population \(ERP\)](#) for the reference period June 2024, released 12 December 2024.

# Primary care surveillance

Primary care surveillance monitors the number and characteristics of people who have presented to their general practitioner with influenza-like-illness and provides insight on the different respiratory pathogens that are causing illness in the community.

- Sentinel general practice surveillance indicates general practice consultations for influenza-like-illness have continued to increase since May, following a period of relatively stable influenza-like-illness consultation rates from mid-January to late April 2025.
- In the last fortnight (16 June to 29 June 2025), there were more general practice consultations for influenza-like illness (9.7 notifications per 1,000 consultations per fortnight) than in the previous fortnight (7.6 notifications per 1,000 consultations per fortnight) (Figure 10).
- From mid-January to May 2025, influenza-like-illness notification rates remained relatively consistent with rates observed in 2024 but have been slightly higher than observed rates in the same period in previous years and the five-year average. Influenza-like-illness consultation rates increased in the last fortnight and are now above the five-year average; however, rates remain lower than the same time period last year (Figure 10).

**Figure 10: Rate of influenza-like-illness notifications per 1,000 consultations per week with sentinel general practice sites compared with the five-year average by year and week of consultation\*†, Australia, 2022 to 29 June 2025**



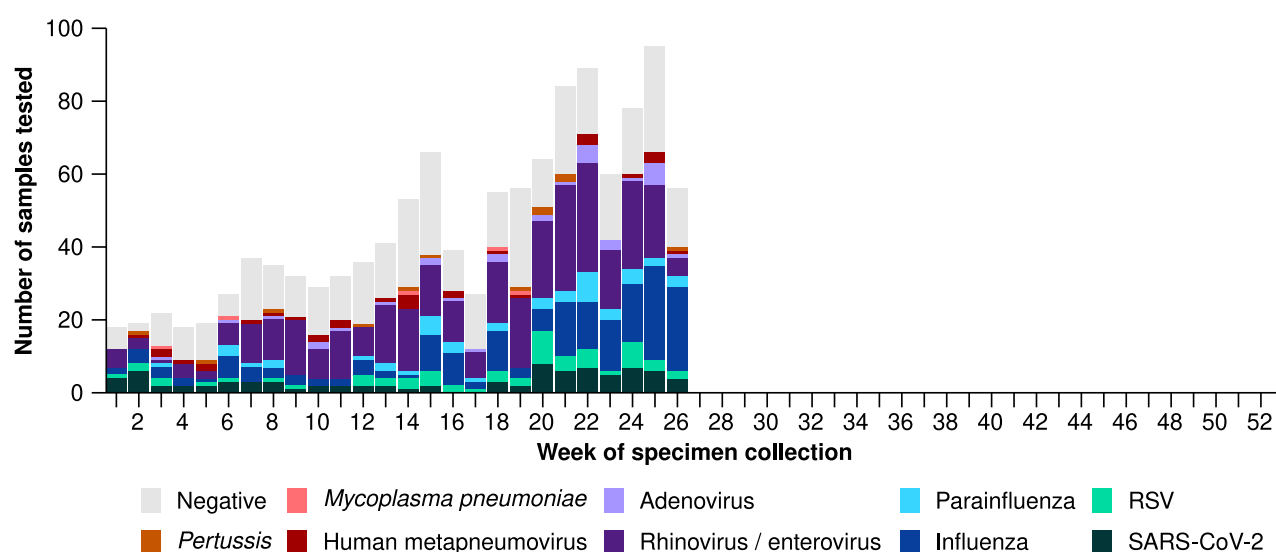
Source: Australian Sentinel Practice Research Network (ASPREN)

\* The years 2020 and 2021 are excluded when comparing the current season to historical periods when influenza virus has circulated without public health restrictions. As such, the five-year average includes the years 2018 to 2019 and 2022 to 2024. Please refer to the [Technical Supplement](#) for interpretation of the five-year average.

† Please refer to the [Technical Supplement](#) for notes on impact of COVID-19 on ASPREN data.

- In the last fortnight, 70.2% (106/151) of people attending general practice with influenza-like-illness who were tested have then tested positive for a respiratory pathogen.
- In the last fortnight, influenza (46.2%; 49/106) was the most commonly detected pathogen, followed by rhinovirus (23.6%; 25/106) and SARS-CoV-2 (9.4%; 10/106) (Figure 11).
- In the year to date, 66.7% (792/1,187) of people attending general practice with influenza-like-illness who were tested have then tested positive for a respiratory pathogen.
- In the year to date, rhinovirus (42.0%; 333/792) has been the most commonly detected pathogen, followed by influenza (23.6%; 187/792), SARS-CoV-2 (10.7%; 85/792), RSV (7.6%; 60/792), and adenovirus (4.0%; 32/792) (Figure 11).

**Figure 11: Number of samples tested for respiratory pathogens among people with influenza-like-illness attending sentinel general practice sites by respiratory pathogen and week of specimen collection, Australia, 1 January to 29 June 2025**



Source: Australian Sentinel Practice Research Network (ASPREN)

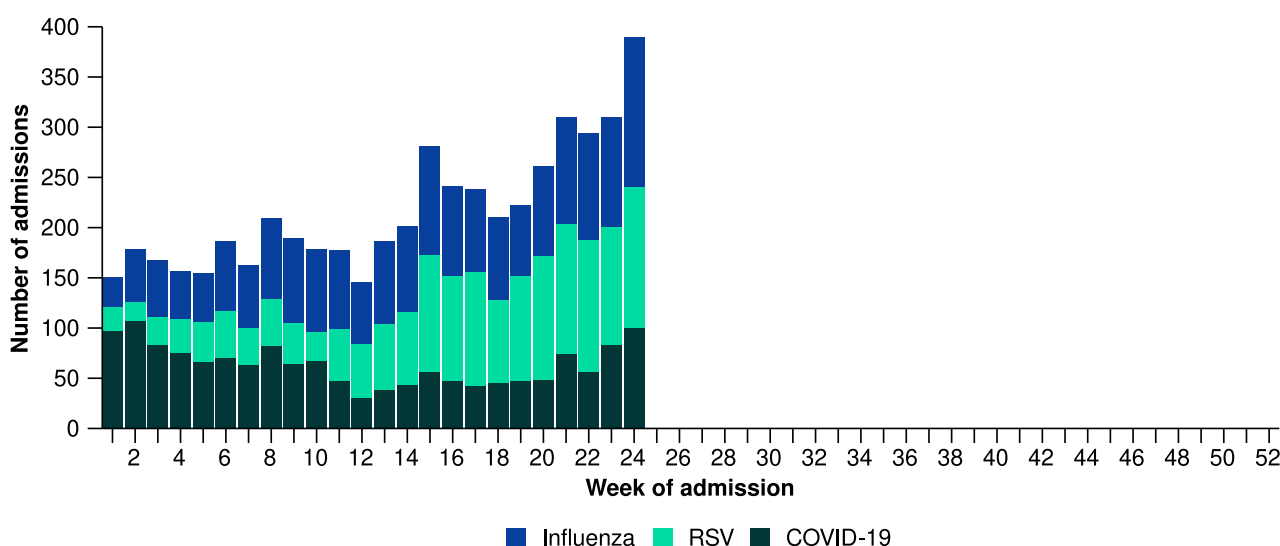
Note: All ASPREN swab samples are transported to the SA Pathology laboratory in Adelaide to be tested for viral and bacterial respiratory pathogens via a multiplex real-time reverse transcription polymerase chain reaction (RT-PCR) assay using in-house primers.

# Hospital-based surveillance

Hospital-based surveillance monitors persons with more severe illness who have been admitted to hospital for their respiratory illness (severe acute respiratory infections). Hospital-based surveillance also measures the ability of the health system to cope with the number of severe acute respiratory infection admissions to ensure delivery of safe, timely and quality health care.

- Sentinel hospital-based surveillance from the Influenza Complications Alert Network (FluCAN) shows the number of patients admitted with severe acute respiratory infections this year have followed an overall increasing trend since late March, after a period of relatively low and stable weekly admissions between January and March 2025. The length of hospital stay continues to vary only slightly between illnesses and the proportion of patients with a severe acute respiratory infection who were admitted directly to an intensive care has remained low.
- In the last severity reporting period (2 June to 15 June 2025), more patients were admitted to a sentinel hospital with a severe acute respiratory infection (n=700), than in the previous severity reporting period (n=604).
  - In the last severity reporting period, at sentinel hospitals there was a 40.8% increase in admissions with COVID-19 (from 130 to 183), a 21.5% increase in admissions with influenza (from 214 to 260), and a 1.2% decrease in admissions with RSV (from 260 to 257), compared to the previous severity reporting period.
- In the year to date for severity reporting (1 January to 15 June 2025), there have been 5,195 admissions with severe acute respiratory infections at sentinel hospitals. Most patients with a severe acute respiratory infection have been admitted with influenza (n=1,911) followed by RSV (n=1,736) (Figure 12).

**Figure 12: Total number of patients (children and adults) admitted with a severe acute respiratory infection to sentinel hospitals by disease and week of admission\*†‡, Australia, 1 January to 15 June 2025**

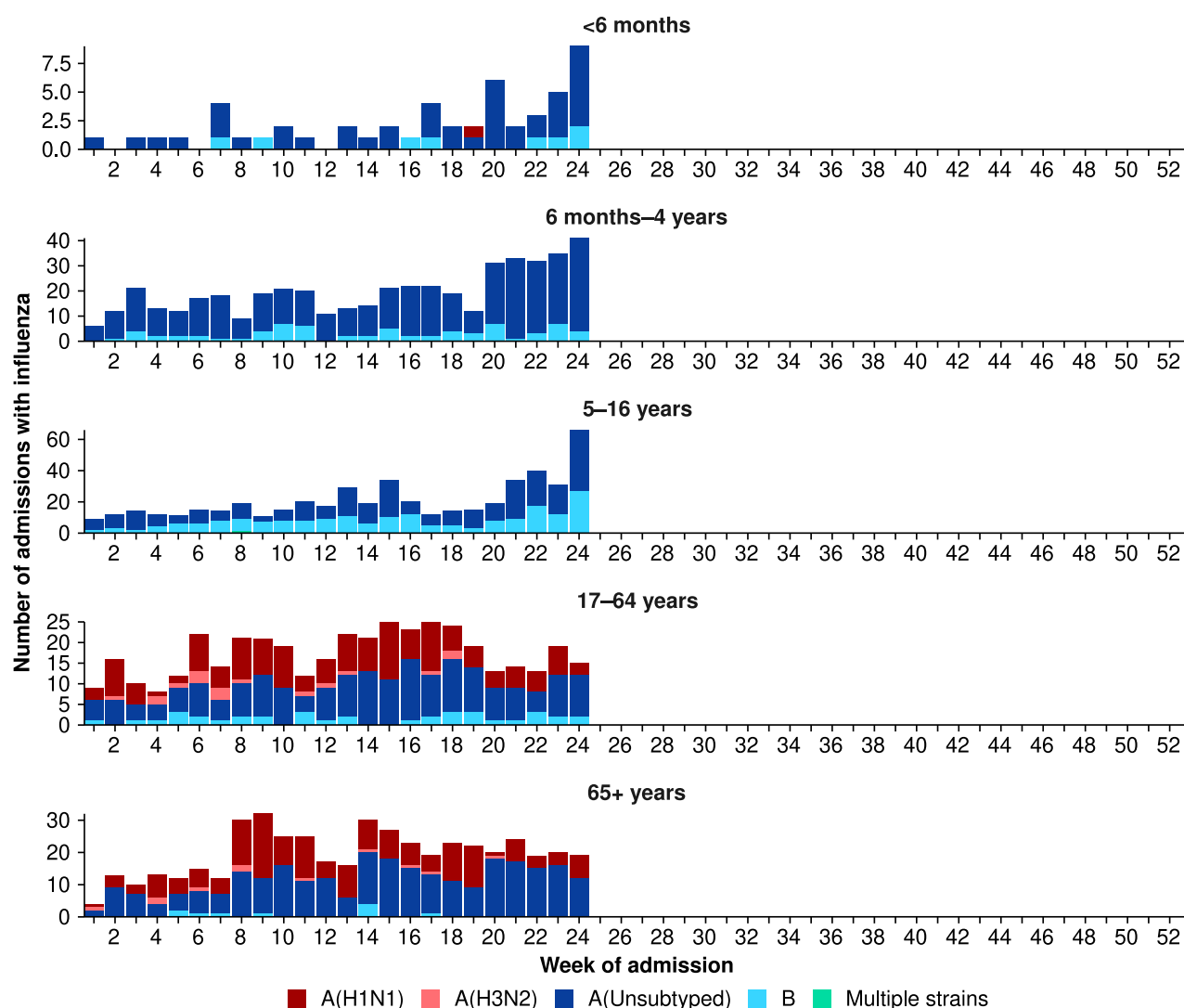


Source: Influenza Complications Alert Network (FluCAN)

- Patients admitted to sentinel hospitals with influenza have mostly been admitted with influenza A (83.0%; 1,587/1,911), while 16.9% (323/1,911) were admitted with influenza B.
  - Most hospital admissions with influenza A have been with influenza A(Unsubtyped) (76.9%; 1,220/1,587), followed by influenza A(H1N1) (21.4%; 339/1,587), and then influenza A(H3N2) (1.8%; 28/1,587).

- In the year to date for severity reporting, influenza A was the most commonly detected influenza subtype in all age groups. Influenza A(H1N1) and influenza A(H3N2) were more commonly observed in adults than children, while influenza B was more commonly observed in children. Of note, school aged children (5–16 years) had the highest proportion of influenza B compared with influenza A (Figure 13).
  - Trends in influenza subtypes should be interpreted with care as there may be differences in the number and selection of influenza samples that undergo typing.
  - While influenza B is often a good match with the seasonal influenza vaccine strain, influenza B can result in more severe infections in children.

**Figure 13: Number of patients admitted with influenza to sentinel hospitals by influenza subtype, age group, and week of admission\*, Australia, 1 January to 15 June 2025**



Source: Influenza Complications Alert Network (FluCAN)

\* Axis varies between age groups. The age distribution of admissions with influenza may not reflect the age distribution of all patients.



- In the year to date for severity reporting, more children (those aged 16 years and younger) have been admitted to sentinel hospitals with RSV than with influenza or COVID-19 (Table 2a).
- Children admitted to sentinel hospitals with influenza tended to be older than children admitted with COVID-19 or RSV (Table 2a).
- Children admitted to sentinel hospitals with RSV had a slightly longer length of hospital stay compared to children with influenza or COVID-19; however, the difference in the length of stay was minor. The proportion of children admitted directly to intensive care was relatively similar across COVID-19, influenza and RSV (Table 2a).

**Table 2a: Demographic characteristics and outcomes for children admitted with a severe acute respiratory infection to a sentinel hospital by disease, Australia, 1 January to 15 June 2025**

	COVID-19	Influenza	RSV
	Year to date for severity reporting (n=498)	Year to date for severity reporting (n=1,028)	Year to date for severity reporting (n=1,495)
<b>Age (years)</b>			
Median [IQR]	1 [0–4]	4 [1–8]	1 [0–2]
<b>Age group (years)</b>			
< 6 months	159 (31.9%)	52 (5.1%)	367 (24.5%)
6 months – 4 years	229 (46.0%)	474 (46.1%)	1022 (68.4%)
5–16 years	110 (22.1%)	502 (48.8%)	106 (7.1%)
<b>Indigenous status</b>			
Aboriginal and Torres Strait Islander	51 (10.2%)	80 (7.8%)	107 (7.2%)
<b>Length of hospital stay (days)†</b>			
Median [IQR]	1 [1–3]	1 [1–2]	2 [1–3]
<b>Patient admission location‡</b>			
Admitted to hospital ward	471 (94.6%)	979 (95.2%)	1413 (94.5%)
Admitted to intensive care directly	27 (5.4%)	49 (4.8%)	82 (5.5%)
<b>Discharge status†</b>			
Alive	375 (75.3%)	809 (78.7%)	1051 (70.3%)
Died	–	2 (0.2%)	1 (0.1%)
Incomplete/missing	123 (24.7%)	217 (21.1%)	443 (29.6%)

Source: Influenza Complications Alert Network (FluCAN)

\* Does not include patients with missing age; therefore, the sum of age-specific totals above may not equal the total number of patients.

† For patients who are still in hospital data may not be complete; therefore, these data are not included in the length of stay or discharge status. In addition, length of stay data excludes patients that acquired their infection in hospital.

‡ Admission location reflects the initial admission ward. Some patients may be initially admitted to general ward then later admitted to an intensive care and this is not reflected here. Does not include patients with missing admission location; therefore, the sum of admission location specific totals above may not equal the total number of patients.

The Paediatric Active Enhanced Disease Surveillance (PAEDS) network carries out enhanced sentinel hospital surveillance for some acute respiratory infections or conditions in children. PAEDS data for acute respiratory infections in children are presented in the Australian Respiratory Surveillance Reports in the sentinel hospital data from FluCAN. For additional information on [COVID-19 in children](#), [Paediatric Inflammatory Multisystem Syndrome \(PIMS-TS\) following COVID-19](#), [influenza in children](#), or [RSV in children](#) please visit the [PAEDS](#) webpages and dashboards.

- In the year to date for severity reporting, more adults (those aged 17 years and over) have been admitted to sentinel hospitals with COVID-19 than with influenza or RSV (Table 2b).
- Adults admitted to sentinel hospitals with COVID-19 or RSV were predominately 65 years and over, whereas the proportion of admissions with influenza was only slightly higher in the 65 years and over age group compared to the 17–64 years age group (Table 2b).
- Adults admitted to sentinel hospitals with COVID-19 had a slightly longer length of hospital stay compared to adults with influenza or RSV. A higher proportion of adults with influenza were admitted directly to intensive care, compared to adults admitted with COVID-19 or RSV (Table 2b).
- Sadly, there have been a number of adults admitted with a severe acute respiratory infection who have died in hospital (Table 2b).

**Table 2b: Demographic characteristics and outcomes for adults admitted with a severe acute respiratory infection to a sentinel hospital by disease, Australia, 1 January to 15 June 2025**

	COVID-19	Influenza	RSV
	Year to date for severity reporting (n=1,050)	Year to date for severity reporting (n=883)	Year to date for severity reporting (n=241)
<b>Age (years)</b>			
Median [IQR]	75 [62–84]	66 [53–77]	73 [61–82]
<b>Age group (years)</b>			
17–64 years	303 (28.9%)	413 (46.8%)	74 (30.7%)
65 years and over	747 (71.1%)	470 (53.2%)	167 (69.3%)
<b>Indigenous status</b>			
Aboriginal and Torres Strait Islander	69 (6.6%)	66 (7.5%)	21 (8.7%)
<b>Length of hospital stay (days)†</b>			
Median [IQR]	5 [2–9]	4 [2–7]	4 [2–8]
<b>Patient admission location‡</b>			
Admitted to hospital ward	986 (93.9%)	792 (89.7%)	223 (92.5%)
Admitted to intensive care directly	64 (6.1%)	91 (10.3%)	18 (7.5%)
<b>Discharge status†</b>			
Alive	781 (74.4%)	648 (73.4%)	150 (62.2%)
Died	37 (3.5%)	18 (2.0%)	10 (4.1%)
Incomplete/missing	232 (22.1%)	217 (24.6%)	81 (33.6%)

Source: Influenza Complications Alert Network (FluCAN)

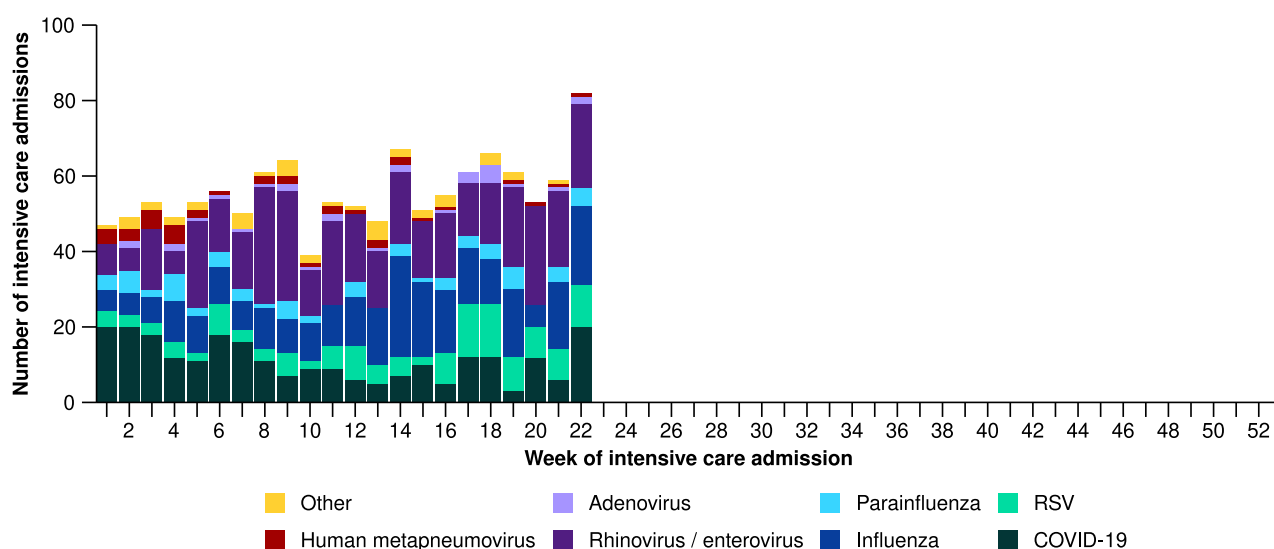
\* Does not include patients with missing age; therefore, the sum of age-specific totals above may not equal the total number of patients.

† For patients who are still in hospital data may not be complete; therefore, these data are not included in the length of stay or discharge status. In addition, length of stay data excludes patients that acquired their infection in hospital.

‡ Admission location reflects the initial admission ward. Some patients may be initially admitted to general ward then later admitted to an intensive care and this is not reflected here. Does not include patients with missing admission location; therefore, the sum of admission location specific totals above may not equal the total number of patients.

- Please note, sentinel intensive care data are updated each month, as such the sentinel intensive care surveillance data presented here have not been updated since the previous report.
- Sentinel intensive care surveillance shows the number of patients admitted to intensive care with severe acute respiratory infections has remained low and stable, though now appears to be increasing.
- In the last severity reporting period (5 May to 1 June 2025), more patients have been admitted to a sentinel intensive care with a severe acute respiratory infection (n=242), than in the previous severity reporting period (n=221) (Figure 14).
- In the year to date for severity reporting (1 January to 1 June 2025), most patients were admitted to sentinel intensive care with rhinovirus / enterovirus, followed by influenza (Figure 14; Table 3).

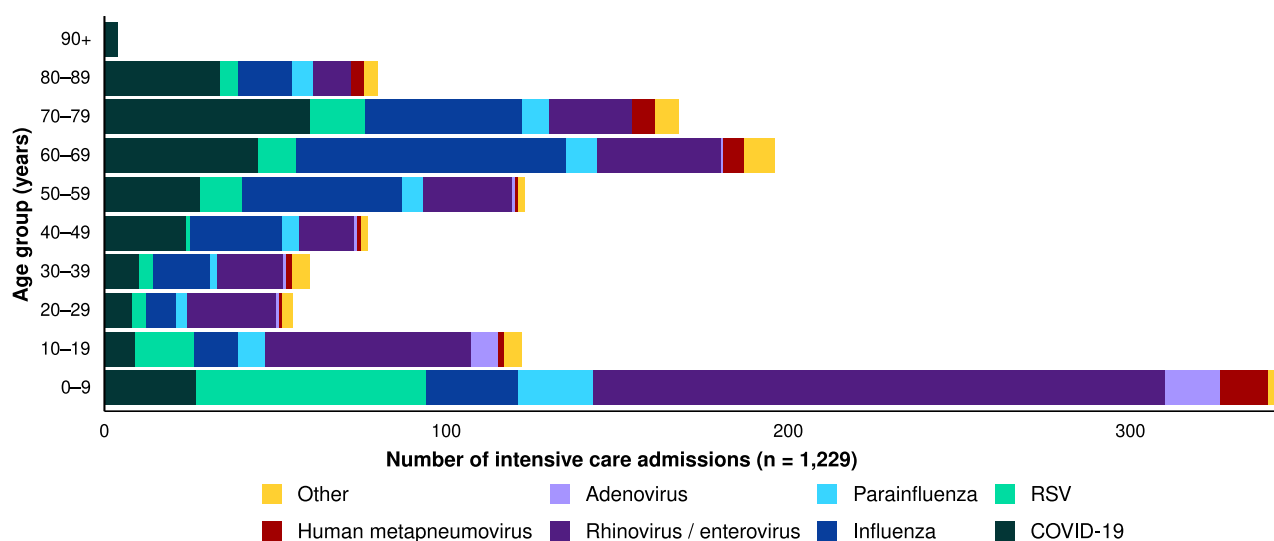
**Figure 14: Number of patients admitted with severe acute respiratory infections to a sentinel intensive care by disease and week of admission, Australia, 1 January to 1 June 2025**



Source: Short Period Incidence Study of Severe Acute Respiratory Infection (SPRINT-SARI) Australia

Note: A range of diagnostic testing procedures are utilised across hospitals in Australia. SPRINT-SARI does not specify which diagnostic testing method should be utilised as this is the domain of the hospital and treating clinicians. Therefore, virological data from SPRINT-SARI should be interpreted with care.

**Figure 15: Number of patients admitted with severe acute respiratory infections to a sentinel intensive care by disease and age group\*, Australia, 1 January to 1 June 2025**



Source: Short Period Incidence Study of Severe Acute Respiratory Infection (SPRINT-SARI) Australia

Note: 4.4% (52/1,173) of patients had co-infections of respiratory pathogens; therefore, the sum of pathogen-specific totals above may not equal the total number of severe acute respiratory infection patients.

\* The age distribution of severe acute respiratory infection intensive care admissions may not reflect the age distribution of all patients.

- In the year to date for severity reporting, admissions to a sentinel intensive care with COVID-19 or influenza have generally been among older people. In contrast, admissions to a sentinel intensive care with rhinovirus or RSV have been among younger people, primarily those aged 0–9 years old (Figure 15; Table 3).
- A higher proportion of patients with COVID-19, influenza and parainfluenza required invasive mechanical ventilation, and the length of ventilation was highest among those with influenza. The length of intensive care stay was relatively similar across diseases (Table 3).
- Most patients admitted to a sentinel intensive care with a severe acute respiratory infection have been discharged home. Sadly, a number of patients have died in hospital (Table 3).

**Table 3: Demographic characteristics and outcomes of patients admitted with a severe acute respiratory infection to a sentinel intensive care by disease\*†, Australia, 1 January to 1 June 2025**

	COVID-19	hMPV	Influenza	Parainfluenza	Rhinovirus	RSV	Other
	Year to date for severity reporting (n=249)	Year to date for severity reporting (n=38)	Year to date for severity reporting (n=281)	Year to date for severity reporting (n=69)	Year to date for severity reporting (n=385)	Year to date for severity reporting (n=137)	Year to date for severity reporting (n=70)
<b>Age (years)</b>							
Median [IQR]	65 [42–75]	41 [3–72]	60 [41–68]	38 [8–68]	12 [5–50]	10 [2–59]	22 [9–64]
<b>Indigenous status</b>							
Aboriginal and Torres Strait Islander	28 (11.2%)	3 (7.9%)	34 (12.1%)	5 (7.2%)	43 (11.2%)	22 (16.1%)	6 (8.6%)
Non-Indigenous	221 (88.8%)	35 (92.1%)	247 (87.9%)	64 (92.8%)	342 (88.8%)	115 (83.9%)	64 (91.4%)
<b>Received invasive mechanical ventilation</b>							
Number (%)	76 (30.5%)	10 (26.3%)	97 (34.5%)	28 (40.6%)	90 (23.4%)	22 (16.1%)	28 (40.0%)
<b>Length of invasive mechanical ventilation (days)*</b>							
Median [IQR]	3 [1–6]	2 [2–7]	5 [2–10]	3 [1–12]	3 [1–6]	3 [1–5]	2 [1–6]
<b>Length of intensive care stay (days)*</b>							
Median [IQR]	3 [2–6]	3 [1–6]	3 [2–6]	2 [1–6]	2 [1–5]	2 [1–4]	3 [2–7]
<b>Length of hospital stay (days)*</b>							
Median [IQR]	7 [4–15]	9 [6–19]	8 [4–16]	6 [3–12]	5 [3–11]	5 [3–10]	8 [4–17]
<b>Patient outcome†</b>							
Ongoing care in intensive care	11 (4.4%)	–	14 (5.0%)	3 (4.3%)	10 (2.6%)	5 (3.6%)	1 (1.4%)
Ongoing care in hospital ward	9 (3.6%)	1 (2.6%)	9 (3.2%)	4 (5.8%)	19 (4.9%)	3 (2.2%)	3 (4.3%)
Transfer to other hospital / facility	40 (16.1%)	5 (13.2%)	48 (17.1%)	7 (10.1%)	40 (10.4%)	16 (11.7%)	7 (10.0%)
Discharged home	146 (58.6%)	29 (76.3%)	181 (64.4%)	49 (71.0%)	295 (76.6%)	107 (78.1%)	49 (70.0%)
Died in hospital	42 (16.9%)	3 (7.9%)	28 (10.0%)	5 (7.2%)	20 (5.2%)	6 (4.4%)	10 (14.3%)

Source: Short Period Incidence Study of Severe Acute Respiratory Infection (SPRINT-SARI) Australia

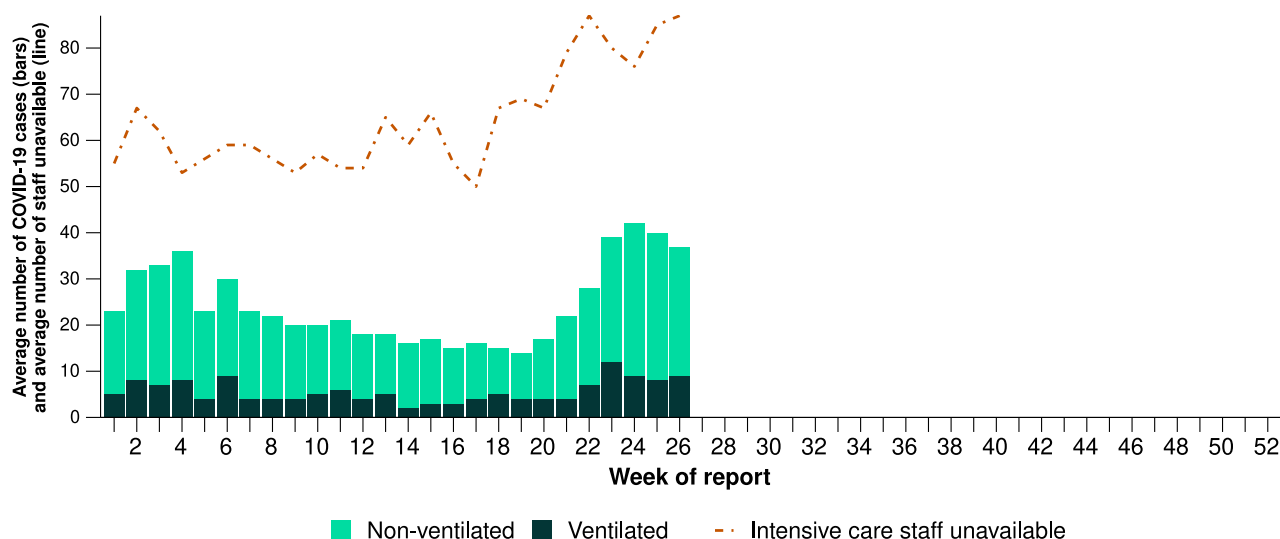
Note: 4.4% (52/1,173) of patients had co-infections of respiratory pathogens; therefore, the sum of pathogen-specific totals above may not equal the total number of severe acute respiratory infection patients.

\* For patients receiving ongoing care in intensive care data may not be complete; therefore, data are not included in the length of ventilation or stay.

† Patients who have been admitted with no discharge information for less than 90 days have been assumed to have ongoing care in the hospital. Patients who have no outcome entered or have been admitted for more than 90 days with no discharge information have been treated as missing.

- Intensive care occupancy surveillance shows the number of COVID-19 cases occupying intensive care beds and number of staff unavailable to work have generally continued to increase since May, following a decreasing or fluctuating trend from mid-January to early May 2025.
- In the last fortnight (16 June to 29 June 2025), there were less COVID-19 cases in intensive care across Australia than in the previous fortnight (Figure 16).
- In the last fortnight, there were more intensive care staff unavailable to work due to COVID-19 exposure or illness across Australia than in the previous fortnight (Figure 16).

**Figure 16: Average number of COVID-19 cases in intensive care and the average number of intensive care staff unavailable to work due to COVID-19 exposure or illness by week of report<sup>\*\*†</sup>, Australia, 1 January to 29 June 2025**



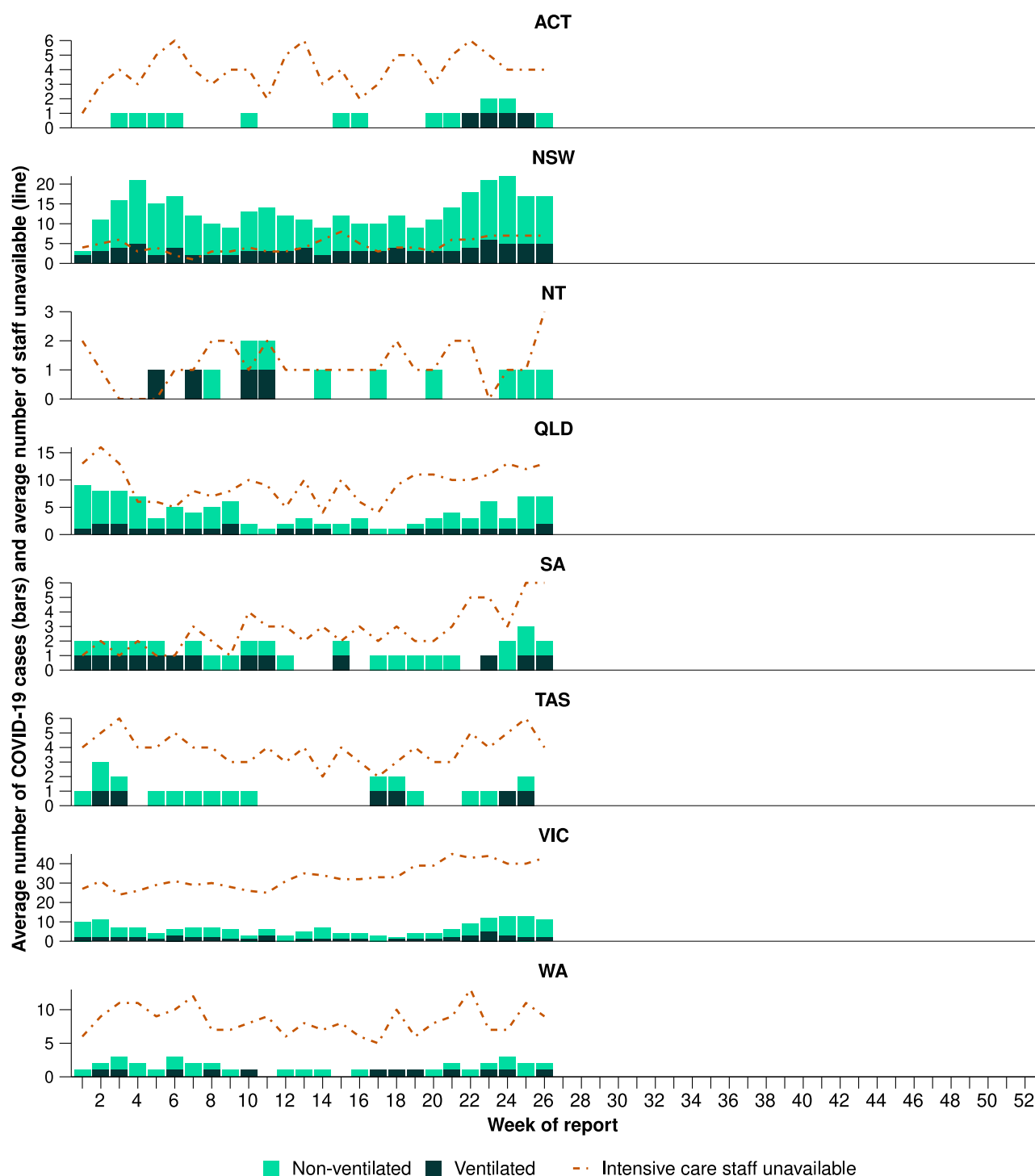
Source: Critical Health Resource Information System (CHRIS)

\* Average number of ventilated and non-ventilated COVID-19 cases in intensive care includes only active COVID-19 cases (those in isolation) and does not include cleared COVID-19 cases.

† Intensive care staff include both medical and nursing staff. Staff unavailability will be underestimated in NSW as most public hospitals in NSW do not report staff unavailability.

- In the last fortnight, the number of COVID-19 cases in intensive care decreased in NSW, and Vic, remained stable in Tas, the NT, SA, the ACT, and WA, and increased in Qld compared with the previous fortnight (Figure 17).
- In the last fortnight, the number of intensive care staff unavailable to work due to COVID-19 exposure or illness remained stable in NSW, Vic, Qld, and the ACT, and increased in Tas, the NT, SA, and WA compared with the previous fortnight (Figure 17).

**Figure 17: Average number of COVID-19 cases in intensive care and the average number of intensive care staff unavailable to work due to COVID-19 exposure or illness by jurisdiction and week of report\*\*†‡, Australia, 1 January to 29 June 2025**



Source: Critical Health Resource Information System (CHRIS)

\* Axis varies between jurisdictions.

† Average number of ventilated and non-ventilated COVID-19 cases in intensive care includes only active COVID-19 cases (those in isolation) and does not include cleared COVID-19 cases.

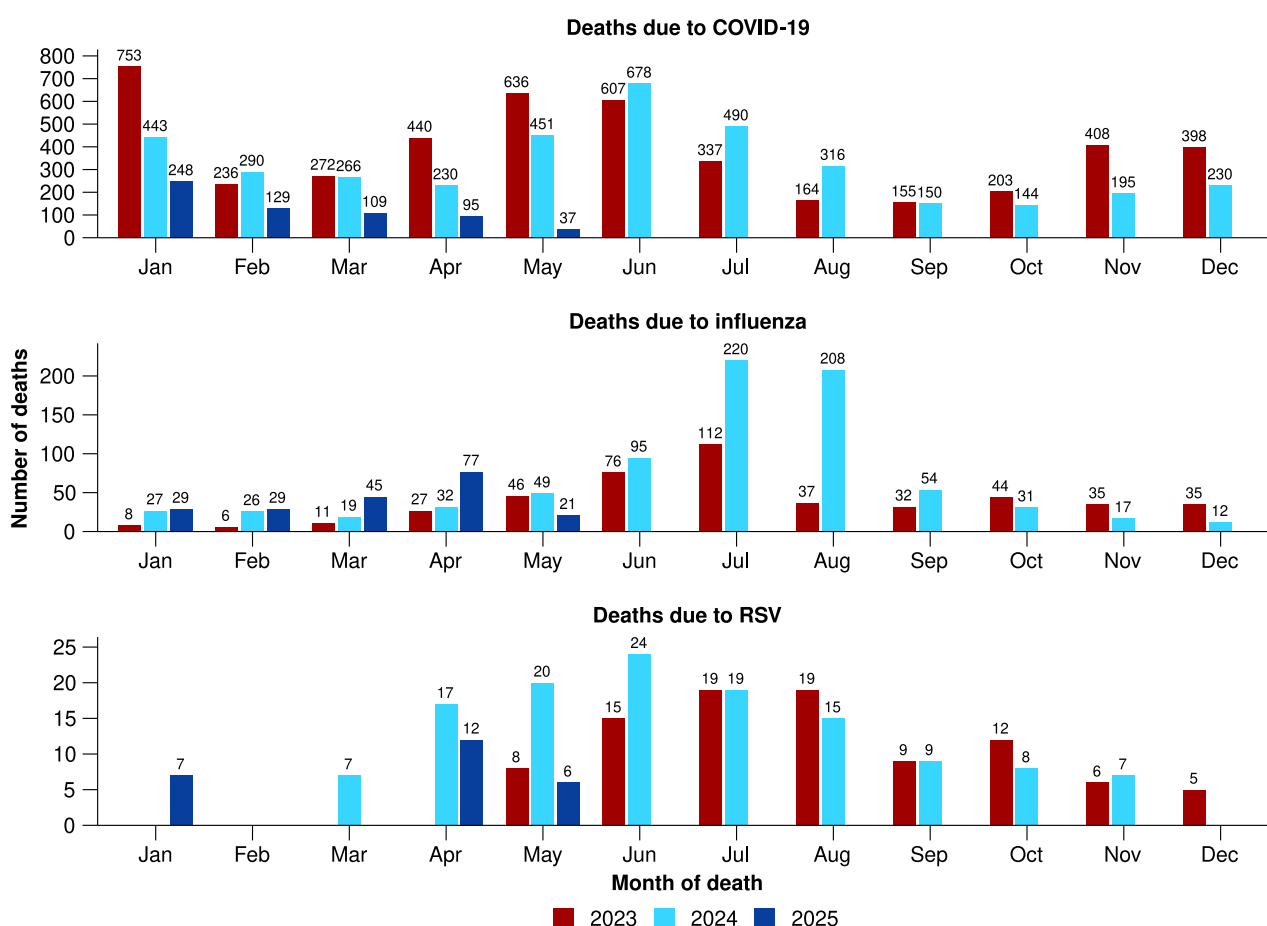
‡ Intensive care staff include both medical and nursing staff. Staff unavailability will be underestimated in NSW as most public hospitals in NSW do not report staff unavailability.

# Mortality surveillance

Death registrations can provide information on the scale and severity of disease associated with acute respiratory infections. For more information on death registrations including completeness, timeliness, and definitions of deaths involving (both *due to* and *with*), *due to* and *with* acute respiratory infections, refer to the [Technical Supplement](#).

- COVID-19 has been the leading cause of acute respiratory infection mortality across 2023–2025.
- Since the end of 2021, a pattern has been observed for COVID-19 where there are two peaks of mortality during the year - one occurring between November and January and the other occurring between May and August. Following an increased number of deaths occurring between November 2024 and January 2025 (which was much lower than in previous years), deaths *due to* COVID-19 in February and March 2025 were substantially lower than in January 2025 (Figure 18a).
- There were 581 deaths *due to* COVID-19 in the first four months of 2025. This is 52.7% fewer deaths *due to* COVID-19 than the 1,229 deaths *due to* COVID-19 that occurred between January and April 2024 and 65.8% fewer than the 1,701 deaths *due to* COVID-19 that occurred between January and April 2023 (Figure 18a).

**Figure 18a: Provisional numbers of deaths *due to* an acute respiratory infection\*† by month, year, and disease, Australia, 1 January 2023 to 31 May 2025**



Source: Australian Bureau of Statistics, [Provisional Mortality Statistics](#), released 27 June 2025.

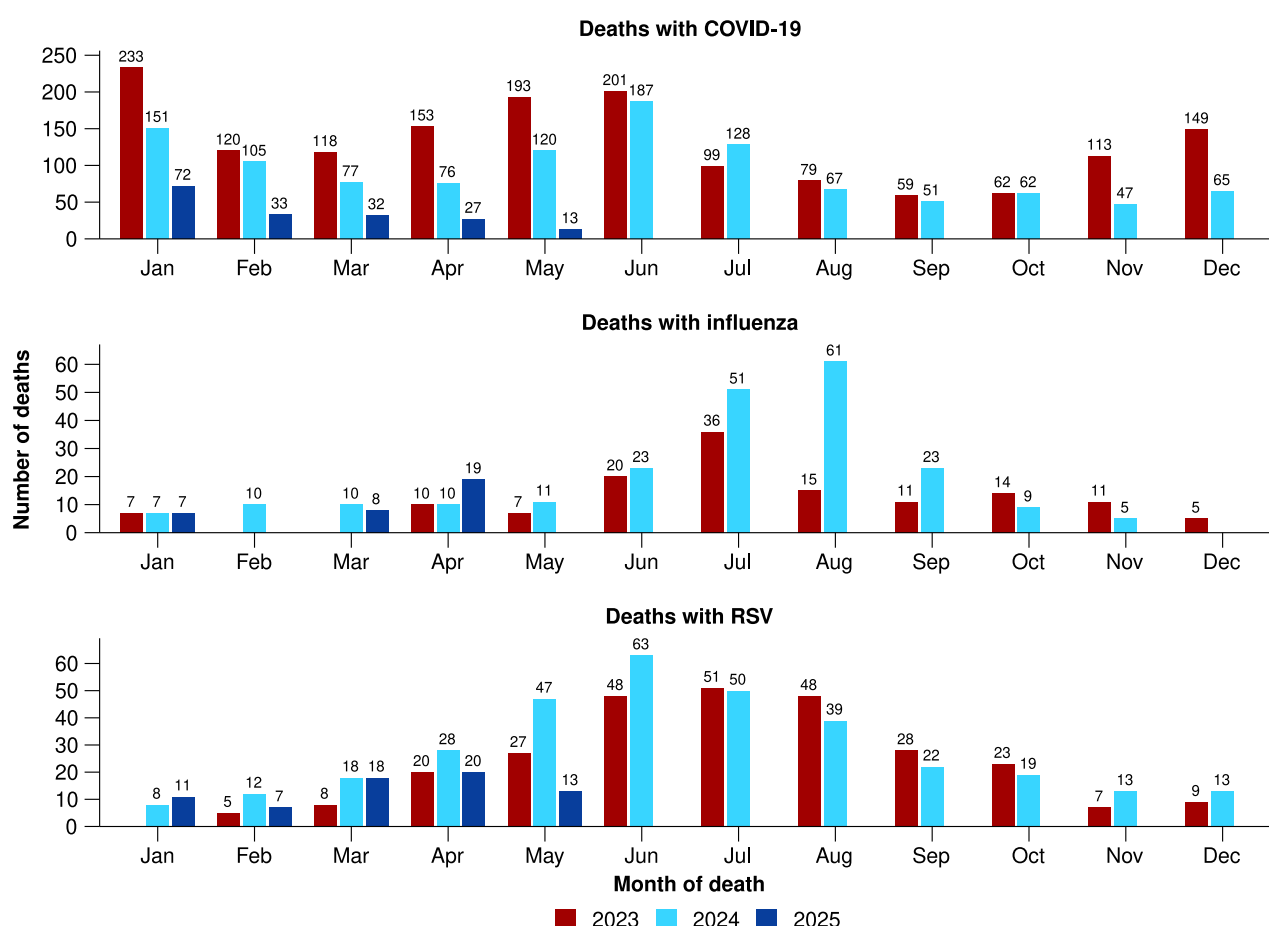
\* Axis varies between acute respiratory infections.

† Data is provisional and subject to change. It can take several weeks for death registrations to be reported, processed, coded, validated, and tabulated. Therefore, the data shown here may be incomplete. Data for some months were not published by the ABS due to small counts, and therefore not reported here. Data includes all deaths (both doctor and coroner certified) that occurred and were registered by 31 May 2025.



- There have been 180 deaths *due to* influenza in the first four months of 2025. This is 73.1% more deaths *due to* influenza than the 104 deaths *due to* influenza that occurred between January and April 2024 and substantially more than the 52 deaths *due to* influenza that occurred between January and April 2023 (Figure 18a). As noted previously, however, there was also a higher number of influenza cases notified in this period than in previous years.
- There have been 19 deaths *due to* RSV published in the first four months of 2025 (Figure 18a).
- The mortality rate for deaths *due to* COVID-19 or influenza for Aboriginal and Torres Strait Islander people was higher than for non-Indigenous people across each year in 2022–2024.
- Since November 2024, there have been substantially fewer deaths *with* COVID-19 each month than there were in the corresponding months of the previous years (Figure 18b).
- Deaths *with* influenza have increased in April 2025 and are higher than in 2023 and 2024 (Figure 18b).
- Deaths *with* RSV are at similar levels in 2025 to in 2023 and 2024 (Figure 18b).
- All three of these acute respiratory infections are more likely to cause death in older age groups than younger age groups.

**Figure 18b: Provisional numbers of deaths *with* an acute respiratory infection\*† by month, year, and disease, Australia, 1 January 2023 to 31 May 2025**



Source: Australian Bureau of Statistics, [Provisional Mortality Statistics](#), released 27 June 2025.

\* Axis varies between acute respiratory infections.

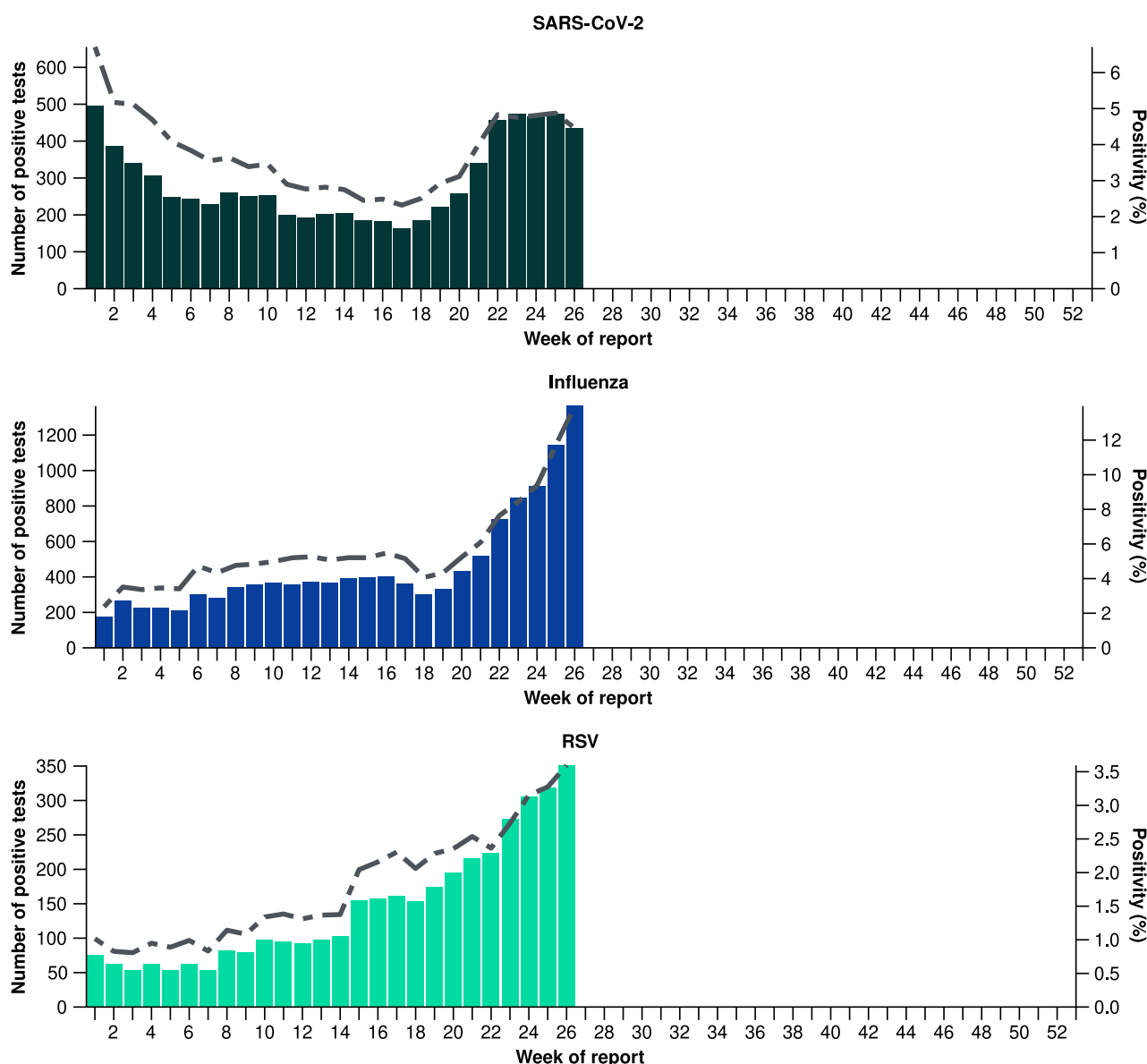
† Data is provisional and subject to change. It can take several weeks for death registrations to be reported, processed, coded, validated, and tabulated. Therefore, the data shown here may be incomplete. Data for some months were not published by the ABS due to small counts, and therefore not reported here. Data includes all deaths (both doctor and coroner certified) that occurred and were registered by 31 May 2025.

# Laboratory surveillance

Sentinel laboratory surveillance monitors and characterises respiratory pathogens to provide information on what pathogens are circulating, potential changes in the pathogens that might affect their infectiousness, severity, ability to evade vaccine and/or infection-acquired immunity, or resistance to antivirals.

- In the last fortnight (16 June to 29 June 2025), SARS-CoV-2 test positivity increased to 4.6% (766/16,632), influenza test positivity increased to 12.8% (2,504/19,509), and RSV test positivity increased to 3.3% (550/16,632) (Figure 19).

**Figure 19: Number of tests positive (bars) and test positivity (line) for SARS-CoV-2, influenza or RSV of those specimens tested by sentinel laboratories by week of report\*†, Australia, 1 January to 29 June 2025**



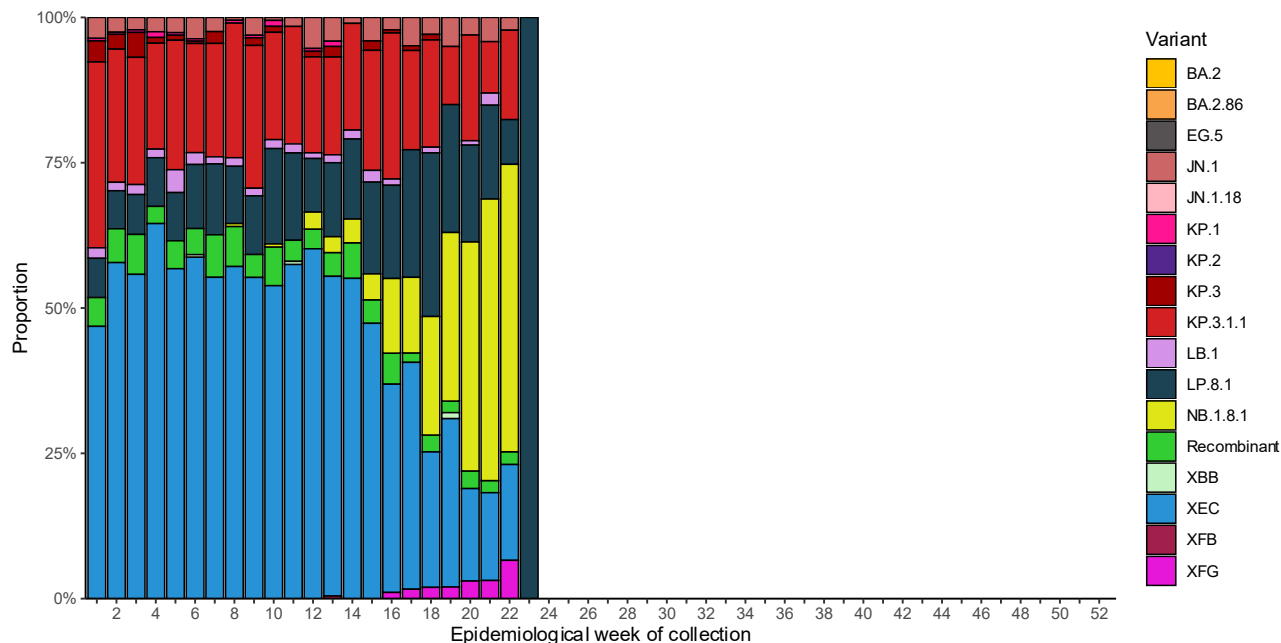
Source: Sentinel laboratories, including National Influenza Centres

\* Number of specimens tested excludes data from WA as testing denominator data are different for the three pathogens in Western Australia.

† A small minority of total samples from Victoria are tested only by respiratory panel (influenza, parainfluenza, adenovirus, human metapneumovirus, seasonal coronaviruses, RSV, and some picornaviruses) but not for SARS-CoV-2. These minority samples include only forensic materials; all other samples are tested by respiratory panel and SARS-CoV-2 assay.

- *Please note, AusTrakka SARS-CoV-2 sequencing data are updated each month, as such SARS-CoV-2 sequence data presented here have not been updated since the previous report.*
- There were 284 SARS-CoV-2 sequences uploaded to AusTrakka with dates of collection in the last 28 days (19 May to 15 June 2025). These sequences were from NSW, Qld, SA, Tas, Vic and WA, with the most recent collection date 2 June 2025.
- All sequences were assigned to the BA.2.86 sub-lineage within B.1.1.529 (Omicron), or recombinants consisting of one or more Omicron sub-lineages (Figure 20a/b). In the last 28 days:
  - 29.6% (84/284) of sequences were from the sub-sub-lineages JN.1 (BA.2.86.1.1), including KP.3 (31/284)
  - 70.4% (200/284) of sequences were recombinant or recombinant sub-lineages, including XEC (44/284) and the most recently designated World Health Organization (WHO) variant under monitoring (VUM) NB.1.8.1 (138/284)
  - NB.1.8.1, a newly designated World Health Organization (WHO) Variant Under Monitoring (VUM) is the dominant sub-lineage in the past 28 days, accounting for 48.6% (138/284) of sequences
  - there were no BA.1, BA.3, BA.4, BA.5 or other BA.2 sub-sub-lineage sequences.
- NB.1.8.1 is the dominant sub-lineage in the last 28 days, accounting for 48.6% (138/284) of sequences (Figure 20a).
- The WHO have identified certain sub-sub-lineages and recombinants as VUM or variants of interest (VOI) because of their epidemiological, pathological, or immunological features of concern. A select number are highlighted below due to their relevance in the Australian context:
  - there are 313 NB.1.8.1 sequences in AusTrakka, with 138 collected in the last 28 days. The overall risk evaluation by the WHO is considered low for this variant.
  - there are 617 LP.8.1 sequences in AusTrakka, with 39 collected in the last 28 days.
  - there are 368 LB.1 sequences in AusTrakka, with four sequences collected in the last 28 days.
  - there are 2,954 KP.3.1.1 sequences in AusTrakka, with 31 sequences collected in the last 28 days.
  - there are 3,343 XEC sequences in AusTrakka, with 44 sequences collected in last 28 days.
  - there are 24 XFG sequences in AusTrakka, with 12 sequences collected in the last 28 days.

**Figure 20a: SARS-CoV-2 Omicron sub-lineage\* sequences by sample collection date, showing the proportions of sequences per week<sup>†‡</sup>, Australia, 1 January to 15 June 2025**

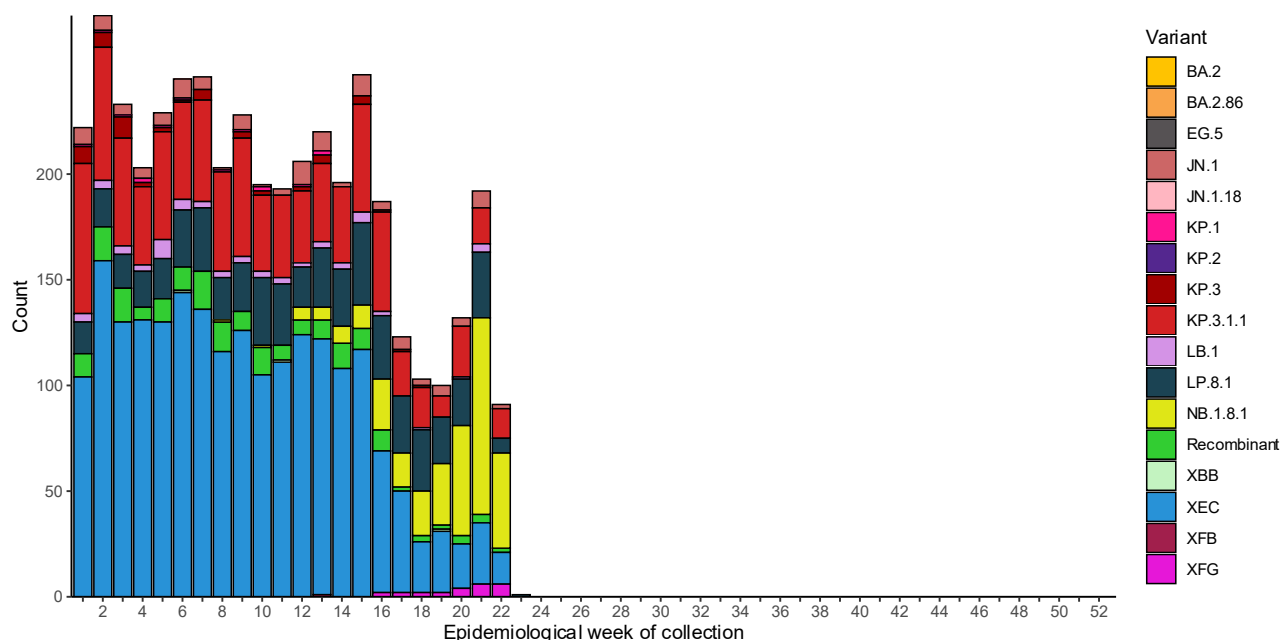


Source: AusTrakka

\* Some sub-sublineages are shown alongside their parent lineage, but not included in the parent lineage totals. For instance, KP.2 and KP.3 are sub-sub lineages of JN.1, so the total of JN.1 sequences will be higher than shown in the corresponding colour alone, and should include the KP.2 and KP.3 totals.  
<sup>†</sup> Sequences in AusTrakka aggregated by week and reported based on date of sample collection, not date of sequencing.

<sup>‡</sup> Proportions in Figure 20a may not be representative when sequence numbers are small; refer to Figure 20b. Data for earlier weeks may change between reporting periods as sequences with older collection dates are uploaded. These numbers are not equivalent to number of cases, as there are many cases which may not be sequenced. Non-VOI and non-VUM Omicron sub-lineages have been collapsed into parent lineages BA.1, BA.2, BA.3, BA.4 and BA.5.

**Figure 20b: SARS-CoV-2 Omicron sub-lineage\* sequences by sample collection date, showing the count of sequences per week<sup>†‡</sup>, Australia, 1 January to 15 June 2025**



Source: AusTrakka

\* Some sub-sublineages are shown alongside their parent lineage, but not included in the parent lineage totals. For instance, KP.2 and KP.3 are sub-sub lineages of JN.1, so the total of JN.1 sequences will be higher than shown in the corresponding colour alone, and should include the KP.2 and KP.3 totals.  
<sup>†</sup> Sequences in AusTrakka aggregated by week and reported based on date of sample collection, not date of sequencing.

<sup>‡</sup> Data for earlier weeks may change between reporting periods as sequences with older collection dates are uploaded. These numbers are not equivalent to number of cases, as there are many cases which may not be sequenced. Non-VOI and non-VUM Omicron sub-lineages have been collapsed into parent lineages BA.1, BA.2, BA.3, BA.4 and BA.5.

- In the year to date, the WHO Collaborating Centre for Reference and Research on Influenza has antigenically characterised 2,249 influenza viruses from Australia (Table 4), of which:
  - 75.0% (1,686/2,249) have been influenza A(H1N1)
  - 11.2% (252/2,249) have been influenza A(H3N2)
  - 13.8% (311/2,249) have been influenza B/Victoria.
- In the year to date, there have been no influenza B/Yamagata viruses characterised (Table 4). The last influenza B/Yamagata virus characterised in Australia was in a sample from 2020.
- Of the influenza A(H1N1) samples tested for neuraminidase inhibitor resistance, 0.2% (1/507) demonstrated highly reduced inhibition to Oseltamivir. None of the influenza A(H3N2) samples tested for neuraminidase inhibitor resistance demonstrated highly reduced inhibition to Oseltamivir.
- None of the samples tested demonstrated highly reduced inhibition to Zanamivir.

**Table 4: Australian influenza viruses typed by haemagglutination inhibition assay and jurisdiction\*†, 1 January to 29 June 2025**

Strain	ACT	NSW	NT	Qld	SA	Tas	Vic	WA	Total
A(H1N1)	219	149	466	72	40	256	455	29	<b>1,686</b>
A(H3N2)	11	21	109	17	4	17	68	5	<b>252</b>
B/Victoria lineage	46	42	53	7	17	26	105	15	<b>311</b>
B/Yamagata lineage	0	0	0	0	0	0	0	0	<b>0</b>
<b>Total</b>	<b>276</b>	<b>212</b>	<b>628</b>	<b>96</b>	<b>61</b>	<b>299</b>	<b>628</b>	<b>49</b>	<b>2,249</b>

Source: World Health Organization (WHO) Collaborating Centre for Reference and Research on Influenza

\*Viruses tested by the WHO Collaborating Centre for Reference and Research on Influenza are not necessarily a random sample of all those in the community and early-year data may be based on limited samples received. There may be up to a month delay on reporting of samples.

† Jurisdiction indicates the residential location for the individual tested, not the submitting laboratory.

# Vaccine coverage, effectiveness and match

Vaccine coverage, effectiveness and match for acute respiratory infections are monitored from several data sources in Australia. Refer to the [Technical Supplement](#) for more information.

## Vaccine coverage

- In Australia, regular COVID-19 vaccinations are the best way to maintain protection against severe disease and death from COVID-19. Most adults should receive a COVID-19 vaccine each year to stay protected against severe illness, hospitalisation and death. Adults aged 75 years and over should get vaccinated every six months.
  - More information on COVID-19 vaccines in Australia is available via the [department's COVID-19 webpages](#) or from the [National Centre for Immunisation Research and Surveillance \(NCIRS\)](#).
- Nationally, 8.8% of adults (aged 18 years and over) have received a COVID-19 vaccine in the past six months (Table 5).
- Nationally, less adults have received a COVID-19 vaccine in the past 12 months (11.1%; Table 5), compared to the 12 months prior (14.6% from 26 June 2023 to 23 June 2024).
  - In the past 12 months, vaccine coverage decreased in all age groups, with the largest decrease seen in 65–74 years age group (from 35% in the 12 months prior to 26.6% in the past 12 months).
- There is substantial variation in COVID-19 vaccine coverage across age groups, ranging from 4.8% in adults aged 18–64 years to 42.2% in adults aged 75 years and over. Vaccine coverage increases with increasing age (Table 5).
- There is some variation in vaccine coverage across jurisdictions, ranging from 4.1% in the NT to 18.9% in Tas (Table 5).

**Table 5: COVID-19 vaccine coverage<sup>\*†‡</sup> by age group and jurisdiction, Australia, 24 June 2024 to 29 June 2025**

Age group	ACT	NSW	NT	Qld	SA	Tas	Vic	WA	Total
<b>Past 12 months (24 June 2024 to 29 June 2025)</b>									
18–64 years	10.5	4.1	2.1	4.5	4.6	8.6	5.2	4.6	4.8
65–74 years	47.5	24.8	14.7	25.4	27.3	39.5	27.2	27.1	26.6
≥ 75 years	65.9	40.1	26.6	40.9	41.8	56.9	41.5	44.0	42.2
All ages (18 years and over)	18.4	10.3	4.1	10.6	12.0	18.9	11.2	10.8	11.1
<b>Past 6 months (30 December 2024 to 29 June 2025)</b>									
18–64 years	7.8	2.9	1.3	3.3	3.4	6.4	3.8	3.6	3.5
65–74 years	39.8	20.1	10.7	20.6	22.6	33.0	22.4	22.7	21.9
≥ 75 years	56.5	33.0	20.1	33.8	35.0	47.7	34.2	37.0	34.9
All ages (18 years and over)	14.7	8.1	2.9	8.3	9.7	15.2	8.9	8.9	8.8

Source: Australian Immunisation Register (AIR) as at 29 June 2025

\* COVID-19 vaccine coverage uses the Australian Bureau of Statistics June 2023 Estimated Resident Population (ERP) as denominator.

† COVID-19 vaccination uptake and coverage are influenced by changes in COVID-19 vaccine recommendations and eligibility criteria. For this reason, caution should be used when comparing coverage rates in the current 12 month period to previous 12 month periods.

‡ Jurisdiction is based on the state or territory in which a vaccine was administered and may differ from a person's residential. Population denominator data used to calculate COVID-19 vaccine coverage are based on an individual's residential address.

- Nationally, 3.5% of Aboriginal and Torres Strait Islander adults (aged 18 years or over) have received a COVID-19 vaccine in the past six months (Table 6).
- Nationally, less Aboriginal and Torres Strait Islander adults have received a COVID-19 vaccine in the past 12 months (4.6%; Table 6), compared to the 12 months prior (7.1% from 26 June 2023 to 23 June 2024).
  - In the past 12 months, vaccine coverage decreased in all age groups of Aboriginal and Torres Strait Islander people, with the largest decrease seen in ≥ 75 years age group (from 34.2% in the 12 months prior to 26.8% in the past 12 months).
- Among Aboriginal and Torres Strait Islander people there is substantial variation in COVID-19 vaccine coverage across age groups, ranging from 2.7% in adults aged 18–64 years to 26.8% in adults aged 75 years and over. Vaccine coverage increases with increasing age (Table 6).
- Among Aboriginal and Torres Strait Islander people, there is slight variation in vaccine coverage across jurisdictions, ranging from 2.5% in the NT to 9.4% in Tas (Table 6).

**Table 6: COVID-19 vaccine coverage\*†‡ among Aboriginal and Torres Strait Islander populations by age group and jurisdiction, Australia, 24 June 2024 to 29 June 2025**

Age group	ACT	NSW	NT	Qld	SA	Tas	Vic	WA	Total
<b>Past 12 months (24 June 2024 to 29 June 2025)</b>									
18–64 years	6.3	2.7	1.9	2.5	2.8	5.5	4.1	2.2	2.7
65–74 years	34.5	18.4	8.5	16.8	17.7	31.2	20.5	15.1	17.4
≥ 75 years	48.0	28.9	12.6	25.0	28.8	40.8	31.5	26.3	26.8
All ages (18 years and over)	9.3	5.0	2.5	4.2	4.9	9.4	6.7	3.7	4.6
<b>Past 6 months (30 December 2024 to 29 June 2025)</b>									
18–64 years	4.7	1.9	1.1	1.8	1.9	4.1	2.8	1.5	1.9
65–74 years	27.1	14.8	5.6	13.2	13.7	25.5	16.4	12.3	13.8
≥ 75 years	38.0	23.1	9.7	19.8	24.1	33.9	25.5	20.7	21.4
All ages (18 years and over)	7.1	3.8	1.6	3.2	3.6	7.4	5.0	2.7	3.5

Source: Australian Immunisation Register (AIR) as at 29 June 2025

\* COVID-19 vaccine coverage uses the AIR population as the denominator.

† COVID-19 vaccination uptake and coverage are influenced by changes in COVID-19 vaccine recommendations and eligibility criteria. For this reason, caution should be used when comparing coverage rates in the 12 month period to previous 12 month periods.

‡ Jurisdiction is based on the state or territory in which a vaccine was administered and may differ from a person's residential. Population denominator data used to calculate COVID-19 vaccine coverage are based on an individual's residential address.



- Influenza virus strains change year to year, so annual vaccination before the peak of the influenza season provides Australians with the best protection against influenza and its complications. The seasonal influenza vaccine is recommended for everyone aged six months and over.
  - More information on influenza vaccines in Australia is available via the [department's influenza vaccine webpages](#) or from [NCIRS](#).
- Nationally, influenza vaccine coverage is 26.4% for the 2025 seasonal campaign so far (Table 7); however, influenza vaccine coverage this year to date remains lower than vaccine coverage at the same time in the last two years.
- There is substantial variation in influenza vaccine coverage across age groups, ranging from 12.9% in children aged 5–14 years to 50.7% in adults aged 65 years and over (Table 7).
  - The current trend should be interpreted with care as people aged 5–64 years are generally not eligible for free seasonal influenza vaccines under the National Immunisation Program.
- There is some variation in influenza vaccine coverage across jurisdictions, ranging from 19.8% in the NT to 35.7% in the ACT (Table 7).
- Among Aboriginal and Torres Strait Islander populations, there is substantial variation in influenza vaccine coverage across age groups, ranging from 9.6% in children aged 5–14 years to 49.6% in adults aged 65 years and over (Table 7).
- Among Aboriginal and Torres Strait Islander populations, there is some variation in influenza vaccine coverage across jurisdictions, ranging from 15.6% in WA to 26.6% in the ACT (Table 7).

**Table 7: Influenza vaccine coverage\*†‡ by age group and jurisdiction, Australia, 1 March to 29 June 2025**

	ACT	NSW	NT	Qld	SA	Tas	Vic	WA	Total
<b>Age groups</b>									
6 months to <5 years	42.5	22.1	25.5	17.4	23.2	24.9	26.1	19.3	22.2
5–14 years	22.0	12.3	9.2	11.8	13.7	13.3	14.5	13.7	12.9
15–49 years	28.9	17.8	17.4	16.6	21.6	21.1	21.2	17.2	18.4
50–64 years	41.4	28.2	22.8	29.1	33.4	36.2	31.9	28.7	29.2
≥ 65 years	57.3	50.2	31.7	52.6	56.9	58.1	52.9	52.0	50.7
All ages (6 months and over)	35.7	26.0	19.8	25.4	31.1	32.3	28.8	25.4	26.4
<b>Aboriginal and Torres Strait Islander populations</b>									
6 months to <5 years	26.5	14.4	23.1	10.6	13.5	17.8	16.9	12.4	13.8
5–14 years	15.4	9.7	14.3	8.5	10.2	10.8	10.8	9.1	9.6
15–49 years	22.8	14.8	21.4	13.2	16.4	17.1	16.9	12.3	14.8
50–64 years	40.7	32.6	33.9	30.2	33.5	40.5	32.6	27.3	31.1
≥ 65 years	61.3	52.9	38.4	51.0	53.4	58.3	54.5	44.8	49.6
All ages (6 months and over)	26.6	19.2	23.3	16.7	19.9	23.0	21.3	15.6	18.4

Source: Australian Immunisation Register (AIR) as at 29 June 2025

\* Influenza vaccine coverage uses the AIR population as the denominator. Coverage data in these tables may differ slightly from coverage estimates in other reports due to differences in calculation methodologies and/or different data download dates.

† Age is calculated based on the person's age as at 1 July of the reporting year.

‡ Jurisdiction is based on the state or territory in which a vaccine was administered and may differ from a person's residential. Population denominator data used to calculate influenza vaccine coverage are based on an individual's residential address as recorded on Medicare.

- Infants can be protected against severe RSV through the vaccination of pregnant people or the direct administration of monoclonal antibodies like nirsevimab.
- On 3 February 2025, the National RSV Maternal and Infant Protection Program commenced with the roll-out of the National Immunisation Program funded maternal RSV vaccine. The other component of the program is the state and territory nirsevimab (Beyfortus) for infants program. This comprehensive program provides multiple opportunities for infants and young children to be protected.
  - More information on RSV immunisation in Australia is available via the [department's RSV vaccine webpages](#) or from [NCIRS](#).
- While high maternal vaccine uptake is a positive indicator of maternal program success, it may result in lower nirsevimab uptake rates in infants. This is because maternal antibodies passed to the infant can provide protection against RSV, potentially reducing the need for infant immunisation.
  - Maternal RSV vaccine data will be included in future reports. Until maternal RSV vaccine data is included in this report, please refer to the [Respiratory Syncytial Virus Mother and Infant Protection Program \(RSV-MIPP\) data](#) published by NCIRS.
- Nationally, 19.7% of infants (aged < 8 months) have received nirsevimab (Table 8).
- There is substantial variation in nirsevimab uptake in infants across jurisdictions, ranging from 11% in NSW to 33.1% in WA (Table 8).
  - The current trend is likely due to variation in the seasonality and eligibility criteria between state and territory programs, as well as the presence of previous nirsevimab programs. Some state and territory programs are seasonal (from 1 April to 30 September), whereas others are year-round. In states with seasonal programs (SA, Tas, Vic, and parts of WA), uptake may appear disproportionately lower at this time of the year. In addition, Qld and WA had nirsevimab programs in 2024, which may contribute to higher nirsevimab uptake in 2025 in these states.

**Table 8: Nirsevimab (Beyfortus) uptake<sup>\*†‡</sup> by age group and jurisdiction, Australia, 3 February to 29 June 2025**

	ACT	NSW	NT	Qld	SA	Tas	Vic	WA	Total
<b>Age group</b>									
Infants (aged < 8 months)	15.1	11.0	27.4	29.7	25.4	28.4	15.2	33.1	19.7
Young children (aged ≥ 8 to 24 months)	1.0	0.4	0.3	0.1	0.9	1.1	0.6	0.7	0.5

Source: Australian Immunisation Register (AIR) as at 29 June 2025

\* Reporting of RSV monoclonal antibodies to the AIR is not compulsory; therefore, uptake estimates are likely to be underestimated. Nirsevimab is recommended for infants whose mother did not receive the RSV vaccine during pregnancy and therefore nirsevimab uptake should be interpreted with care.

† For infants and young children vaccinated, age in months is calculate as months between the immunisation encounter and date of birth rounded down. For the infant and young children population, age in months is calculated as months between the AIR data extract date and date of birth rounded down.

‡ Jurisdiction is based on the state or territory in which a vaccine was administered and may differ from a person's residential. Population denominator data used to calculate nirsevimab uptake are based on an individual's residential address as recorded on Medicare.

## Vaccine effectiveness

- It is too early to assess vaccine effectiveness for the 2025 influenza season.

## Vaccine match

- Refer to the [Technical Supplement](#) for information on the 2025 southern hemisphere influenza vaccines composition.
- In the year to date, 99.2% (1,673/1,686) of influenza A(H1N1) isolates, 99.6% (251/252) of influenza A(H3N2) isolates and 99.7% (310/311) of influenza B/Victoria lineage isolates characterised have been antigenically similar to the corresponding 2025 vaccine components.