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## **Community and Hospital Surveillance**

ILI, SARI, Influenza and Respiratory Pathogens

#### 2017 Influenza Season, Week 38, ending 24 September 2017

### SUMMARY

- Influenza-like illness (ILI) consultation rates decreased compared to the previous week, and were below the seasonal threshold level.
- Influenza associated severe acute respiratory illness (SARI) hospitalisation rates were similar to last week, at a low level and below the seasonal threshold level.
- Influenza A(H3N2) are the predominant viruses in New Zealand this year.
- Influenza B/Victoria lineage viruses also co-circulated with B/Yamagata lineage viruses with more B/Yamagata viruses this year.
- Rhinovirus and respiratory syncytial virus (RSV) are the most commonly detected noninfluenza viruses this week, with numbers remaining high.

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 January 2017.

Note: Data in this report are provisional and may change as more cases are assessed and information is updated. Data were extracted on 27 September 2017.

## **INFLUENZA-LIKE ILLNESS and SEVERE ACUTE RESPIRATORY ILLNESS**

#### Influenza-like illness (ILI)

During week 38, ending 24 September 2017, 101 patients with influenza-like illness consulted sentinel general practices in 20 DHBs. The weekly ILI incidence was 24.6 per 100 000 patient population (Figures 1 and 2). Of the 58 tested ILI cases, 14 were positive for influenza viruses. This gives an ILI related influenza incidence (adjusted) of 5.9 per 100 000 patient population.

#### Figure 1. Weekly resident ILI and influenza incidence since 2 January 2017



Week 2017







The weekly consultation rates for influenza-like illness by different age groups and ethnicities are shown in Figures 3 and 4.



Figure 3. Weekly ILI incidence by age group since 2 January 2017

Figure 4. Weekly ILI incidence by ethnicity since 2 January 2017





Figure 5 compares the consultation rates for influenza-like illness for each DHB over the past week. Waitemata (189.8 per 100 000, 24 cases), Auckland (81.0 per 100 000, 46 cases), and Whanganui (60.2 per 100 000, 1 case) DHBs had the highest consultation rates.



Figure 5. Rate of ILI consultations per 100 000 registered by DHB per week since 2 January 2017

\*Results that have some uncertainty, with less than 5% of the DHB population covered (see Notes on Interpretation). Note: Outliers have been omitted from this graph.

ILI consultation rates for any particular DHB should be treated with caution as they may not be representative of the real situation for a particular community or setting, especially if the surveillance system has a small number of participating General Practices in the DHB, or the GP enrolled patient population is small, the calculated ILI rates are subject to greater fluctuation.

Since 2 January 2017, a total of 2813 ILI cases were identified. This gives a cumulative ILI incidence of 684.2 per 100 000 patient population (Table 1). Among the 2262 tested ILI cases, 949 (42.0%) were positive for influenza viruses. This gives an ILI related (adjusted) influenza incidence of 287.0 per 100 000 patient population.

	I	LI & influen	za cases among	sentinel practi	ces
Characteristics	ILI cases	Influenza cases	Prop Influenza positive <sup>1</sup> (%)	ILI incidence (per 100 000)	Influenza incidence <sup>2</sup> (per 100 000)
Overall	2813	949	42.0 (100.0)	684.2	287.0
Age group (years)					
<1	40	3	9.4 (0.3)	605.1	56.7
1-4	189	38	25.9 (4.0)	771.1	199.3
5–19	735	307	51.3 (32.3)	868.7	445.2
20–34	452	137	36.6 (14.4)	526.1	192.7
35–49	655	238	45.4 (25.1)	799.0	362.9
50–64	487	149	38.5 (15.7)	664.1	255.7
65–79	215	59	36.0 (6.2)	526.5	189.4
>80	40	18	51.4 (1.9)	299.5	154.0
Unknown	0	0	0.0		
Ethnicity					
Māori	265	85	41.7 (9.0)	465.6	194.0
Pacific peoples	119	43	47.3 (4.5)	398.0	188.1
Asian	277	127	51.8 (13.4)	924.5	479.2
European and Other	2150	694	40.3 (73.1)	732.1	295.2
Unknown	2	0	0.0	0.0	
Sex					
Female	1602	529	41.4 (55.7)	750.6	310.7
Male	1207	420	42.7 (44.3)	610.5	260.6
Unknown	4	0	0.0		

<sup>1</sup>Proportion of cases tested which were positive for influenza viruses

<sup>2</sup>Adjusted to positivity of tested cases



## Severe acute respiratory illness (SARI)

There were 2920 acute admissions to ADHB and CMDHB hospitals during week 38, ending 24 September 2017. A total of 102 patients with acute respiratory illness were assessed in these hospitals. Of these, 59 (57.8%) patients met the SARI case definition. Seven cases were admitted to ICU and one SARI related death was reported this week. Of the 43 tested residents with SARI, two were positive for influenza viruses, giving a SARI related influenza incidence of 0.3 per 100 000 population this week.

# Figure 6. Weekly resident SARI and influenza incidence since 2 May 2016 and previous seasons SARI incidence



Week 2017

Since 1 May 2017, a total of 1677 SARI cases were identified. This gives a SARI proportion of 28.2 per 1000 acute hospitalisations (Table 2). One hundred and sixteen SARI cases have been admitted to ICU and 28 SARI related deaths were reported during this period.

Of the 1677 SARI cases, 1314 were ADHB and CMDHB residents, giving a SARI incidence of 145.1 per 100 000 population (Table 2). Among the 1197 tested SARI cases who were ADHB and CMDHB residents, 349 (29.2%) had positive influenza virus results. This gives a SARI (adjusted) related influenza incidence of 42.3 per 100 000 population.

			SARI & inf	luenza cases amor	ng all hospital	SARI & i	nfluenza cases a	mong ADHB	& CMDHB
				patients			reside	ents	
Characteristics	Admissions	Assessed	SARI Cases (%)	Cases per 1000 hospitalisations	Influenza positive <sup>1</sup> (%)	SARI cases	SARI incidence (per 100 000)	Influenza Cases	Influenza incidence (per 100 000)
Overall	59545	3053	1677 (54.9)	28.2	372 (29.3)	1314	145.1	349	42.3
Age group (years)									
<1	2381		313	131.5	23 (8.5)	295	2184.2	22	186.3
1–4	4080		201	49.3	33 (20.0)	182	344.2	31	71.1
5–19	7023		93	13.2	21 (27.6)	78	40.5	15	9.5
20–34	11222		84	7.5	30 (37.0)	81	38.9	29	14.4
35–49	8668		93	10.7	32 (36.4)	86	45.0	30	16.7
50–64	10114		192	19.0	82 (44.8)	187	124.2	80	55.8
65–79	9668		248	25.7	82 (34.7)	236	322.9	75	107.6
>80	6389		172	26.9	69 (41.6)	168	717.0	67	296.5
Unknown	0		281			1		0	
Ethnicity									
Māori	8129		286	35.2	52 (20.2)	258	259.4	47	52.3
Pacific peoples	12700		492	38.7	125 (27.8)	478	346.4	118	94.0
Asian	9985		113	11.3	37 (37.4)	109	51.8	36	19.0
European and Other	28320		504	17.8	158 (34.3)	468	116.5	148	40.0
Unknown	411		282	686.1		1		0	
Hospitals									
ADHB	34794	1126	807 (71.7)	23.2	182 (35.0)	519	118.9	164	42.1
CMDHB	24751	1927	870 (45.1)	35.2	190 (25.3)	795	169.4	185	42.7
Sex									
Female	31582		717	22.7	196 (30.1)	670	144.0	186	43.6
Male	27960		677	24.2	174 (28.4)	642	145.8	162	40.7
Unknown	3		283			2		1	

#### Table 2. Demographic characteristics of SARI cases and related influenza cases, since 1 May 2017

<sup>1</sup>Proportion of cases tested which were positive for influenza viruses Note. A specimen may be positive for more than one virus; a patient may have more than one specimen tested.

## **RESPIRATORY PATHOGEN SURVEILLANCE**

#### Influenza virus

During week 38, 58 ILI specimens were tested; 14 were positive for influenza viruses. In addition, 45 SARI specimens were tested; two were positive for influenza viruses.

Since 1 May 2017, 2247 ILI specimens were tested, 948 (42.2%) were positive for influenza with the following viruses. In addition, 1431 SARI specimens were tested, 410 (28.7%) were positive for influenza viruses (see Table 3).

Table 5. Initidenza viruses anong izi and OARI c	<b>Way 2017</b>						
Influenza viruses	ILI	SARI	SARI and non-SARI				
	Cases (%)	Cases (%)	ICU (%)	Deaths (%)			
No. of specimens tested	2247	1431	263	31			
No. of positive specimens (%) <sup>1</sup>	948 (42.2)	410 (28.7)	24 (9.1)	13 (41.9)			
Influenza A	545	299	15	8			
A (not subtyped)	11	114	5	1			
A(H1N1)pdm09	70	40	3	0			
A(H1N1)pdm09 by PCR	50	36	2	0			
A/Michigan/45/2015 (H1N1)pdm09 - like	16	4	1	0			
A/California/7/2009 (H1N1)pdm09 - like	4	0	0	0			
A(H3N2)	464	145	7	7			
A(H3N2) by PCR	413	140	6	7			
A/Hong Kong/4801/2014 (H3N2) - like	51	5	1	0			
Influenza B	403	111	9	5			
B (lineage not determined)	18	54	7	3			
B/Yamagata lineage	370	56	2	2			
B/Yamagata lineage by PCR	182	29	2	2			
B/Phuket/3073/2013 - like	188	27	0	0			
B/Victoria lineage	15	1	0	0			
B/Victoria lineage by PCR	7	1	0	0			
B/Brisbane/60/2008 - like	8	0	0	0			
Influenza and non-influenza co-detection (% +ve)	56 (5.9)	29 (7.1)	3 (12.5)	1 (7.7)			

#### Table 3. Influenza viruses among ILI and SARI cases since 1 May 2017

<sup>1</sup>Number of specimens positive for at least one of the listed viruses

Note. A specimen may be positive for more than one virus; a patient may have more than one specimen tested.

The recommended influenza vaccine formulation for trivalent vaccine for New Zealand in 2017 is:

A(H1N1) an A/Michigan/45/2015 (H1N1)pdm09-like virus

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

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

Quadrivalent vaccines contain the above three viruses plus one more vaccine component: B/Phuket/3073/2013-like virus (belonging to B/Yamagata lineage)

Note: Antigenic characterization of the current A(H3N2) viruses have been technically challenging because many viruses had low or undetectable haemagglutination activity. This phenomenon has been well recognized globally and documented in WHO's Weekly Epidemiological Record: http://apps.who.int/iris/bitstream/10665/254756/1/WER9211.pdf?ua=1



#### Non-influenza respiratory pathogens

Since 1 May 2017, 2186 ILI specimens were tested for non-influenza viruses, 499 (22.8%) were positive with the following viruses. One thousand and two hundred ninety SARI specimens were tested for non-influenza viruses, 477 (37.0%) were positive with the following viruses (see Table 4).

#### Table 4. Non-influenza viruses among ILI and SARI cases since 1 May 2017

Non-influenza respiratory viruses	ILI	SARI	SARI and n	on-SARI
	Cases (%)	Cases (%)	ICU (%)	Deaths (%)
No. of specimens tested	2186	1290	206	32
No. of positive specimens (%) <sup>1</sup>	499 (22.8)	477 (37.0)	134 (65.0)	3 (9.4)
Respiratory syncytial virus (RSV)	130	250	70	2
Parainfluenza 1 (PIV1)	6	1	0	0
Parainfluenza 2 (PIV2)	24	16	4	0
Parainfluenza 3 (PIV3)	63	52	11	0
Rhinovirus (RV)	163	150	59	1
Adenovirus (AdV)	59	54	14	0
Human metapneumovirus (hMPV)	51	38	5	1
Enterovirus	36	31	11	0
Single virus detection (% of positives)	470 (94.2)	378 (79.2)	99 (73.9)	0 (-)
Multiple virus detection (% of positives)	29 (5.8)	99 (20.8)	35 (26.1)	0 (-)

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





Figure 8. Temporal distribution of the number and proportion of influenza viruses from SARI specimens by type and week<sup>1</sup>







Figure 10. Temporal distribution of the number and proportion of non-influenza viruses from SARI specimens by type and week<sup>1</sup>



<sup>1</sup>Figures for recent weeks will be underestimates due to time lag in receiving laboratory test results.



## APPENDIX

#### Table 5. Influenza-like illness count by DHB by week 18–38, 2017

рпв											Week										
ОПВ	18 19 20 21 22 23 24 25 26 27 28								28	29	30	31	32	33	34	35	36	37	38		
Auckland	15	14	15	18	30	25	45	71	58	83	78	62	82	69	58	60	64	47	38	49	46
Bay of Plenty	0	0	0	0	2	1	1	6	7	11	4	4	5	6	7	7	11	9	2	2	7
Canterbury	2	5	10	3	3	1	11	16	9	12	9	20	18	15	22	18	20	5	8	6	10
Capital and Coast	2	2	3	4	11	9	13	6	8	25	18	22	13	7	16	23	7	5	6	6	3
Counties Manukau	1	2	0	2	1	0	3	1	6	3	6	1	2	3	0	1	0	0	0	0	0
Hawke's Bay	0	0	1	3	1	1	3	0	4	4	8	2	3	8	7	3	1	2	3	2	0
Hutt Valley	4	0	0	0	0	1	1	1	0	0	0	0	0	1	0	0	0	0	0	0	0
Lakes	0	0	0	1	0	0	0	0	0	0	3	0	0	0	0	0	0	0	0	0	0
MidCentral	0	0	0	0	0	0	0	1	1	1	0	0	1	1	1	0	1	1	0	0	0
Nelson Marlborough	0	3	2	0	1	0	4	3	2	2	4	9	3	4	7	3	2	0	1	1	1
Northland	2	1	4	2	0	0	1	8	4	4	2	2	0	0	1	0	0	2	0	0	1
South Canterbury	1	3	1	2	0	0	2	1	1	1	0	5	10	8	8	16	23	6	6	9	0
Southern	1	2	0	5	4	5	1	2	4	4	7	11	4	10	11	4	6	5	6	4	7
Tairawhiti	0	0	1	2	4	2	2	7	3	2	2	4	2	2	2	2	1	0	0	0	0
Taranaki	1	0	0	1	0	0	1	3	1	2	1	1	0	0	2	1	0	1	2	0	0
Waikato	0	5	3	3	4	6	4	8	4	13	9	6	5	3	7	4	1	7	2	0	0
Wairarapa	0	0	0	0	0	0	0	0	5	2	1	1	0	0	0	0	0	0	0	0	1
Waitemata	1	6	2	8	22	16	24	44	41	37	30	44	50	52	37	39	24	40	36	26	24
West Coast	1	1	0	0	0	2	0	1	3	2	0	0	0	1	2	0	0	0	1	1	0
Whanganui	0	0	1	4	0	1	2	0	5	5	5	0	6	4	1	1	1	0	0	0	1
New Zealand	31	44	43	58	83	70	118	179	166	213	187	194	204	194	189	182	162	130	111	106	101

#### Table 6. Influenza-like illness rate by DHB by week 18–38, 2017

סחט	Rate per 100 000																				
ИПВ	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38
Auckland	26.4	24.7	26.4	31.7	52.8	44.0	79.2	125.0	102.1	146.2	137.4	109.2	144.4	121.5	102.1	105.7	112.7	82.8	66.9	86.3	81.0
Bay of Plenty	0.0	0.0	0.0	0.0	13.5	6.7	6.7	40.4	47.1	74.1	26.9	26.9	33.7	40.4	47.1	47.1	74.1	60.6	13.5	13.5	47.1
Canterbury	2.8	7.1	14.2	4.3	4.3	1.4	15.6	22.8	12.8	17.1	12.8	28.4	25.6	21.3	31.3	25.6	28.4	7.1	11.4	8.5	14.2
Capital and Coast	7.8	7.8	11.7	15.6	42.8	35.0	50.6	23.4	31.2	97.3	70.1	85.7	50.6	27.3	62.3	89.6	27.3	19.5	23.4	23.4	11.7
Counties Manukau	3.0	6.0	0.0	6.0	3.0	0.0	9.0	3.0	18.0	9.0	18.0	3.0	6.0	9.0	0.0	3.0	0.0	0.0	0.0	0.0	0.0
Hawke's Bay	0.0	0.0	5.2	15.7	5.2	5.2	15.7	0.0	20.9	20.9	41.7	10.4	15.7	41.7	36.5	15.7	5.2	10.4	15.7	10.4	0.0
Hutt Valley	15.2	0.0	0.0	0.0	0.0	3.8	3.8	3.8	0.0	0.0	0.0	0.0	0.0	3.8	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Lakes*	0.0	0.0	0.0	22.3	0.0	0.0	0.0	0.0	0.0	0.0	66.8	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
MidCentral*	0.0	0.0	0.0	0.0	0.0	0.0	0.0	35.5	35.5	35.5	0.0	0.0	35.5	35.5	35.5	0.0	35.5	35.5	0.0	0.0	0.0
Nelson Marlborough	0.0	29.2	19.5	0.0	9.7	0.0	39.0	29.2	19.5	19.5	39.0	87.7	29.2	39.0	68.2	29.2	19.5	0.0	9.7	9.7	9.7
Northland	15.7	7.9	31.5	15.7	0.0	0.0	7.9	62.9	31.5	31.5	15.7	15.7	0.0	0.0	7.9	0.0	0.0	15.7	0.0	0.0	7.9
South Canterbury	9.3	27.8	9.3	18.6	0.0	0.0	18.6	9.3	9.3	9.3	0.0	46.4	92.8	74.2	74.2	148.5	213.4	55.7	55.7	83.5	0.0
Southern	2.2	4.4	0.0	10.9	8.7	10.9	2.2	4.4	8.7	8.7	15.3	24.0	8.7	21.8	24.0	8.7	13.1	10.9	13.1	8.7	15.3
Tairawhiti	0.0	0.0	14.4	28.9	57.7	28.9	28.9	101.0	43.3	28.9	28.9	57.7	28.9	28.9	28.9	28.9	14.4	0.0	0.0	0.0	0.0
Taranaki	4.4	0.0	0.0	4.4	0.0	0.0	4.4	13.1	4.4	8.7	4.4	4.4	0.0	0.0	8.7	4.4	0.0	4.4	8.7	0.0	0.0
Waikato*	0.0	32.5	19.5	19.5	26.0	39.0	26.0	52.1	26.0	84.6	58.6	39.0	32.5	19.5	45.5	26.0	6.5	45.5	13.0	0.0	0.0
Wairarapa	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	200.0	80.0	40.0	40.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	40.0
Waitemata*	7.9	47.5	15.8	63.3	174.0	126.6	189.8	348.0	324.3	292.7	237.3	348.0	395.5	411.3	292.7	308.5	189.8	316.4	284.7	205.6	189.8
West Coast	6.3	6.3	0.0	0.0	0.0	12.7	0.0	6.3	19.0	12.7	0.0	0.0	0.0	6.3	12.7	0.0	0.0	0.0	6.3	6.3	0.0
Whanganui*	0.0	0.0	60.2	241.0	0.0	60.2	120.5	0.0	301.2	301.2	301.2	0.0	361.4	241.0	60.2	60.2	60.2	0.0	0.0	0.0	60.2
New Zealand	7.5	10.7	10.5	14.1	20.2	17.0	28.7	43.5	40.4	51.8	45.5	47.2	49.6	47.2	46.0	44.3	39.4	31.6	27.0	25.8	24.6

\*Results that have some uncertainty, with less than 5% of the DHB population covered (see Notes on Interpretation).



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

We acknowledge the support of the New Zealand Ministry of Health. SARI surveillance was established and funded by the US CDC, and continues to operate through funding from the New Zealand Ministry of Health.

#### DESCRIPTION OF ILI ACTIVITY THRESHOLDS

The values for the different intensity levels for 2017 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	Seas	Above seasonal							
(baseline, per 100,000)	low	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"* (<u>http://www.who.int/influenza/resources/documents/WHO\_Epidemiological\_Influenza\_Surveillance\_Standards\_2014.pdf</u>).



#### NOTES ON INTERPRETATION

- SARI case definition: "An acute respiratory illness with a history of fever or measured fever of ≥38°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)". A non-SARI case is a hospitalised respiratory patient who does not meet the SARI case definition.
- ILI case definition: "An acute respiratory illness with a history of fever or measured fever of ≥38°C, AND cough, AND onset within the past 10 days, AND requiring GP consultation".
- ILI sentinel general practices: a total of 74 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 (<u>http://www.accessdata.fda.gov/cdrh\_docs/pdf8/k080570.pdf</u>.);
- 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).
- ILI consultation rates for any particular DHB should be treated with caution. If the surveillance system has a small number of participating General Practices in the DHB, or the GP enrolled patient population is small, the calculated ILI rates are subject to greater fluctuation.

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