

# British Columbia Influenza Surveillance Bulletin

Influenza Season 2013-14, Number 17, Weeks 15-16

April 6 to 19, 2014

## Table of Contents:

### British Columbia:

Sentinel Physicians	<a href="#">Page 2</a>
Children's Hospital ER	<a href="#">Page 2</a>
Medical Services Plan	<a href="#">Page 3</a>
Laboratory Surveillance	<a href="#">Page 5</a>
ILI Outbreaks	<a href="#">Page 7</a>

### Canada:

FluWatch Activity levels	<a href="#">Page 8</a>
NML Strain Characterization	<a href="#">Page 8</a>
NML Antiviral Resistance	<a href="#">Page 8</a>

### International:

USA and WHO	<a href="#">Page 9</a>
Avian Influenza A(H7N9)	<a href="#">Page 9</a>
MERS-CoV	<a href="#">Page 9</a>
WHO 2013-14 Recommended Vaccine Components	<a href="#">Page 10</a>
WHO 2014-15 Recommended Vaccine Components	<a href="#">Page 10</a>

### Additional Information:

List of Acronyms	<a href="#">Page 11</a>
Web Sites	<a href="#">Page 11</a>
Outbreak Report Form	<a href="#">Page 12</a>

## Ongoing influenza B activity in BC

In weeks 15-16 (April 6 to 19, 2014), influenza activity increased slightly in BC, driven by ongoing influenza B circulation during this period.

At the BC provincial laboratory, the overall influenza positivity rate increased from  $\leq 15\%$  in weeks 10-14 to 17% in week 15 and 22% in week 16. Influenza B viruses comprised  $>85\%$  of all influenza positive specimens in weeks 15-16.

Consultation rates among patients presenting to sentinel physicians or the BC Children's and Women's Hospital ER for influenza-like illness increased slightly from week 15 to 16, while MSP service claims for influenza illness remained at low levels throughout the province.

Two lab-confirmed influenza B school outbreaks were reported during this period from Northern Health Authority: one in week 15 and one in week 16.

Prepared by BCCDC Influenza & Emerging Respiratory Pathogens Team  
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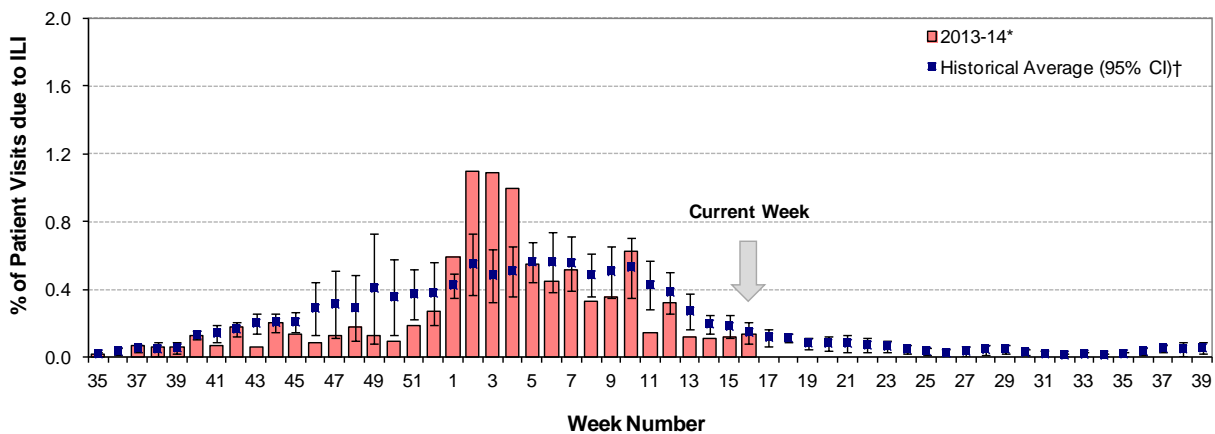
Report Disseminated: April 24, 2014

## British Columbia

### Sentinel Physicians

The proportion of patients with influenza-like illness (ILI) among those presenting to sentinel physicians has remained relatively stable in recent weeks. Rates increased slightly from 0.12% in week 15 to 0.13% in week 16 but remained within expected ranges for this time of year. For weeks 15-16, 46-59% of sentinel sites have reported data.

**Percent of patient visits to sentinel physicians due to influenza-like illness (ILI) compared to historical average, British Columbia, 2013-14**



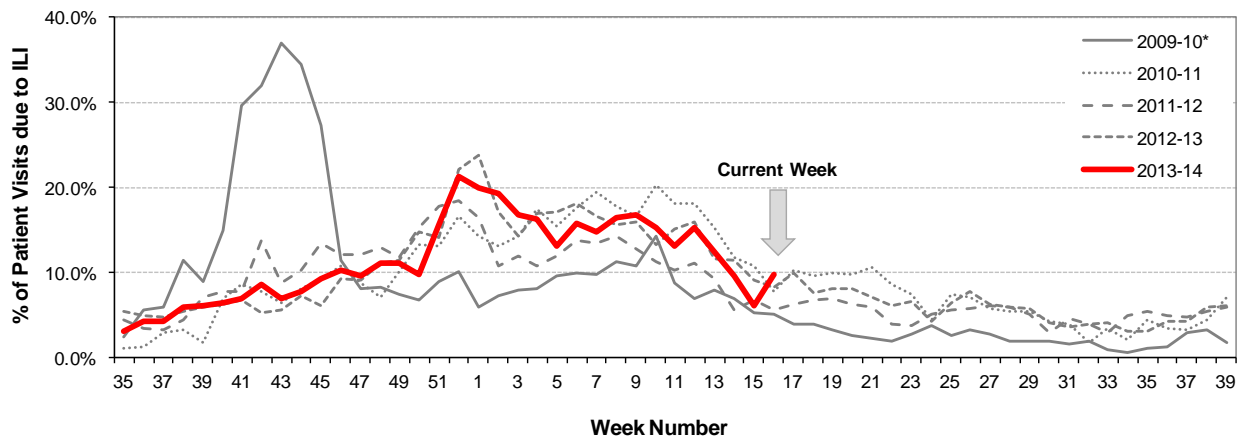
\* Data are subject to change as reporting becomes more complete.

† Historical average based on 2001-02 to 2012-13 seasons, excluding 2008-09 and 2009-10 due to atypical seasonality; CI=confidence interval.

### BC Children's Hospital Emergency Room

The proportion of visits to BC Children's Hospital Emergency Room (ER) attributed to ILI increased slightly from 6% in week 15 to 10% in week 16 but remained comparable to rates seen in previous seasons for this time of year.

**Percent of patients presenting to BC Children's Hospital ER with triage chief complaint of "flu," or "influenza" or "fever/cough," British Columbia, 2013-14**



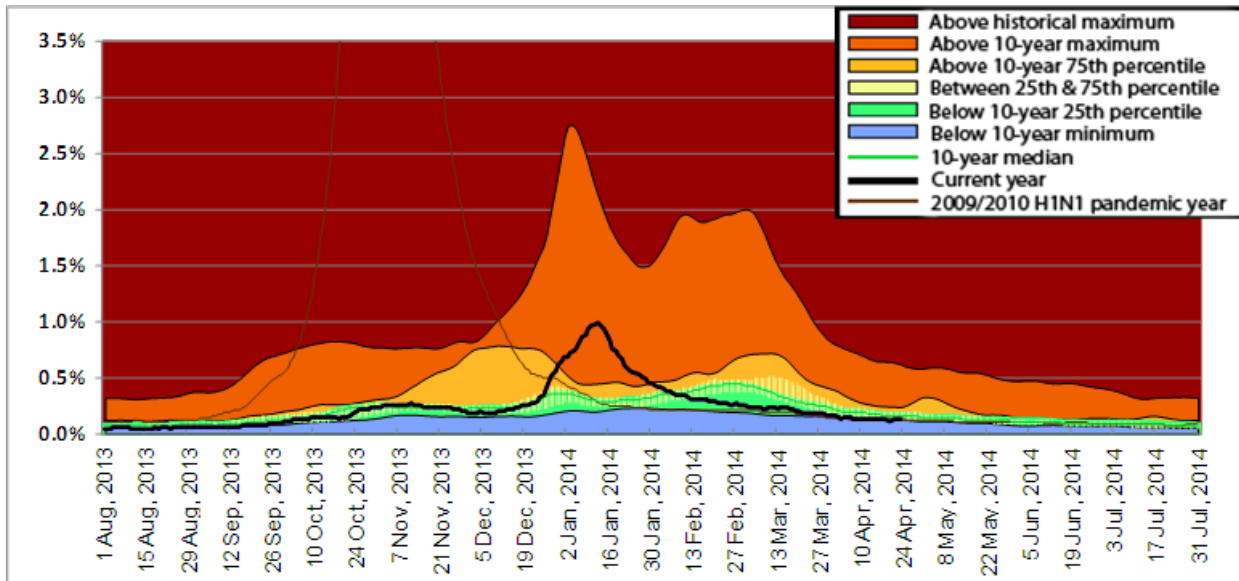
Source: BCCH Admitting, discharge, transfer database, ADT

\* Data from 2010-11 to 2013-14 is based on new system (Triage Chief Complaint) not directly comparable to data for 2009-10. In bulletins before week 9 of 2011-12 season, data is based on old system.

**Medical Services Plan**

In weeks 15-16, BC Medical Services Plan (MSP) general practitioner claims for influenza illness (II), as a proportion of all submitted MSP claims, remained at low levels throughout the province.

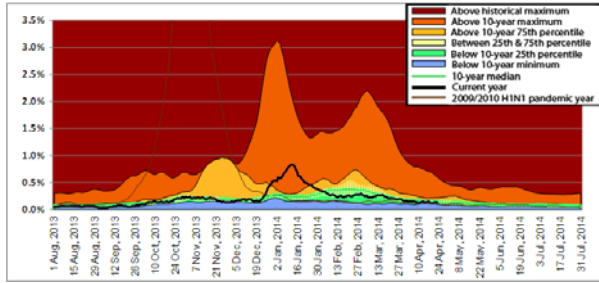
**Service claims submitted to MSP for influenza illness (II)\* as a proportion of all submitted general practitioner service claims, British Columbia, 2013-14**



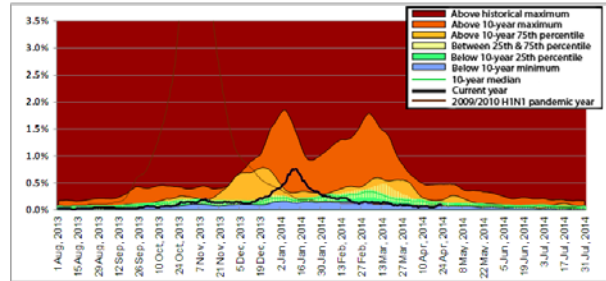
\* Influenza illness is tracked as the percentage of all submitted MSP general practitioner claims with ICD-9 code 487 (influenza). Data provided by Population Health Surveillance and Epidemiology, BC Ministry of Health Services

**Note:** MSP week beginning 1 August 2013 corresponds to sentinel ILI week 31; data current to 23 April 2014.

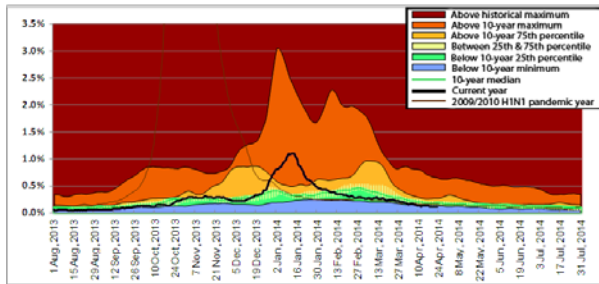
**Interior**



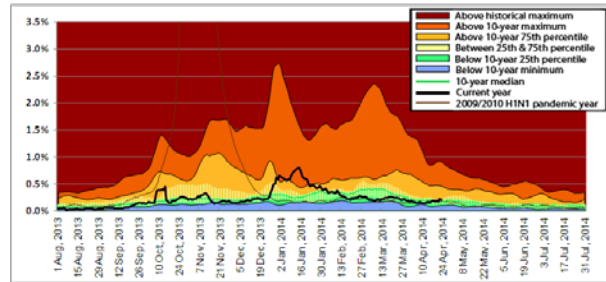
**Vancouver Island**



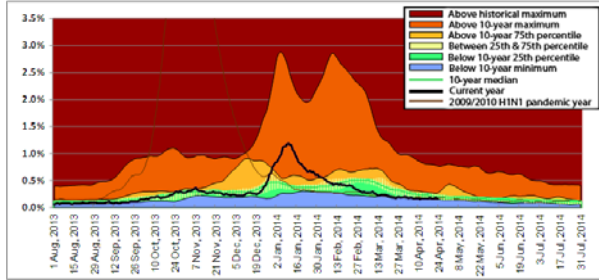
**Fraser**



**Northern**



**Vancouver Coastal**

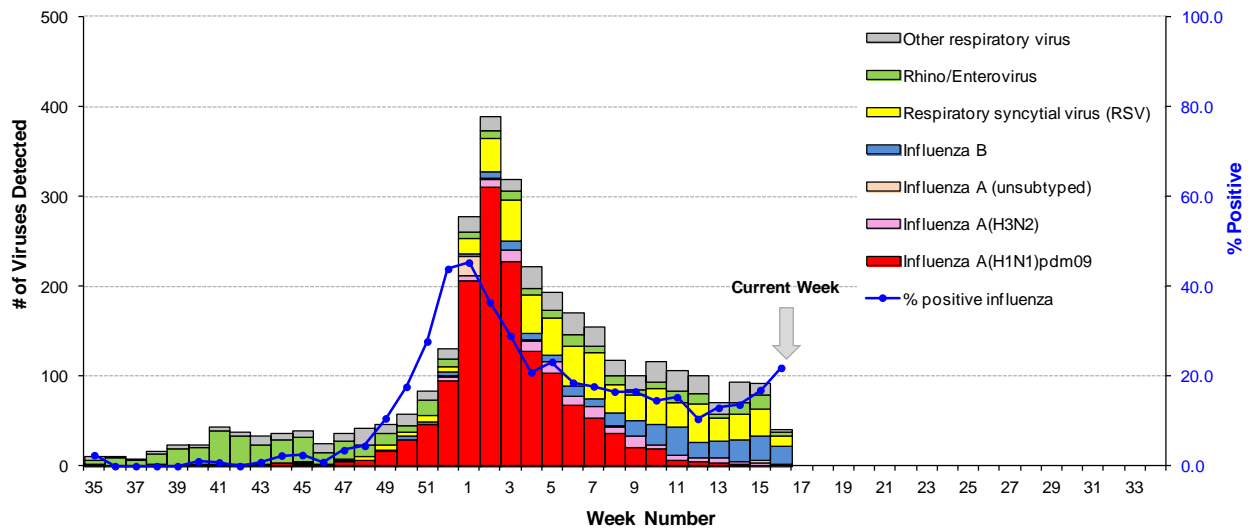


### Laboratory Reports

The proportion of specimens testing positive for influenza at the BC Public Health Microbiology & Reference Laboratory (PHMRL), PHSA, increased slightly from  $\leq 15\%$  in weeks 10-14 to 17% in week 15 and 22% in week 16, driven by ongoing influenza B circulation during this period concurrent with decreasing test volumes. Influenza B continued to predominate during this period, comprising 85-95% of all influenza positive specimens. Of the 291 respiratory specimens tested in weeks 15-16, 54 (19%) were positive for influenza, including 48/54 (89%) influenza B and 6/54 (11%) influenza A [3 A(H3N2) and 3 pending subtype]. RSV positivity decreased slightly from 15% in week 15 to 13% in week 16.

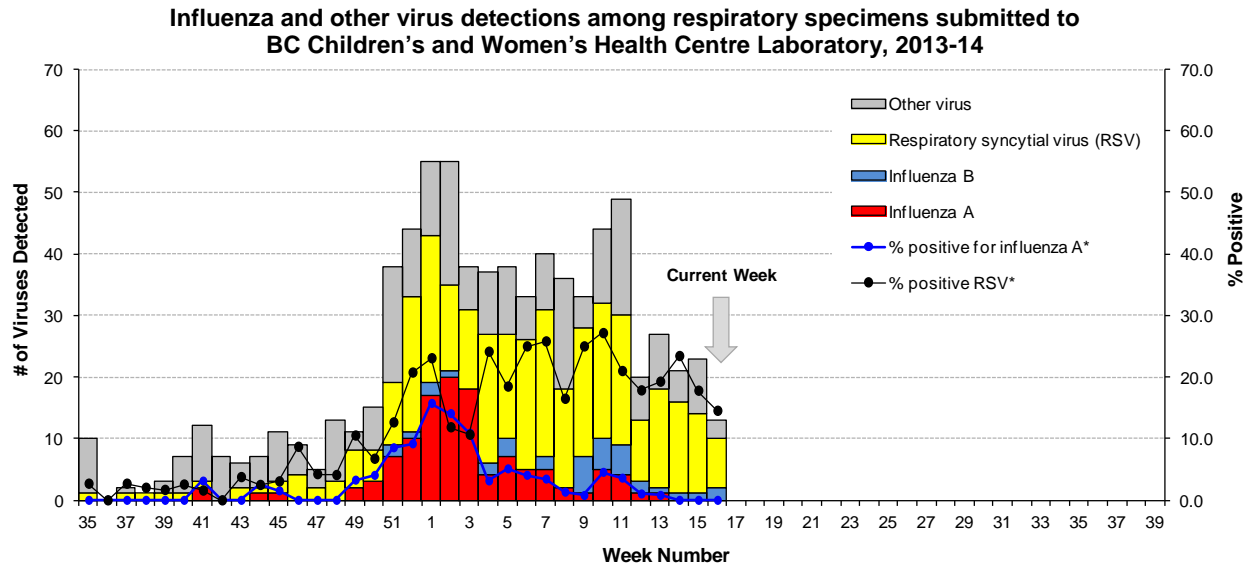
The 2013/14 influenza season to date has been characterized by predominant influenza A(H1N1)pdm09 activity, with ongoing late-season but less substantial influenza B circulation. Since week 40 (September 29 – October 5, 2013), 1,797 specimens have tested positive for influenza at the BC PHMRL. Of the 1,765 specimens with subtype information available, 1,378 (78%) were influenza A(H1N1)pdm09, 128 (7%) were influenza A(H3N2), and 259 (15%) were influenza B.

**Influenza and other virus detections among respiratory specimens submitted to BC Public Health Microbiology & Reference Laboratory, PHSA, 2013-14**



Note: PHMRL data current to April 21, 2014

At the BC Children's and Women's Health Centre Laboratory, the proportion of tests positive for influenza B increased from 1% in week 15 to 4% in week 16, reflecting ongoing influenza B circulation in BC. No influenza A viruses have been detected since week 14. RSV remained the most commonly detected respiratory virus; however, the proportion of tests positive for this virus decreased from 18% in week 15 to 15% in week 16.



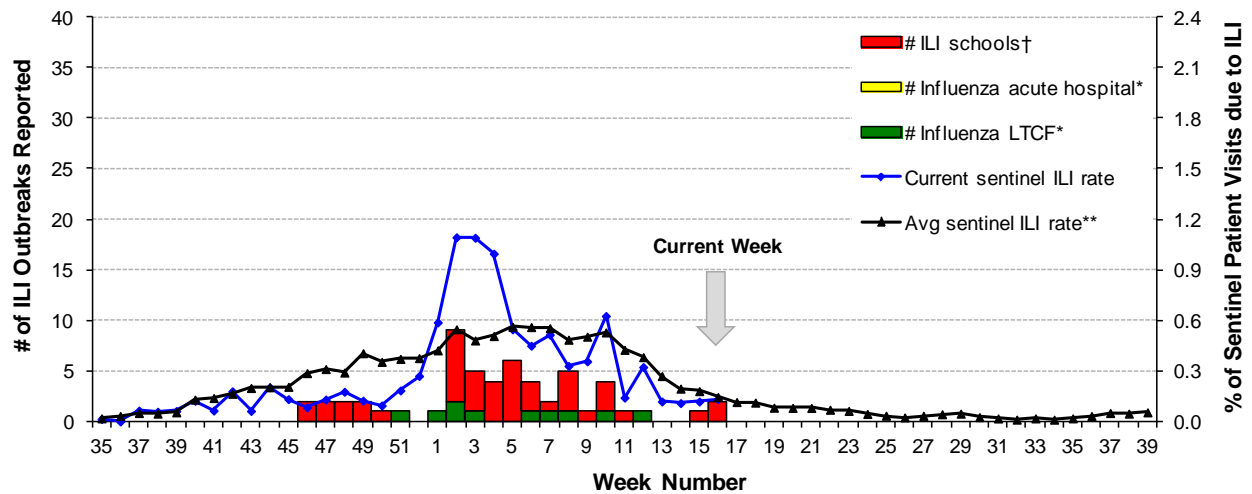
\* Positive rates were calculated using aggregate data. The denominators for each rate represent the total number of tests; multiple tests may be performed for a single specimen and/or patient.

### Influenza-like Illness (ILI) Outbreaks

In weeks 15-16, 5 ILI outbreaks were reported, including 3 school outbreaks from NHA (2 due to influenza B and 1 pending lab result) and 2 outbreaks from long-term care facilities (LTCF) in IHA (1 due to HMPV and 1 due to coronavirus). So far in week 17, three LTCF outbreaks have been reported from VIHA, IHA, and FHA, all with laboratory results pending.

In total during the 2013-14 season, 46 LTCF ILI outbreaks have been reported, including 10 outbreaks due to influenza viruses: 6 due to A(H1N1)pdm09 (2 in FHA, 3 in IHA, and 1 in VCHA); 2 due to A(H3N2) (both in FHA); 1 influenza A with subtype unknown due to insufficient viral copies in IHA; and 1 influenza B in FHA. In addition, 46 ILI outbreaks have been reported from schools so far this season, including one due to A(H1N1)pdm09 in week 47 and three due to influenza B in weeks 11, 15 and 16, all from NHA.

**Number of influenza-like illness (ILI) outbreaks reported, compared to current sentinel ILI rate and historical average sentinel ILI rate, British Columbia 2013-14**



\* Facility-based influenza outbreaks defined as 2 or more ILI cases within 7-day period, with at least one laboratory-confirmed case of influenza.

† School-based ILI outbreak defined as >10% absenteeism on any day, most likely due to ILI.

\*\* Historical values exclude 2008-09 and 2009-10 seasons due to atypical seasonality.

## National

### FluWatch (week 15):

In week 15, influenza activity in Canada remains within expected levels for this time of year but is being sustained by continued circulation of influenza B. Ontario reported a marked increase in influenza activity and some western provinces have also reported increases in detections of influenza B. The influenza positivity rate was 15% in week 15. Of the 722 positive influenza tests, 89 (12%) were influenza A [half A(H1N1)pdm09 and half A(H3N2) among those subtyped] and 633 (88%) influenza B. While the influenza A(H1N1) virus has mostly affected adults 20-64 years of age this season, influenza B is having a greater impact on adults 65 years of age and older, as well as young persons 5 to 19 years of age. The number of hospitalizations and deaths reported from participating regions so far this season are comparable to reports in past influenza seasons. Details are available at: [http://www.phac-aspc.gc.ca/fluwatch/13-14/w15\\_14/index-eng.php](http://www.phac-aspc.gc.ca/fluwatch/13-14/w15_14/index-eng.php).

### National Microbiology Laboratory (NML): Strain Characterization

From September 1, 2013 to April 24, 2014, 1,814 isolates were collected from provincial and hospital laboratories for antigenic characterization at the NML:

- 93 A/Texas/50/2012-like A(H3N2)<sup>¶</sup> from NS, NB, QC, ON, SK, AB, BC and YT
- 1,320 A/California/07/2009-like [A(H1N1)pdm09]\* from NL, PE, NS, NB, QC, ON, MB, SK, AB, BC, NT and NU; of these, 2 viruses showed reduced titres with antiserum produced against A/California/7/2009 signalling possible antigenic change
- 381 B/Massachusetts/02/2012-like<sup>†</sup> from NL, NS, NB, QC, ON, MB, SK, AB and BC
- 20 B/Brisbane/60/2008-like\*\* from QC, ON, MB, SK, AB, and BC

<sup>¶</sup> Virus most closely related to the recommended H3N2 reference virus for the 2013-14 northern hemisphere influenza vaccine.

\* Virus most closely related to the recommended H1N1 reference virus for the 2013-14 northern hemisphere influenza vaccine.

<sup>†</sup> Virus most closely related to the recommended influenza B component for the 2013-14 northern hemisphere influenza vaccine; belongs to the B Yamagata lineage.

\*\* Virus most closely related to the recommended influenza B component for the 2011-2012 northern hemisphere influenza vaccine; belongs to the B Victoria/02/87 lineage.

### NML: Antiviral Resistance

From September 1, 2013 to April 24, 2014, drug susceptibility testing was performed at the NML for influenza viruses: 1,474 influenza A [119 A(H3N2) and 1,355 A(H1N1)pdm09] viruses were tested for resistance to amantadine; 1,557 influenza viruses [75 A(H3N2), 1,216 A(H1N1)pdm09, and 266 B] were tested for resistance to oseltamivir; and 1,479 influenza viruses [70 A(H3N2), 1,180 A(H1N1)pdm09, and 229 B] were tested for resistance to zanamivir. All tested influenza A viruses were resistant to amantadine. All but two tested viruses were sensitive to oseltamivir, and all were sensitive to zanamivir. Both viruses resistant to oseltamivir were A(H1N1)pdm09 viruses with a H275Y mutation.



## International

**USA (week 15):** Influenza activity continued to decrease in most regions of the United States in week 15. Of the 4,653 specimens tested, 675 (15%) were positive for influenza viruses, of which 45% were influenza A [7% A(H1N1)pdm09, 46% A(H3N2), 47% untyped] and 55% were influenza B. While overall detections of influenza have been relatively low, the proportion of influenza B continues to surge. The proportion of deaths attributed to pneumonia and influenza was below the epidemic threshold and the proportion of outpatient visits for influenza-like illness (ILI) was 1.5%, below the national baseline of 2%. Widespread influenza activity was reported from six states over this period. Details are available at: [www.cdc.gov/flu/weekly/](http://www.cdc.gov/flu/weekly/).

**WHO (as of 22 April 2014):** Globally, the northern hemisphere influenza season appeared to be approaching interseasonal levels in most countries. As influenza detections declined, the proportion of influenza B detections increased slightly in many regions, especially Asia, the Middle East, and North America. In Europe, influenza activity continued to decrease, as the influenza season appeared to be winding down in the region. A rise in the percentage of influenza specimens testing positive was observed, but the overall number of specimens declined. Influenza A(H3N2) and A(H1N1)pdm09 co-circulated, with low numbers of influenza B virus detected. In Eastern Europe, influenza activity was higher later in the season compared to the north and the south-west, but detections have begun to decline as well. In Eastern Asia, influenza activity approached interseasonal levels, and influenza B comprised the majority of influenza detections. In Tropical Asia, influenza activity continued to decline. In Northern Africa and Western Asia, influenza activity remained low in most countries, with influenza B the predominant virus detected. In the Southern Hemisphere, influenza activity remained low and detections were sporadic. During weeks 13 and 14 (23 March 2014 to 5 April 2014), WHO GISRS laboratories tested more than 44,319 specimens. Of these, 6,717 were positive for influenza viruses: 4,163 (62%) were typed as influenza A and 2,554 (38%) as influenza B. Of the sub-typed influenza A viruses, 1,149 (47%) were influenza A(H1N1)pdm09 and 1,287 (53%) were influenza A(H3N2). The majority of characterized B viruses were found to belong to the B-Yamagata lineage. Details are available at: [www.who.int/influenza/surveillance\\_monitoring/updates/latest\\_update\\_GIP\\_surveillance/en/](http://www.who.int/influenza/surveillance_monitoring/updates/latest_update_GIP_surveillance/en/).

**Avian Influenza A(H7N9) Virus:** Since our last surveillance bulletin, 16 new cases of human infection with avian influenza A(H7N9) have been reported, including one travel-related case reported from Hong Kong and one (unconfirmed) travel-related case from Taiwan. To date (as of 23 April 2014), the WHO has been informed of 427 laboratory-confirmed cases and 146 deaths. At this time, there is no evidence of sustained human-to-human transmission and the risk assessment remains unchanged. Clinicians should remain vigilant for patients presenting with severe acute respiratory illness (SARI) with recent travel or epidemiological links to affected areas. Details are available at: [www.who.int/csr/don/en/](http://www.who.int/csr/don/en/).

**Middle East Respiratory Syndrome Coronavirus (MERS-CoV):** Since the beginning of April 2014, there has been a substantial increase in case reports of MERS-CoV. Most of these reports have been associated with large nosocomial outbreaks in Jeddah, Saudi Arabia, and Abu Dhabi, United Arab Emirates (UAE), including asymptomatic cases identified through case tracking and affecting mostly health care workers, as well as ongoing case reports from the Riyadh region in persons with chronic comorbidity. Also this month, Malaysia and Greece reported their first imported cases of MERS-CoV in persons with recent travel to affected regions of the Middle East. Globally, from September 2012 to date (as of 23 April 2014), the WHO has confirmed 254 cases of MERS-CoV, including 93 deaths. However, these official WHO counts do not include a large number of cases recently reported by the Saudi Ministry of Health, which would put the global tally at over 300 cases, an increase of more than 160 cases so far this month. Given ongoing activity in affected regions, clinicians are reminded to stay alert for possible importations among patients presenting with severe acute respiratory illness (SARI) and links to the Middle East. Details are available at: [www.who.int/csr/don/en/](http://www.who.int/csr/don/en/).

### **WHO Recommendations for 2013-14 Northern Hemisphere Influenza Vaccine**

On February 21, 2013, the WHO announced the recommended strain components for the 2013-14 northern hemisphere vaccine:

- A/California/7/2009 (H1N1)pdm09 virus
- A/Victoria/361/2011 (H3N2)-like virus\*
- B/Massachusetts/2/2012-(Yamagata lineage)-like virus\*\*

\*It is recommended that A/Texas/50/2012 be used as the A(H3N2) vaccine component because of antigenic changes in earlier A/Victoria/361/2011-like vaccine viruses (such as IVR-165) resulting from adaptation to propagation in eggs.

\*\* This one of the three recommended components is different from the northern hemisphere seasonal TIV vaccines produced and administered in 2012-13 (although remaining of the same lineage).

For further details, see:

[www.who.int/influenza/vaccines/virus/recommendations/2013\\_14\\_north/en/index.html](http://www.who.int/influenza/vaccines/virus/recommendations/2013_14_north/en/index.html).

### **WHO Recommendations for 2014-15 Northern Hemisphere Influenza Vaccine**

On February 20, 2014, the WHO announced the recommended strain components for the 2014-15 northern hemisphere vaccine:

- A/California/7/2009 (H1N1)pdm09 virus
- A/Texas/50/2012 (H3N2)-like virus
- B/Massachusetts/2/2012-(Yamagata lineage)-like virus

These recommended strains are the same as those used for the 2013-14 northern hemisphere vaccine.

For further details, see: [www.who.int/influenza/vaccines/virus/recommendations/2014\\_15\\_north/en/](http://www.who.int/influenza/vaccines/virus/recommendations/2014_15_north/en/).

## **Additional Information**

### **List of Acronyms:**

**ACF:** Acute Care Facility

**AI:** Avian influenza

**FHA:** Fraser Health Authority

**HBoV:** Human bocavirus

**HMPV:** Human metapneumovirus

**HSDA:** Health Service Delivery Area

**IHA:** Interior Health Authority

**ILI:** Influenza-Like Illness

**LTCF:** Long-Term Care Facility

**MSP:** BC Medical Services Plan

**NHA:** Northern Health Authority

**NML:** National Microbiological Laboratory

**A(H1N1)pdm09:** Pandemic H1N1 influenza (2009)

**RSV:** Respiratory syncytial virus

**VCHA:** Vancouver Coastal Health Authority

**VIHA:** Vancouver Island Health Authority

**WHO:** World Health Organization

### **Recently updated AMMI Canada Guidelines on the Use of Antiviral Drugs for Influenza:**

[www.ammi.ca/guidelines](http://www.ammi.ca/guidelines)

### **Web Sites:**

BCCDC Emerging Respiratory Pathogen Updates:

[www.bccdc.ca/dis-cond/DiseaseStatsReports/EmergingRespiratoryVirusUpdates.htm](http://www.bccdc.ca/dis-cond/DiseaseStatsReports/EmergingRespiratoryVirusUpdates.htm)

### **Influenza Web Sites**

Canada – Flu Watch: [www.phac-aspc.gc.ca/fluwatch/](http://www.phac-aspc.gc.ca/fluwatch/)

Washington State Flu Updates: [www.doh.wa.gov/Portals/1/Documents/5100/fluupdate.pdf](http://www.doh.wa.gov/Portals/1/Documents/5100/fluupdate.pdf)

USA Weekly Surveillance Reports: [www.cdc.gov/flu/weekly/](http://www.cdc.gov/flu/weekly/)

European Influenza Surveillance Scheme:

[ecdc.europa.eu/EN/HEALTHTOPICS/SEASONAL\\_INFLUENZA/EPIDEMIOLOGICAL\\_DATA/Pages/Weekly\\_Influenza\\_Surveillance\\_Overview.aspx](http://ecdc.europa.eu/EN/HEALTHTOPICS/SEASONAL_INFLUENZA/EPIDEMIOLOGICAL_DATA/Pages/Weekly_Influenza_Surveillance_Overview.aspx)

WHO – Weekly Epidemiological Record: [www.who.int/wer/en/](http://www.who.int/wer/en/)

WHO Collaborating Centre for Reference and Research on Influenza (Australia):

[www.influenzacentre.org/](http://www.influenzacentre.org/)

Australian Influenza Report:

[www.health.gov.au/internet/main/publishing.nsf/content/cda-surveil-ozflu-flucurr.htm](http://www.health.gov.au/internet/main/publishing.nsf/content/cda-surveil-ozflu-flucurr.htm)

New Zealand Influenza Surveillance Reports: [www.surv.esr.cri.nz/virology/influenza\\_weekly\\_update.php](http://www.surv.esr.cri.nz/virology/influenza_weekly_update.php)

### **Avian Influenza Web Sites**

WHO – Influenza at the Human-Animal Interface: [www.who.int/csr/disease/avian\\_influenza/en/](http://www.who.int/csr/disease/avian_influenza/en/)

World Organization for Animal Health: [www.oie.int/eng/en\\_index.htm](http://www.oie.int/eng/en_index.htm)

### **Contact Us:**

Tel: (604) 707-2510

Fax: (604) 707-2516

Email: [InfluenzaFieldEpi@bccdc.ca](mailto:InfluenzaFieldEpi@bccdc.ca)

Communicable Disease Prevention and Control Services (CDPACS)

BC Centre for Disease Control

655 West 12<sup>th</sup> Ave, Vancouver BC V5Z 4R4

Online: [www.bccdc.ca/dis-cond/DiseaseStatsReports/influSurveillanceReports.htm](http://www.bccdc.ca/dis-cond/DiseaseStatsReports/influSurveillanceReports.htm)

# Influenza-Like Illness (ILI) Outbreak Summary Report Form

Please complete and email to [ilioutbreak@bccdc.ca](mailto:ilioutbreak@bccdc.ca)

**Note: This form is for provincial surveillance purposes.  
Please notify your local health unit per local guidelines/requirements.**

**ILI:** Acute onset of respiratory illness with fever and cough and with one or more of the following: sore throat, arthralgia, myalgia, or prostration which *could* be due to influenza virus. In children under 5, gastrointestinal symptoms may also be present. In patients under 5 or 65 and older, fever may not be prominent.

**Schools and work site outbreak:** greater than 10% absenteeism on any day, most likely due to ILI.

**Residential institutions (facilities) outbreak:** two or more cases of ILI within a seven-day period.

<b>A</b>	<b><u>Reporting Information</u></b>	Health unit/medical health officer notified? <input type="checkbox"/> Yes <input type="checkbox"/> No
	Person Reporting: _____	Title: _____
	Contact Phone: _____	Email: _____
	Health Authority: _____	HSDA: _____
	Full Facility Name: _____	
	Is this report:	<input type="checkbox"/> First Notification ( <i>complete section B below; Section D if available</i> ) <input type="checkbox"/> Update ( <i>complete section C below; Section D if available</i> ) <input type="checkbox"/> Outbreak Over ( <i>complete section C below; Section D if available</i> )

<b>B</b>	<b><u>First Notification</u></b>
	Type of facility: <input type="checkbox"/> LTCF <input type="checkbox"/> Acute Care Hospital <input type="checkbox"/> Senior's Residence (if ward or wing, please specify name/number: _____)
	<input type="checkbox"/> Workplace <input type="checkbox"/> School (grades: _____) <input type="checkbox"/> Other (_____)
	Date of onset of first case of ILI (dd/mm/yyyy): <u>DD / MMM / YYYY</u>

Numbers to date	Residents/Students	Staff
<b>Total</b>		
<b>With ILI</b>		
<b>Hospitalized</b>		
<b>Died</b>		

<b>C</b>	<b><u>Update AND Outbreak Declared Over</u></b>
	Date of onset for most recent case of ILI (dd/mm/yyyy): <u>DD / MMM / YYYY</u>
	If over, date outbreak declared over (dd/mm/yyyy): <u>DD / MMM / YYYY</u>

Numbers to date	Residents/Students	Staff
<b>Total</b>		
<b>With ILI</b>		
<b>Hospitalized</b>		
<b>Died</b>		

<b>D</b>	<b><u>Laboratory Information</u></b>
	Specimen(s) submitted? <input type="checkbox"/> Yes (location: _____) <input type="checkbox"/> No <input type="checkbox"/> Don't know If yes, organism identified? <input type="checkbox"/> Yes (specify: _____) <input type="checkbox"/> No <input type="checkbox"/> Don't know