

Community and Hospital Surveillance

ILI, SARI, Influenza and Respiratory Pathogens

2016 Influenza Season, Week 19, ending 15 May 2016

SUMMARY

During week 19 (9–15 May 2016), influenza activity was very low among those consultation-seeking patients nationwide. Influenza activity was also very low among those hospitalised patients in Auckland and Counties Manukau District Health Boards.

- **Influenza-like illness (ILI) and severe acute respiratory illness (SARI) surveillance**

ILI surveillance: Thirty-seven patients with influenza-like illness consulted sentinel general practices in 20 DHBs. The weekly ILI incidence was 8.7 per 100 000 patient population (Figure 1), below the seasonal threshold of ILI consultations. The ILI related influenza incidence was 0.4 per 100 000 patient population.

SARI surveillance: There were 2621 acute admissions to ADHB and CMDHB hospitals this week. Of the 111 patients with suspected respiratory infections, 44 (39.6%) patients met the SARI case definition. Three SARI cases have been admitted to ICU and one SARI related death was reported. The weekly SARI incidence was 3.1 per 100 000 population. The SARI related influenza incidence was 0.0 per 100 000 population.

- **Respiratory pathogen surveillance**

Influenza virus: During this week, 21 ILI specimens were tested, one was positive for influenza viruses. In addition, 16 SARI specimens were tested, none were positive for influenza viruses. For details, see Table 3.

Non-influenza respiratory viruses: For cumulative totals, see Table 4.

The surveillance for community-based influenza-like illness (ILI) and hospital-based severe acute respiratory illness (SARI) provides evidence to inform public health and clinical practice to reduce the impact of influenza virus infection and other important respiratory pathogens. This weekly report summarises data obtained from the ILI and SARI surveillance platforms. The report includes incidence, demographic characteristics, clinical outcomes and aetiologies for community ILI cases as well as hospital SARI cases including ICU admissions and deaths for the past week as well as the cumulative period since 2 May 2016.

Note: Data in this report are provisional and may change as more cases are assessed and information is updated. Data were extracted on 19 May 2016.

INFLUENZA-LIKE ILLNESS and SEVERE ACUTE RESPIRATORY ILLNESS

Influenza-like illness (ILI)

During week 19, ending 15 May 2016, 37 patients with influenza-like illness consulted sentinel general practices in 20 DHBs. The weekly ILI incidence was 8.7 per 100 000 patient population. Of the 21 tested ILI cases, one was positive for influenza viruses. This gives an ILI related influenza incidence of 0.4 per 100 000 patient population (Table 1).

Figure 1. Weekly resident ILI and influenza incidence since 2 May 2016

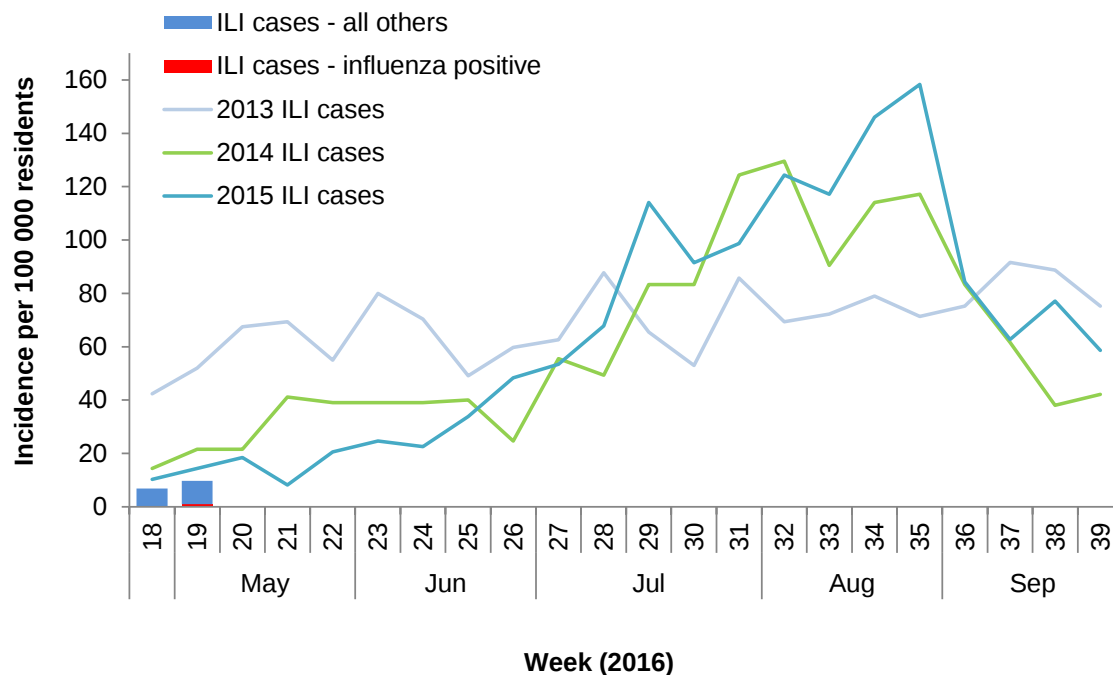


Figure 2. Comparison of 2016 rate with average seasonal rate, and historical thresholds

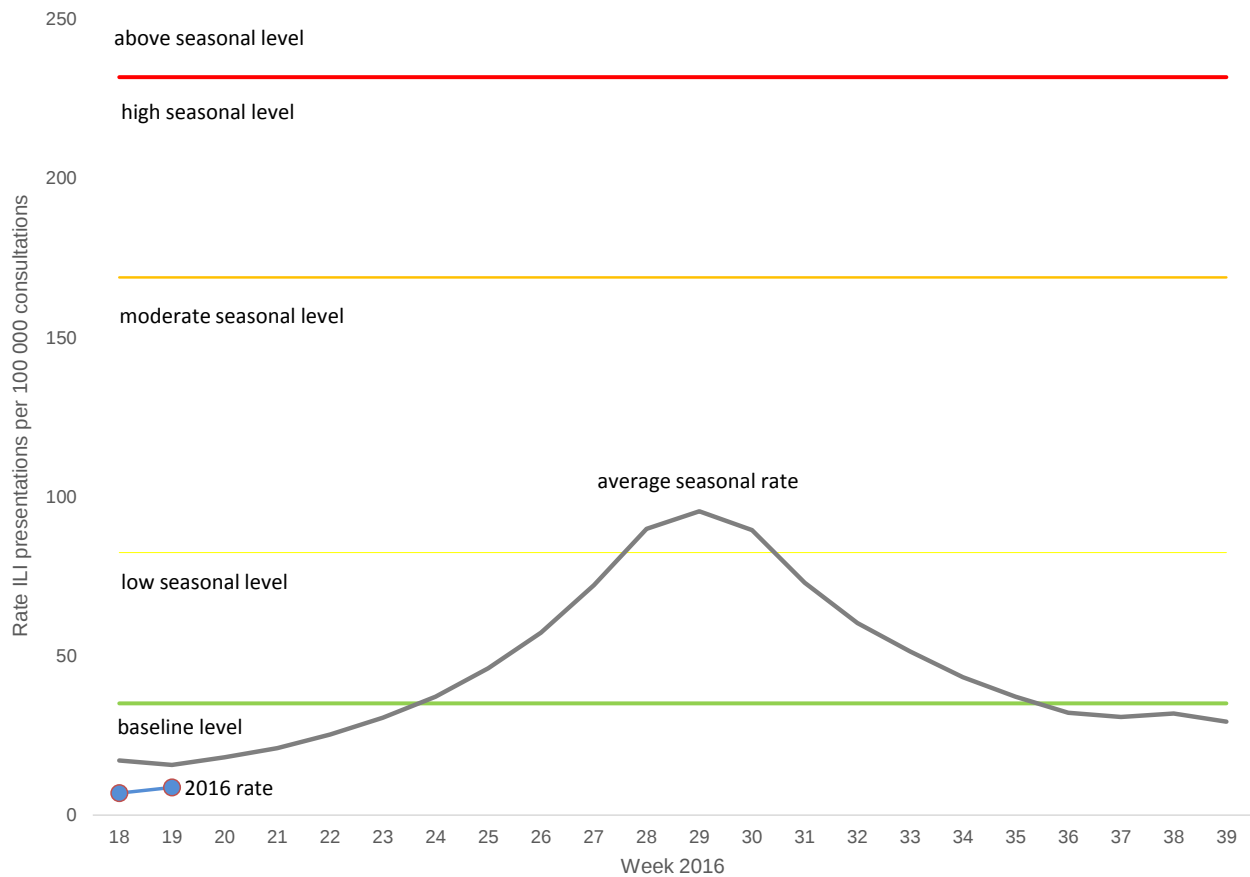
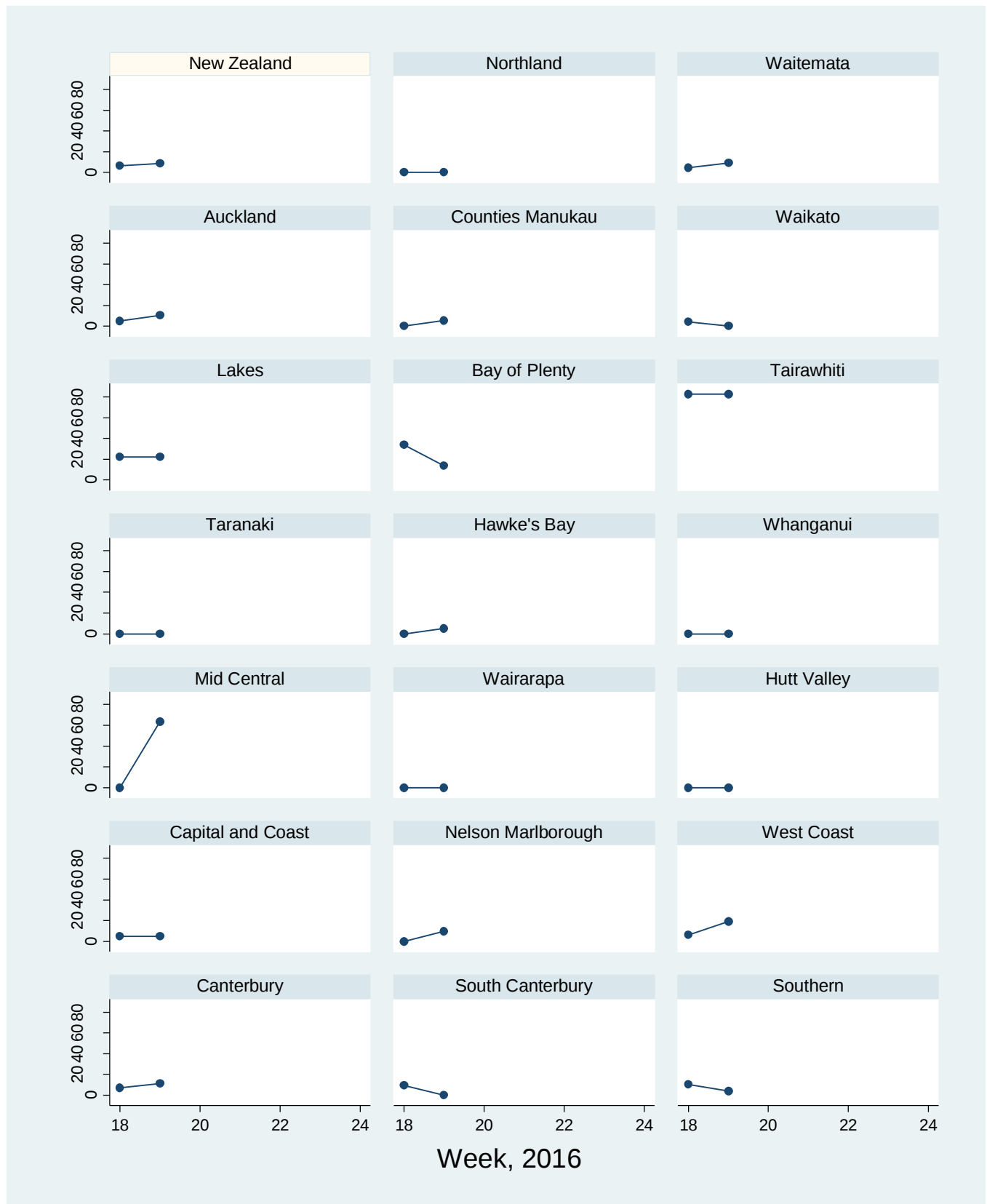


Figure 3 compares the consultation rates for influenza-like illness for each DHB over the past week. Tairāwhiti (82.9 per 100 000, 5 cases), MidCentral (63.6 per 100 000, 5 cases) and Lakes (22.3 per 100 000, 1 case) DHBs had the highest consultation rates.

Figure 3. Rate of ILI consultations by DHB per week since 2 May 2016



Since 2 May 2016, a total of 65 ILI cases were identified. This gives a cumulative ILI incidence of 15.3 per 100 000 patient population (Table 1). Among the 43 tested ILI cases, one (2.3%) was positive for influenza viruses. This gives an ILI related influenza incidence of 0.4 per 100 000 patient population.

Table 1. Demographic characteristics of ILI and influenza cases, since 2 May 2016

Characteristics	ILI & influenza cases among sentinel practices				
	ILI cases	Influenza cases	Prop Influenza positive ¹ (%)	ILI incidence (per 100 000)	Influenza incidence ² (per 100 000)
Overall	65	1	2.3 (100.0)	15.3	0.4
Age group (years)					
<1	0	0	(0.0)	0.0	
1 to 4	0	0	(0.0)	0.0	
5 to 19	11	0	0.0 (0.0)	12.6	0.0
20 to 34	22	0	0.0 (0.0)	24.5	0.0
35 to 49	14	1	12.5 (100.0)	16.8	2.1
50 to 64	8	0	0.0 (0.0)	10.3	0.0
65 to 79	8	0	0.0 (0.0)	17.6	0.0
80 and over	2	0	0.0 (0.0)	13.0	0.0
Unknown	0	0	0.0		
Ethnicity					
Māori	10	0	0.0 (0.0)	18.4	0.0
Pacific peoples	1	0	0.0 (0.0)	3.4	0.0
Asian	1	0	0.0	3.3	
European and Other	53	1	2.9 (100.0)	17.1	0.5
Unknown	0	0	0.0	0.0	
Sex					
Female	39	1	4.0 (100.0)	17.6	0.7
Male	26	0	0.0 (0.0)	12.8	0.0
Unknown	0	0	0.0		

¹Proportion of cases tested which were positive for influenza viruses

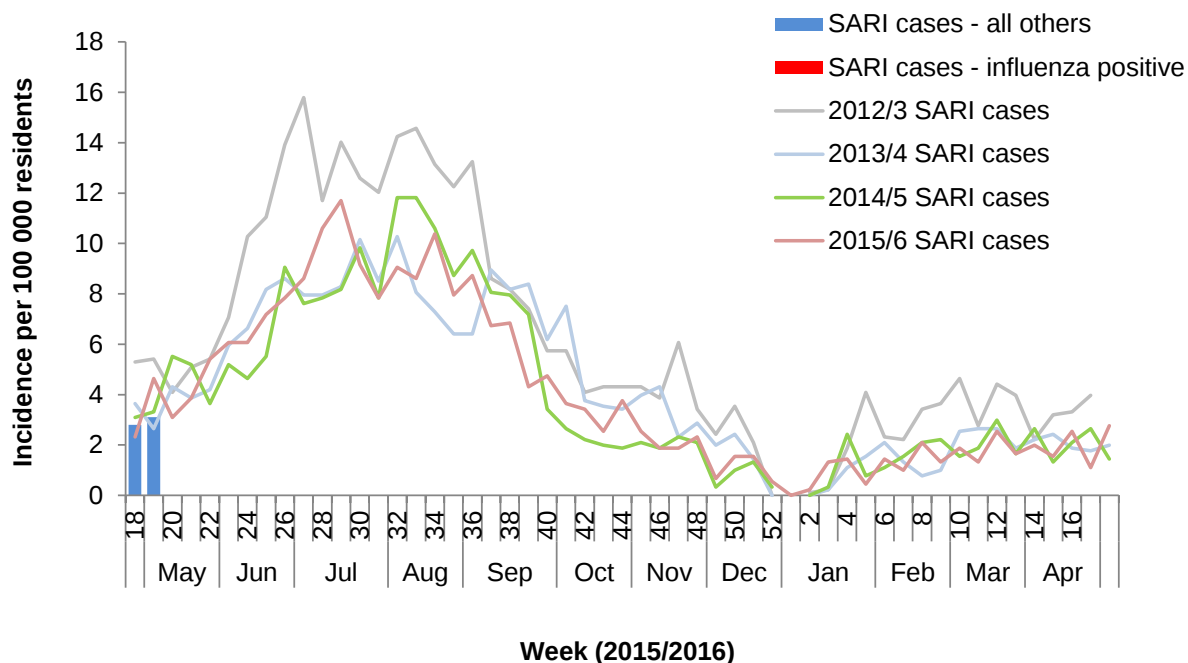
²Adjusted to positivity of tested cases

Severe acute respiratory illness (SARI)

There were 2621 acute admissions to ADHB and CMDHB hospitals during week 19, ending 15 May 2016. A total of 111 patients with suspected respiratory infections were assessed in these hospitals. Of these, 44 (39.6%) patients met the SARI case definition. Three SARI cases have been admitted to ICU and one SARI related death was reported this week.

Of the 44 SARI cases this week, 28 were residents of ADHB and CMDHB. This gives a weekly SARI incidence of 3.1 per 100 000 population (Figure 4). Fifteen SARI residents had specimens tested for influenza viruses, none were positive for influenza viruses. This gives a SARI related influenza incidence of 0.0 per 100 000 patient population (Table 2).

Figure 4. Weekly resident SARI and influenza incidence since 2 May 2016 and previous seasons SARI incidence



Since 2 May 2016, a total of 79 SARI cases were identified. This gives a SARI proportion of 14.8 per 1000 acute hospitalisations (Table 2). Five SARI cases have been admitted to ICU and two SARI related deaths were reported during this period.

Of the 79 SARI cases, 53 were ADHB and CMDHB residents, giving a SARI incidence of 5.9 per 100 000 population (Table 2). Among the 32 tested SARI cases who were ADHB and CMDHB residents, none (0.0%) had positive influenza virus results. This gives a SARI related influenza incidence of 0.0 per 100 000 population.

Table 2. Demographic characteristics of SARI cases and related influenza cases, since 2 May 2016

Characteristics	Admissions	Assessed	SARI & influenza cases among all hospital patients			SARI & influenza cases among ADHB & CMDHB residents			
			SARI Cases (%)	Cases per 1000 hospitalisations	Influenza positive ¹ (%)	SARI cases	SARI incidence (per 100 000)	Influenza Cases	Influenza incidence (per 100 000)
Overall	5327	219	79 (36.1)	14.8	0 (0.0)	53	5.9	0	0.0
Age group (years)									
<1	156		11	70.5	0 (0.0)	9	66.6	0	0.0
1 to 4	365		20	54.8	0 (0.0)	20	37.8	0	0.0
5 to 19	642		5	7.8	0 (0.0)	4	2.1	0	0.0
20 to 34	1079		2	1.9	0 (0.0)	2	1.0	0	0.0
35 to 49	798		6	7.5	0 (0.0)	6	3.1	0	0.0
50 to 64	960		7	7.3	0 (0.0)	6	4.0	0	0.0
65 to 79	834		4	4.8	0 (0.0)	4	5.5	0	0.0
80 and over	493		2	4.1	0 (0.0)	2	8.5	0	0.0
Unknown	0		22			0		0	
Ethnicity									
Māori	753		10	13.3	0 (0.0)	10	10.1	0	0.0
Pacific peoples	1119		28	25.0	0 (0.0)	25	18.1	0	0.0
Asian	866		7	8.1	0 (0.0)	6	2.9	0	0.0
European and Other	2564		12	4.7	0 (0.0)	12	3.0	0	0.0
Unknown	23		22	956.5		0		0	
Hospitals									
ADHB	3149	105	39 (37.1)	12.4	0 (0.0)	20	4.6	0	0.0
CMDHB	2178	114	40 (35.1)	18.4	0 (0.0)	33	7.0	0	0.0
Sex									
Female	2840		27	9.5	0 (0.0)	25	5.4	0	0.0
Male	2487		30	12.1	0 (0.0)	28	6.4	0	0.0
Unknown	0		22			0		0	

¹Proportion of cases tested which were positive for influenza viruses

RESPIRATORY PATHOGEN SURVEILLANCE

Influenza virus

During week 19, 21 ILI specimens were tested; one was positive for influenza viruses. In addition, 16 SARI specimens were tested; none were positive for influenza viruses.

Since 2 May 2016, 43 ILI specimens were tested, one (2.3%) was positive for influenza with the following viruses. In addition, 36 SARI specimens were tested, none were positive for influenza viruses (see Table 3).

Table 3. Influenza viruses among ILI and SARI cases since 2 May 2016

<i>Influenza viruses</i>	ILI	SARI		
	Cases	Cases	ICU	Deaths
No. of specimens tested	43	36	3	0
No. of positive specimens (%) ¹	1 (2.3)	0 (0.0)	0 (0.0)	(-)
Influenza A	0	0	0	0
A (not subtyped)	0	0	0	0
A (H1N1)pdm09	0	0	0	0
A(H1N1)pdm09 by PCR	0	0	0	0
*A/California/7/2009 (H1N1) - like	0	0	0	0
A(H3N2)	0	0	0	0
A(H3N2) by PCR	0	0	0	0
‡A/Switzerland/9715293/2013 (H3N2) - like	0	0	0	0
Influenza B	1	0	0	0
B (lineage not determined)	0	0	0	0
B/Yamagata lineage	1	0	0	0
B/Yamagata lineage by PCR	1	0	0	0
¥B/Phuket/3073/2013 - like	0	0	0	0
B/Victoria lineage	0	0	0	0
B/Victoria lineage by PCR	0	0	0	0
B/Brisbane/60/2008 - like	0	0	0	0
Influenza and non-influenza co-detection (% +ve)	0 (0.0)	(-)	(-)	(-)

¹Number of specimens positive for at least one of the listed viruses; note a specimen may be positive for more than one virus

The recommended influenza vaccine formulation for New Zealand in 2016 is:

A(H1N1) - an A/California/7/2009 (H1N1)pdm09-like strain*

A(H3N2) - an A/Hong Kong/4801/2014 (H3N2)-like strain

B - a B/Brisbane/60/2008-like strain (belonging to B/Victoria lineage)

Note: A/California/7/2009 (H1N1)-like strain is an influenza A(H1N1)pdm09 strain.

* This virus was the A(H3N2) vaccine component for NZ in 2015.

¥ This virus was the B vaccine component for NZ in 2015.

Non-influenza respiratory pathogens

Since 2 May 2016, thirty-six ILI cases were tested for non-influenza viruses, 20 (55.6%) were positive with the following viruses. Nine SARI specimens were tested for non-influenza viruses, six (66.7%) were positive with the following viruses (see Table 4).

Table 4. Non-influenza viruses among ILI and SARI cases since 2 May 2016

<i>Non-influenza respiratory viruses</i>	ILI	SARI		
	Cases	Cases	ICU	Deaths
No. of specimens tested	36	9	0	0
No. of positive specimens (%) ¹	20 (55.6)	6 (66.7)	(-)	(-)
Respiratory syncytial virus (RSV)	5	2	0	0
Parainfluenza 1 (PIV1)	2	2	0	0
Parainfluenza 2 (PIV2)	0	0	0	0
Parainfluenza 3 (PIV3)	0	0	0	0
Rhinovirus (RV)	12	1	0	0
Adenovirus (AdV)	0	2	0	0
Human metapneumovirus (hMPV)	1	0	0	0
Enterovirus	0	0	0	0
Single virus detection (% of positives)	20 (100.0)	5 (83.3)	(-)	(-)
Multiple virus detection (% of positives)	0 (0.0)	1 (16.7)	(-)	(-)

APPENDIX

Recent global experience with pandemic influenza A(H1N1)pdm09 highlights the importance of monitoring severe and mild respiratory disease to support pandemic preparedness as well as seasonal influenza prevention and control. Two active, prospective, population-based surveillance systems were used to monitor influenza and other respiratory pathogens: 1) among those registered patients seeking consultations with influenza-like illness (ILI) at sentinel general practices nation-wide; 2) among those hospitalized patients with severe acute respiratory illness (SARI) in Auckland and Counties Manukau District Health Boards (ADHB and CMDHB).

The aims of ILI and SARI surveillance are: 1) to measure the burden of severe and moderate disease caused by influenza and other respiratory pathogens; 2) to monitor trends in severe and moderate disease caused by influenza and other respiratory pathogens; 3) to identify high risk groups that should be prioritized for prevention and treatment; 4) to monitor antigenic, genetic and antiviral characteristics of influenza viruses associated with severe and mild disease. 5) to provide a study base to estimate the effectiveness of influenza vaccine.

ACKNOWLEDGEMENT

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DESCRIPTION OF ILI ACTIVITY THRESHOLDS

The values for the different intensity levels for 2016 are listed in the table below. This is based on New Zealand's consultation rates from 2000–2015 (excluding the pandemic year, 2009) and WHO's interim guidance severity assessment

Below seasonal level (baseline, per 100,000)	Seasonal level (per 100,000)			Above seasonal level (per 100,000)
	low	moderate	high	
<35.1	35.1-82.5	82.5-168.9	168.9-231.8	>231.8

- The baseline threshold indicates the level of influenza activity that signals the start and end of the annual influenza season and it is based on the Moving Epidemic Method (MEM) (Vega *et al. Influenza and other respiratory viruses* 2013;7(4):546-558).
- Seasonal levels (low, moderate and high) are estimated as the upper limits of the 40%, 90% and 97.5% one-sided confidence intervals of the geometric mean of 30 highest epidemic weekly rates using the MEM method. As many other countries use this method, it allows the NZ data to be interpreted not just at the country level but also comparable with other countries.
- The average seasonal curve indicates the usual seasonal activity that may occur during a typical year using the method described in "Global epidemiological surveillance standards for influenza" (http://www.who.int/influenza/resources/documents/WHO_Epidemiological_Influenza_Surveillance_Standards_2014.pdf).

NOTES ON INTERPRETATION

- SARI case definition: “An acute respiratory illness with a history of fever or measured fever of $\geq 38^{\circ}\text{C}$, AND cough, AND onset within the past 10 days, AND requiring inpatient hospitalisation (defined as a patient who is admitted under a medical team and to a hospital ward or assessment unit).
- ILI case definition: “An acute respiratory illness with a history of fever or measured fever of $\geq 38^{\circ}\text{C}$, AND cough, AND onset within the past 10 days, AND requiring GP consultation”.
- ILI sentinel general practices: a total of 82 sentinel general practices have agreed to participate in community ILI surveillance. These practices have ~400 000 registered patients, covering roughly 9% of the NZ population.
- SARI sentinel hospitals serving a population of 906 000 people: Auckland City Hospital and the associated Starship Children’s Hospital (ADHB), and Middlemore Hospital and the associated Kidz First Children’s Hospital (CMDHB).
- The real-time PCR assay for influenza virus uses CDC’s protocol (http://www.accessdata.fda.gov/cdrh_docs/pdf8/k080570.pdf);
- The real-time PCR assay for non-influenza respiratory viruses (respiratory syncytial virus, parainfluenza virus types 1-3, human metapneumovirus, rhinovirus and adenovirus) uses CDC’s protocol. Note: The rhinovirus PCR detects mostly rhinovirus with slight cross-reactivity against enterovirus.
- The surveillance week is Monday to Sunday inclusive, and data are extracted on the subsequent Tuesday. Results from previous weeks will be revised as data are updated (laboratory test results in particular may be delayed).

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