

British Columbia Influenza Surveillance Bulletin

Influenza Season 2017-18, Number 13, Week 6

February 4 to 10, 2018

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Passed the Epidemic Peak but Influenza Activity Remains above Seasonal Levels

During week 6 (February 4 to 10, 2018), most influenza surveillance indicators continued to decline but influenza activity remained above seasonal levels in most regions.

Influenza positivity at the BCCDC Public Health Laboratory continued to decline, falling to 30% in week 6 from a peak of more than 50% in week 52, driven by declining A(H3N2) activity. Influenza B has predominated among influenza detections (65%) this week with type B positivity remaining stable around 20%.

Since our last bulletin, 11 new lab-confirmed outbreaks were reported, 10 from long-term care facilities (LTCFs) and 1 from an acute care hospital; 1 school ILI outbreak was reported. Of the 11 outbreaks, 9 had influenza B detected, and 2 had influenza A detected; the 1 influenza A outbreak that had subtype information available was A(H3N2).

On February 15, the US CDC published interim estimates of influenza vaccine effectiveness (VE) for the 2017-18 season. Adjusted VE for A(H3N2) was low at 25%, but higher for influenza B at 42%. The full report is available here:

https://www.cdc.gov/mmwr/volumes/67/wr/mm6706a2.htm?s_cid=mm6706a2_e

Prepared by BCCDC Influenza & Emerging Respiratory Pathogens Team

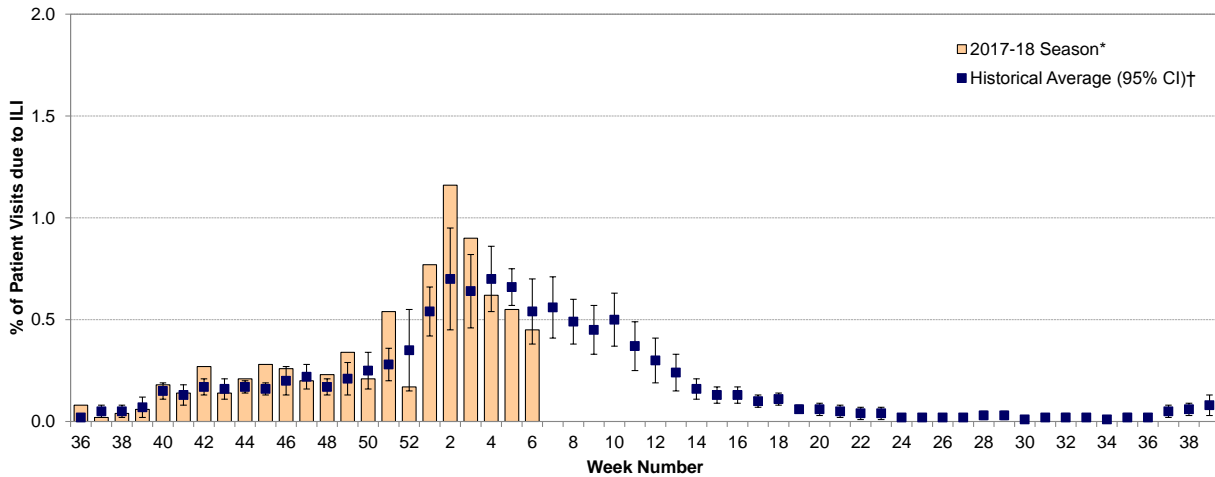
Report Disseminated: February 15, 2018

British Columbia

Sentinel Physicians

The proportion of patients with influenza-like illness (ILI), among those presenting to sentinel sites, is below average but within expected levels for week 6, continuing a downward trend since week 2. Rates are subject to change as reporting becomes more complete. To date, 40% of sentinel sites have reported data for week 6.

Percent of patient visits to sentinel physicians due to influenza-like illness (ILI) compared to historical average, British Columbia, 2017-18



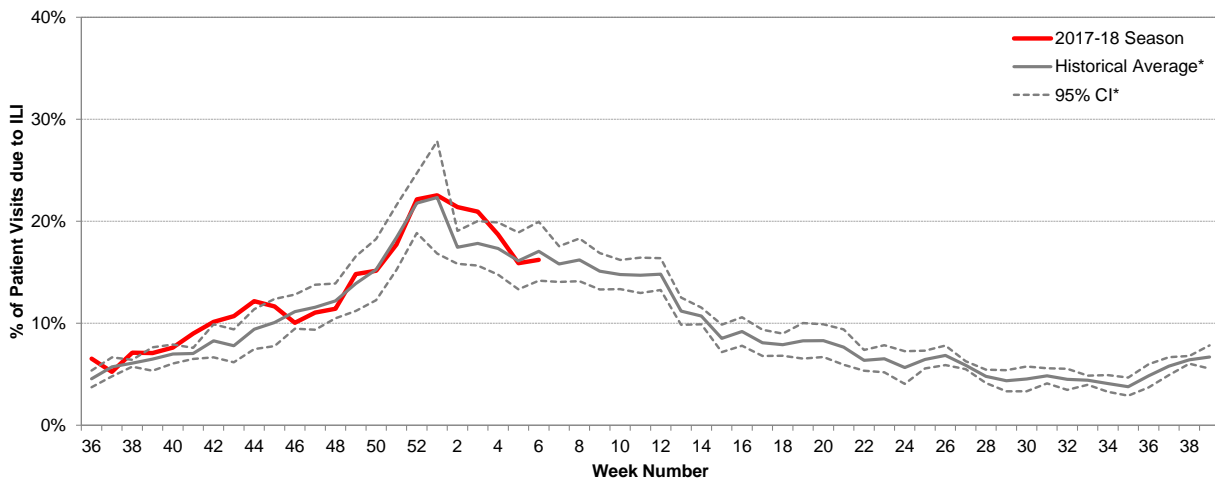
* Data are subject to change as reporting becomes more complete.

† 10-year historical average for 2017-18 season based on 2005-06 to 2016-2017 seasons, excluding 2008-09 and 2009-10 due to atypical seasonality; CI=confidence interval.

BC Children's Hospital Emergency Room

In week 6, the proportion of visits to BC Children's Hospital Emergency Room (ER) attributed to ILI remained within expected levels for this period.

Percent of patients presenting to BC Children's Hospital ER attributed to influenza-like illness (ILI), British Columbia, 2017-18



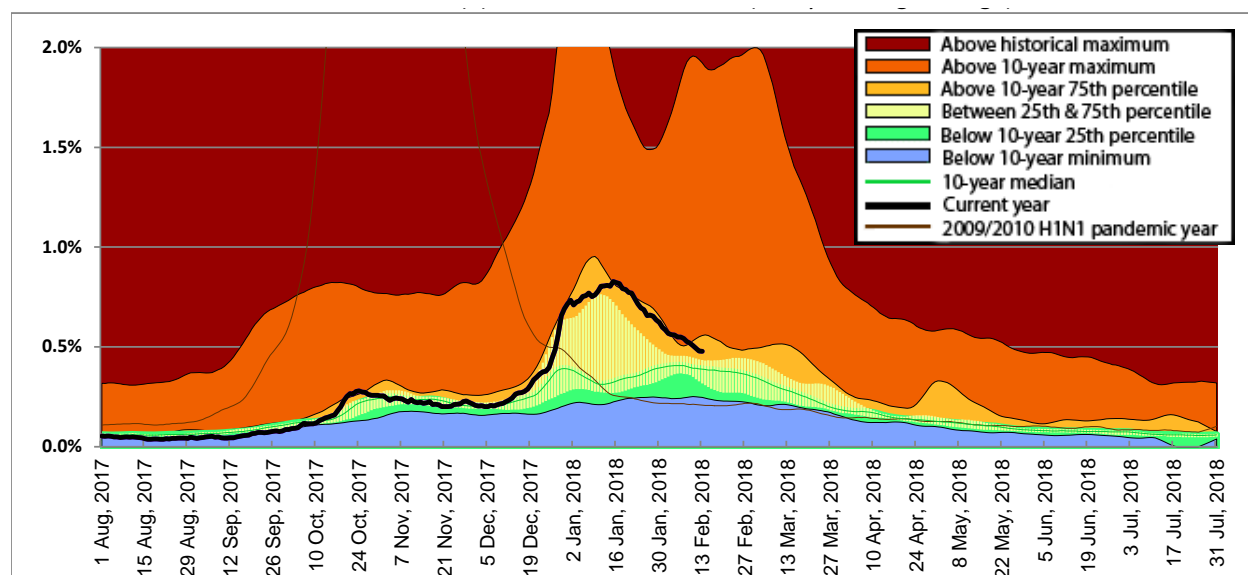
Source: BCCCH Admitting, Discharge, Transfer database (ADT). Data includes records with a triage chief complaint of "flu" or "influenza" or "fever/cough."

* 5-year historical average for 2017-18 season based on 2012-13 to 2016-17 seasons; CI=confidence interval.

Medical Services Plan

In week 6, BC Medical Services Plan (MSP) general practitioner claims for influenza illness (II), as a proportion of all submitted MSP claims continued to decline in most regions of the province following several weeks of elevated activity overall, with the exception of NHA where rates increased and were above 10-year maximum rates for this time of year. In week 6, rates for the province overall and FHA, VCHA and VIHA were above the 10-year 75th percentile, while rates in IHA were at expected levels for this time of year.

Service claims submitted to MSP for influenza illness (II)* as a proportion of all submitted general practitioner service claims, British Columbia, 2017-18

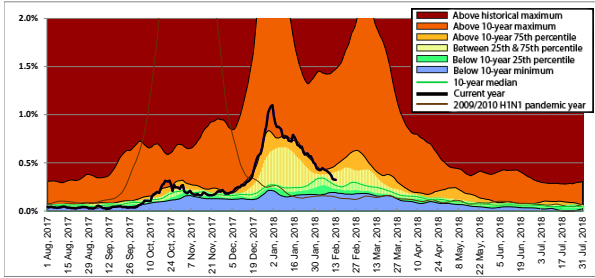


* Influenza illness is tracked as the percentage of all submitted MSP general practitioner claims with ICD-9 code 487 (influenza).

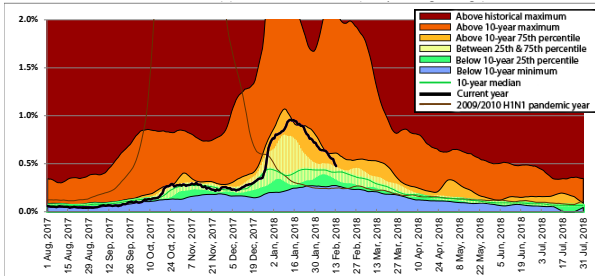
Data for the period August 1, 2009 to July 31, 2010 have been excluded from the 10-year median calculation due to atypical seasonality during the 2009/2010 H1N1 pandemic year. MSP data beginning August 1, 2017 corresponds to sentinel ILI week 31; data are current to February 13, 2018.

Data provided by Population Health Surveillance and Epidemiology, BC Ministry of Health Services.

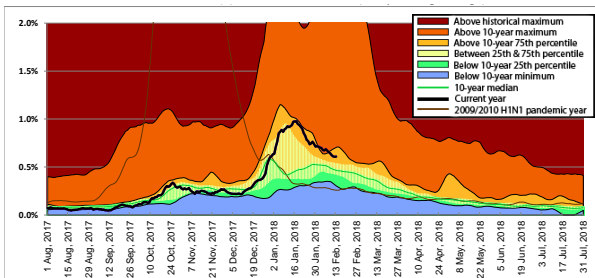
Interior



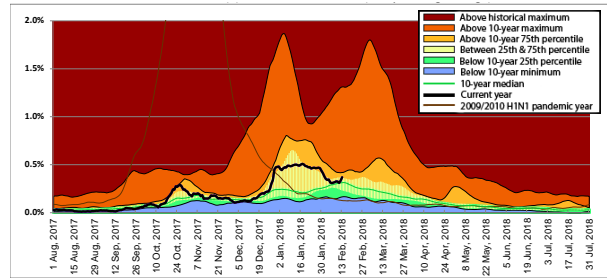
Fraser



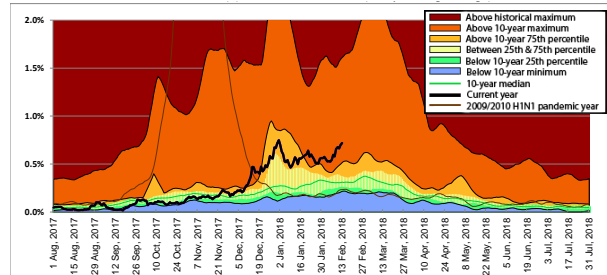
Vancouver Coastal



Vancouver Island



Northern



Laboratory Reports

BCCDC Public Health Laboratory

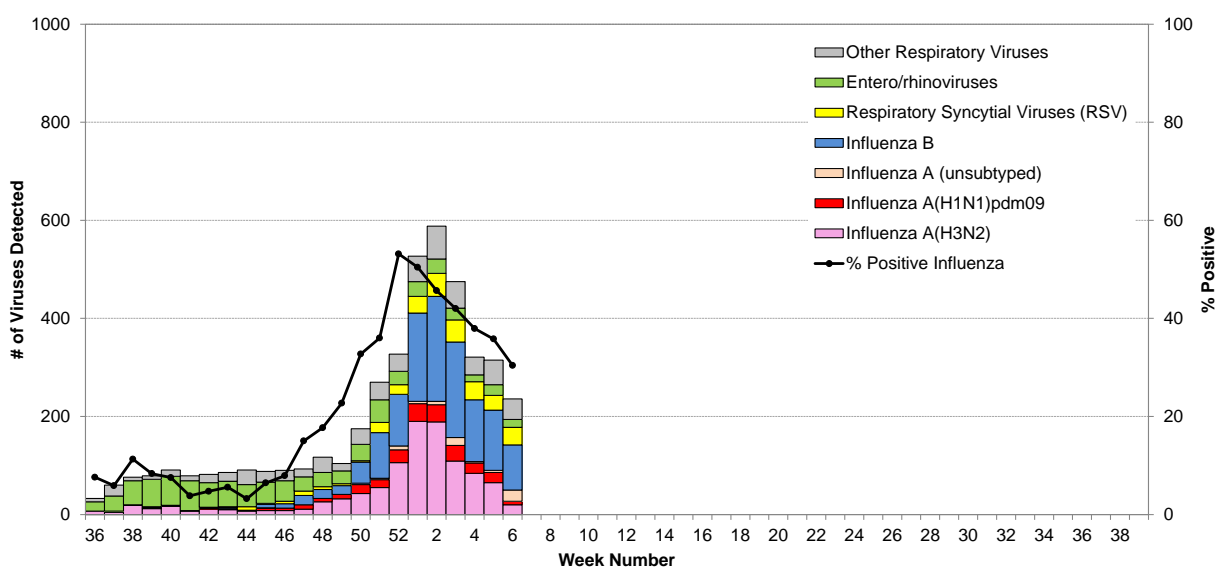
In week 6, 467 patients were tested for respiratory viruses at the BCCDC Public Health Laboratory (PHL). Of these, 141 (30%) tested positive for influenza; 49 (35%) had influenza A detected [19 A(H3N2), 7 A(H1N1)pdm09 and 23 subtype pending], 91 (65%) had influenza B detected and 1 had A(H3N2) and B detected. Influenza positivity at the BCCDC PHL declined to 30% in week 6 from a peak of more than 50% in week 52, driven by declining detection of A(H3N2), the dominant influenza A subtype this season. Influenza B positivity remained greater than in previous years for this period, comprising about two-thirds of all influenza detections in week 6 and with influenza B positivity rates remaining stable around 20%. Based on national surveillance reports, B(Yamagata) is the predominant influenza B lineage circulating in BC and elsewhere in Canada this season.

Cumulatively during the 2017-18 season (since week 40, starting October 1, 2017), 2535 (32%) patients tested positive for influenza at the BCCDC PHL, including 1297 (51%) with influenza A [981 A(H3N2), 246 A(H1N1)pdm09, 70 subtype pending], 1228 (48%) with influenza B and 10 patients with both influenza A [eight with A(H3N2) and two with A(H1N1)pdm09] and B detected.

More than half (60%) of A(H3N2) cases have been detected among elderly adults ≥ 65 years old, with 8% <20 years old, 17% 20-49 years old, and 15% 50-64 years old. Conversely, 40% of influenza B cases have been detected among elderly adults ≥ 65 years old, with 17% <20 years old, 24% 20-49 years old, and 19% 50-64 years old. Among A(H1N1)pdm09 cases, only 16% have been detected among elderly adults ≥ 65 years old, with 29% <20 years old, 38% 20-49 years old, and 17% 50-64 years old.

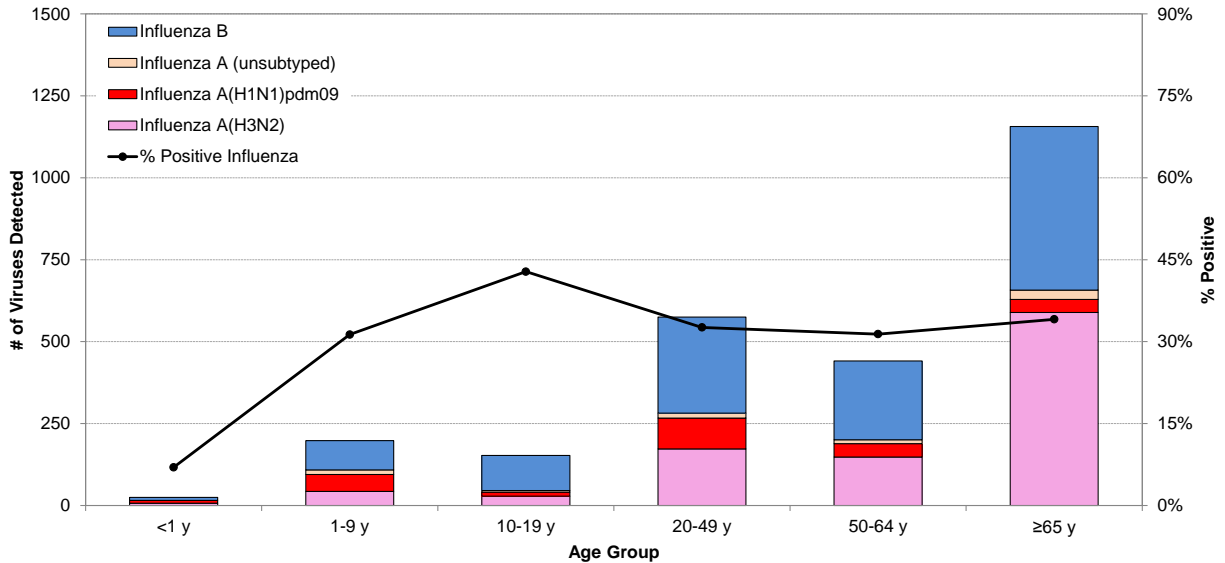
RSV was the most commonly detected non-influenza respiratory virus during this period. RSV detections have been less frequent than in the 2016-17 season; 8% of patients tested positive for RSV in week 6 this season compared to 17% in the 2016-17 season during the same period.

Influenza and other virus detections among respiratory specimens submitted to BCCDC Public Health Laboratory, 2017-18



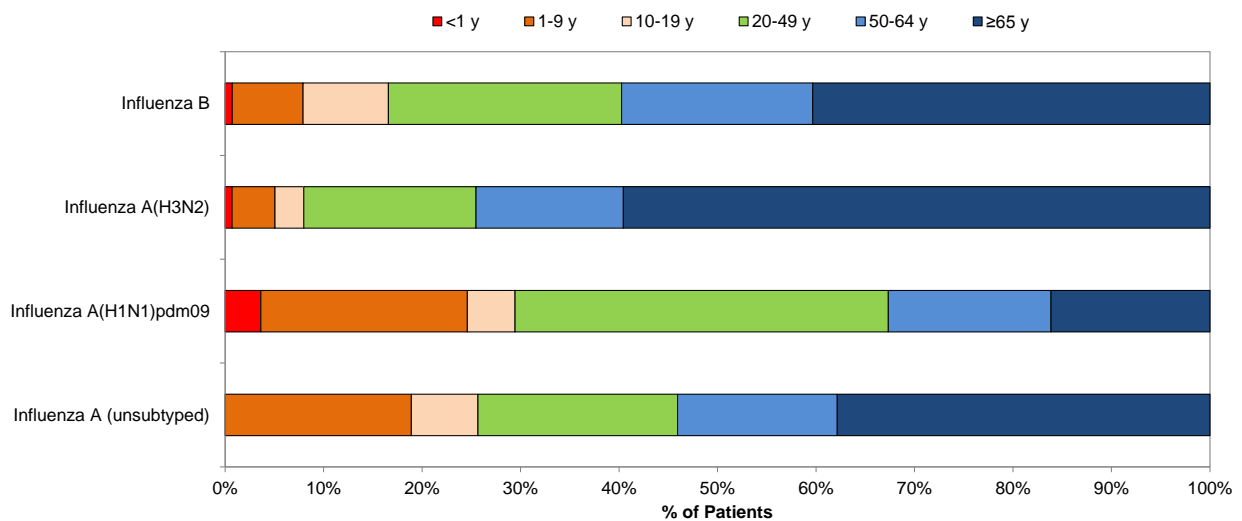
Source: BCCDC Public Health Laboratory (PHDRW); Data are current to February 14, 2018.

Cumulative number (since week 40) of influenza detections by type subtype and age group, BCCDC Public Health Laboratory, 2017-18



Source: BCCDC Public Health Laboratory (PHDRW); Data are current to February 14, 2018; figure includes cumulative influenza detections for specimens collected from weeks 40-6.

Age distribution of influenza detections (cumulative since week 40), BCCDC Public Health Laboratory, 2017-18

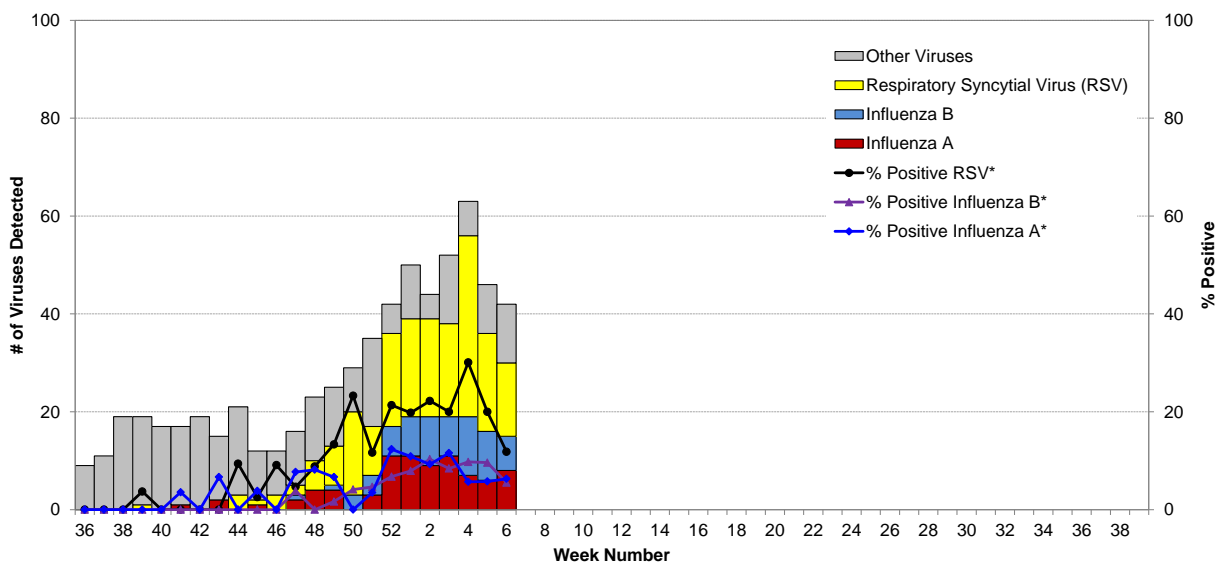


Source: BCCDC Public Health Laboratory (PHDRW); Data are current to February 14, 2018; figure includes cumulative influenza detections for specimens collected from weeks 40-6.

BC Children's and Women's Health Centre Laboratory

In week 6, 127 tests for influenza viruses were conducted at the BC Children's and Women's Health Centre (CWHC) laboratory. Of these, 8 (6%) were positive for influenza A and 7 (6%) were positive for influenza B. Respiratory syncytial virus (RSV) was the most commonly detected respiratory viruses during this period, with 12% positivity in week 6. Akin to observations from the BCCDC PHL, RSV positivity from this week was substantially lower than week 6 in the 2016/17 season where RSV positivity was 25%; this represents a shift from past weeks where RSV positivity was similar to the previous season.

Influenza and other virus detections among respiratory specimens submitted to BC Children's and Women's Health Centre Laboratory, 2017-18



* 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

Since our last bulletin, 11 new lab-confirmed outbreaks were reported, 10 from long-term care facilities (LTCFs) and 1 from an acute care hospital. Of the 11 newly reported outbreaks, 2 had onset in week 4 in VCHA, 5 had onset in week 5 (1 in FHA, 1 in IHA, 3 in VCHA), and 4 had onset in week 6 (2 in FHA, 2 in VCHA). Of the 11 outbreaks, 9 had influenza B detected, and 2 had influenza A detected; the 1 influenza A outbreak that had subtype information available was A(H3N2).

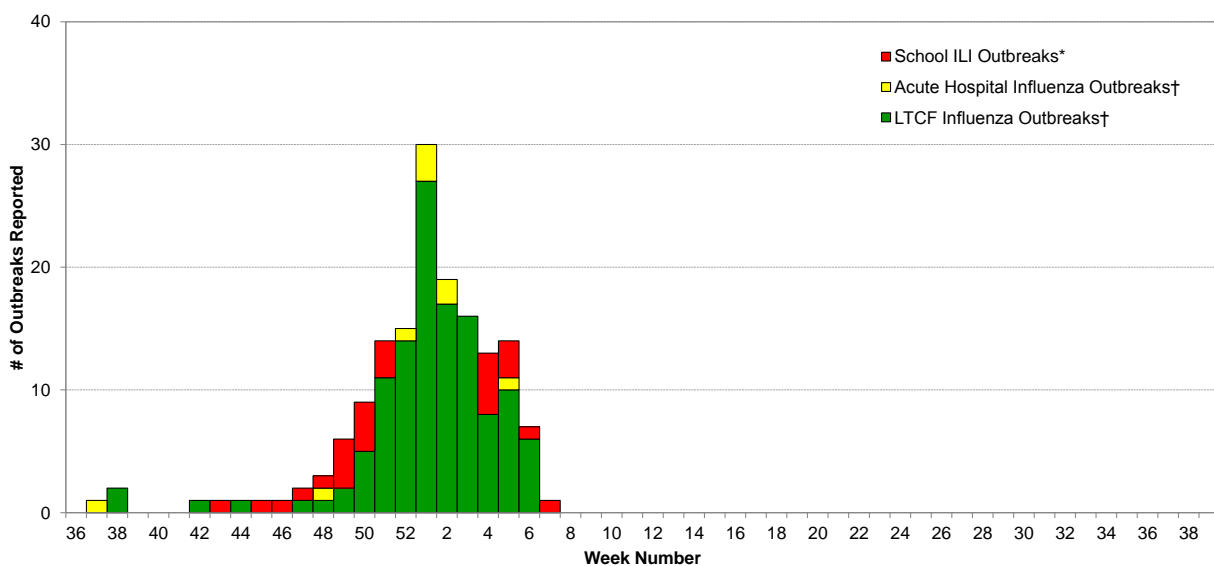
Additionally, 1 school ILI outbreak, with unknown etiology, was reported during week 7. This outbreak occurred in IHA, currently the only health authority routinely reporting school ILI outbreaks to BCCDC.

Influenza outbreak reports appear to have declined in frequency following a peak in week 1; this may reflect declining influenza activity in the province but could also be attributed to delayed reporting. The majority of outbreaks reported in recent weeks have been due to influenza B.

Cumulatively during the 2017-18 season (since week 37, starting September 10, 2017), 131 lab-confirmed influenza outbreaks have been reported, including 44 with influenza A detected [22 A(H3N2) and 22 subtype unknown], 77 with influenza B, 3 with influenza A (H3N2) and influenza B, and 7 with influenza A (unspecified subtype) and influenza B; of these, 122 were reported in LTCFs and 9 were reported from an acute care facility. No influenza A outbreaks have been subtyped as A(H1N1)pdm09 so far this season. Additionally, 26 school ILI outbreaks have occurred without etiologic agent identified.

So far during this season of mixed A(H3N2) and influenza B co-circulation, the number of long term care facility outbreaks reported since week 40 (n=120) is lower than the tally for the same period during recent A(H3N2) dominant epidemics in 2014-15 (n=147) and 2016-17 (n=155) but higher than during recent A(H1N1)pdm09 dominant epidemics in 2013-14 (n=6) and 2015-16 (n=16), bearing in mind variation in the timing of seasonal epidemics that may influence final end-of-season comparisons.

Number of influenza-like illness (ILI) outbreaks reported, British Columbia 2017-18



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

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

Updated Antiviral Guidelines

The Association of Medical Microbiology and Infectious Disease Canada (AMMI Canada) have released updated guidance on the use of antiviral drugs given potential low vaccine effectiveness for the 2017-18 influenza season. These guidelines are available at: <https://www.ammi.ca/Update/79.ENG.pdf>.

National

FluWatch (week 5, January 28 to February 3, 2018)

Overall, influenza activity in Canada remains at peak levels but there are signs that activity is starting to slow down in parts of the country. In week 5, the total number of detections of influenza B was similar to the total number of detections of influenza A. An increasing proportion of weekly pediatric hospitalizations reported by the IMPACT network are due to influenza B. In week 5, influenza A and B accounted for an equal proportion of hospitalizations. To date this season, the majority of lab confirmations, hospitalizations and deaths have been among adults 65 years of age and older. Details are available at: www.canada.ca/en/public-health/services/diseases/flu-influenza/influenza-surveillance/weekly-influenza-reports.html.

National Microbiology Laboratory (NML): Strain Characterization

From September 1, 2017 to February 15, 2018, the National Microbiology Laboratory (NML) received 1,443 influenza viruses from Canadian laboratories for antigenic characterization.

Influenza A(H3N2): Of the 810 influenza A(H3N2) viruses, only 163 (20%) had sufficient haemagglutination titre for antigenic characterization by haemagglutination inhibition (HI) assay. Of the 163 viruses characterized by HI assay, 148 (91%) were considered antigenically similar to a cell culture-propagated A/Hong Kong/4801/2014-like virus, the WHO-recommended A(H3N2) component for the 2017-18 northern hemisphere influenza vaccine, while 15 (9%) viruses (all belonging to genetic clade 3C.3a) showed reduced titre with ferret antisera raised against cell culture-propagated A/Hong Kong/4801/2014. Of the 162 out of 163 viruses that were antigenically characterized with available sequencing information, 132 belonged to genetic clade 3C.2a, 15 belonged to subclade 3C.2a1 and 15 belonged to clade 3C.3a; sequencing is pending for the remaining isolate. Of the 647 viruses genetically characterized, 570 (88%) were reported to belong to genetic clade 3C.2a, which includes the A/Hong Kong/4801/2014 vaccine strain, while 75 (12%) belonged to subclade 3C.2a1 and 2 belonged to clade 3C.3a.

Influenza A(H1N1)pdm09: All of the 72 A(H1N1)pdm09 viruses characterized were antigenically similar to an A/Michigan/45/2015-like virus, the WHO-recommended influenza A(H1N1) component for the 2017-18 northern hemisphere influenza vaccine.

Influenza B: Of the 561 influenza B viruses characterized, 24 (4%) belonged to the B(Victoria) lineage and 537 (96%) belonged to the B(Yamagata) lineage. Among the 24 B(Victoria) viruses, 6 (25%) were characterized as antigenically similar to a B/Brisbane/60/2008(Victoria)-like virus, the WHO-recommended influenza B component for the 2017-18 northern hemisphere trivalent influenza vaccine, while 18 (75%) viruses showed reduced titre with ferret antisera produced against cell-propagated B/Brisbane/60/2008. Sequence analysis showed that these 18 viruses had a two-amino acid deletion in the hemagglutinin (HA) gene. Among the 537 B(Yamagata) viruses, all were antigenically similar to a B/Phuket/3073/2013(Yamagata lineage)-like virus, the WHO-recommended influenza B component for the 2017-18 northern hemisphere quadrivalent influenza vaccine containing two influenza B strains.

National Microbiology Laboratory (NML): Antiviral Resistance

From September 1, 2017 to February 15, 2018, the NML received influenza viruses from Canadian laboratories for drug susceptibility testing.

Amantadine: Of the 866 influenza A viruses [792 A(H3N2) and 74 A(H1N1)pdm09] tested against amantadine, all were resistant.

Oseltamivir: Of the 736 influenza viruses [390 A(H3N2), 60 A(H1N1)pdm09, and 286 B] tested against oseltamivir, all were sensitive except one A(H1N1)pdm09 virus with a H275Y mutation which was resistant.

Zanamivir: Of the 732 influenza viruses [386 A(H3N2), 60 A(H1N1)pdm09, and 286 B] tested against zanamivir, all were sensitive except one B virus which was resistant.

Mid-season 2017-18 Vaccine Effectiveness Estimates

Canada

On February 1, 2018, Canadian researchers published the first estimates of mid-season influenza vaccine effectiveness (VE) for the 2017-18 season. The 2017-18 season to date in Canada has been characterized by an equal mix of influenza A (49%) and influenza B (51%) viruses, the latter being unusual so early in the season. About two-thirds of participants contributing to VE analyses were working-age adults 20–64 years old.

Adjusted VE against A(H3N2), driven by a single genetic subgroup of clade 3C.2a, was low at 17% (95%CI: -14 to 40%) overall and 10% (95%CI: -31 to 39%) in working-age adults. Higher adjusted VE was observed for influenza B at 55% (95%CI: 38 to 68%) overall and 40% (95%CI: 16 to 67) in working-age adults, despite prominent use of lineage-mismatched B(Victoria) trivalent vaccine in most regions against circulating viruses that belonged to the B(Yamagata) lineage.

The full report is available as an open-access publication from *EuroSurveillance*:
<http://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2018.23.5.18-00035>

United States

On February 15, 2018, the US CDC published interim estimates of influenza vaccine effectiveness (VE) for the 2017-18 season. In contrast to the mixed circulation of influenza A(H3N2) and B viruses in Canada, the 2017-18 season in the United States has been characterized by early and widespread influenza activity of predominantly influenza subtype A(H3N2).

Adjusted VE against A(H3N2) was 25% (95% CI: 13 to 36%), comparable to Canadian estimates with both suggesting low protection against the dominant circulating strain. This estimate was driven by young children (6 months–8 years) who comprised about one-quarter of US study participants and for whom adjusted VE was 51% (95% CI: 29 to 66%). Adjusted VE in working-age adults (18–49 years) comprising about one-third of participants was non-significant at 20% (95% CI: -4 to 38%) and was even lower in adults 50-64 years old at 12% (-26 to 39%). Adjusted VE against influenza B was 42% (95% CI: 25 to 56%), somewhat lower than previous Canadian findings despite the more prominent use of quadrivalent vaccines containing both the B(Yamagata) and B(Victoria) lineages in the US.

The full report is available as an open-access publication from *Morbidity and Mortality Weekly Report (MMWR)*: https://www.cdc.gov/mmwr/volumes/67/wr/mm6706a2.htm?s_cid=mm6706a2_e

Spain (Navarre)

On February 15, 2018, Spanish researchers published interim estimates of influenza vaccine effectiveness (VE) for the 2017-18 season. The 2017/18 season in Europe has been characterised by co-circulation of influenza B, A(H3N2), and A(H1N1)pdm09. Unlike the Canadian and American studies which included only primary healthcare patients, researchers included both primary healthcare patients and hospitalized patients; this may impede comparison between these studies. Findings were driven by influenza B which comprised more than three-quarters of all influenza detections. There were too few A(H3N2) cases (118 in total, 43 from the outpatient setting) to reliably compare with other studies.

Despite exclusive use of trivalent influenza vaccine containing lineage-mismatched influenza B(Victoria) antigen, the adjusted VE against influenza B that was predominantly B(Yamagata) was 52% (95% CI: 12 to 74%) in the outpatient setting. This finding suggests moderate, cross-lineage protection against influenza B, which has been observed previously for influenza B and is consistent with Canadian findings.

The full report is available as an open-access publication from *Eurosurveillance*:
<https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2018.23.7.18-00057>

International

USA (week 5, January 28 to February 3, 2018)

During week 5, overall influenza activity increased in the United States. The most frequently identified influenza virus subtype reported by public health laboratories during week 5 was influenza A(H3N2). The percentage of respiratory specimens testing positive for influenza in clinical laboratories remained elevated. The proportion of deaths attributed to pneumonia and influenza (P&I) was above the system-specific epidemic threshold in the National Center for Health Statistics (NCHS) Mortality Surveillance System. Ten influenza-associated pediatric deaths were reported. A cumulative rate of 59.9 laboratory-confirmed influenza-associated hospitalizations per 100,000 population was reported. The proportion of outpatient visits for influenza-like illness (ILI) was 7.7%, which is above the national baseline of 2.2%. The geographic spread of influenza in Puerto Rico and 48 states was reported as widespread; two states reported regional activity; the District of Columbia and Guam reported local activity; and the U.S. Virgin Islands reported sporadic activity. Details are available at: www.cdc.gov/flu/weekly/.

WHO

There have been no WHO influenza updates since our last bulletin. Previous updates are available at: www.who.int/influenza/surveillance_monitoring/updates/en/.

Hong Kong

Recent elevated influenza B activity in Hong Kong has resulted in the precautionary closure of schools in advance of the Lunar New Year holiday due to a high volume of influenza cases among school-aged children. The holiday may serve as a source of amplification through social networks as people travel and get together to celebrate. As such, the BCCDC recommends that individuals who intend to travel to East Asia during the Lunar New Year period take precautions such as washing your hand regularly, eating well, exercising and making sure their vaccinations are up to date. Patients who become ill upon returning to Canada and require medical attention should inform their healthcare provider of recent travel.

WHO Recommendations for Influenza Vaccines

WHO Recommendations for 2017-18 Northern Hemisphere Influenza Vaccine

On March 2, 2017, the WHO announced the recommended strain components for the 2017-18 northern hemisphere trivalent influenza vaccine (TIV):*

- an A/Michigan/45/2015 (H1N1)pdm09-like virus;†
- an A/Hong Kong/4801/2014 (H3N2)-like virus;
- a B/Brisbane/60/2008 (Victoria-lineage)-like virus.

It is recommended that quadrivalent influenza vaccines (QIV) containing two influenza B viruses contain the above three viruses and a B/Phuket/3073/2013 (Yamagata-lineage)-like virus.

* These recommended strains are the same as those recommended for the 2017 southern hemisphere TIV and represent a change for one of the three components used for the 2016-17 northern hemisphere TIV and 2016 southern hemisphere TIV.

† Recommended strain represents a change from an A/California/7/2009-like virus, which had been retained as the A(H1N1)pdm09 component since the 2009 pandemic, to an A/Michigan/45/2015-like virus belonging to the emerging phylogenetic subclade 6B.1.

For further details: www.who.int/influenza/vaccines/virus/recommendations/2017_18_north/en/.

WHO Recommendations for the 2018 Southern Hemisphere Influenza Vaccine

On September 28, 2017, the WHO announced recommended strain components for the 2018 southern hemisphere trivalent influenza vaccine (TIV):*

- an A/Michigan/45/2015 (H1N1)pdm09-like virus;†
- an A/Singapore/INFIMH-16-0019/2016 (H3N2)-like virus;‡
- a B/Phuket/3073/2013-like (Yamagata-lineage) virus.§

It is recommended that quadrivalent influenza vaccines (QIV) containing two influenza B viruses contain the above three viruses and a B/Brisbane/60/2008 (Victoria-lineage)-like virus.

* Recommended strains represent a change for two of the three components used for the 2017 southern hemisphere vaccines.

† Recommended strain is the same as recommended for the 2017 southern hemisphere and 2017-18 northern hemisphere vaccines. The A/Michigan/45/2015-like virus belongs to the emerging phylogenetic subclade 6B.1; it replaces the A/California/7/2009-like virus that had been retained as the previous A(H1N1) component since the 2009 pandemic.

‡ Recommended strain for the A(H3N2) component represents a phylogenetic clade-level change from a clade 3C.2a virus to a clade 3C.2a1 virus containing the amino acid substitution N121K in the HA which is found in the majority of recent A(H3N2) viruses.

§ Recommended strain for the influenza B component represents a lineage-level change from a B(Victoria)-lineage virus to a B(Yamagata)-lineage virus.

For further details: http://www.who.int/influenza/vaccines/virus/recommendations/2018_south/en/.

The European Centre for Disease Prevention and Control has also posted a useful summary of WHO recommendations for the 2018 southern hemisphere influenza season, including rationale, available at: <https://ecdc.europa.eu/en/news-events/who-recommendations-influenza-virus-vaccine-composition-2018-southern-hemisphere>

Additional Information

Explanatory Note:

The surveillance period for the 2017-18 influenza season is defined starting in week 40. Weeks 36-39 of the 2016-17 season are shown on graphs for comparison purposes.

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

Current AMMI Canada Guidelines on the Use of Antiviral Drugs for Influenza:

www.ammi.ca/Update/79.ENG.pdf

Web Sites:

BCCDC Emerging Respiratory Pathogen Updates:

www.bccdc.ca/health-professionals/data-reports/emerging-respiratory-virus-updates

Influenza Web Sites

Canada – Influenza surveillance (FluWatch): <https://www.canada.ca/en/public-health/services/diseases/flu-influenza/influenza-surveillance.html>

Washington State Flu Updates: <http://www.doh.wa.gov/portals/1/documents/5100/420-100-fluupdate.pdf>

USA Weekly Surveillance Reports: www.cdc.gov/flu/weekly/

Joint ECDC – WHO/Europe weekly influenza update (Flu News Europe): flunewseurope.org

WHO – Weekly Epidemiological Record: www.who.int/wer/en/

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

www.influenzacentre.org/

Australian Influenza Report:

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

Avian Influenza Web Sites

WHO – Influenza at the Human-Animal Interface: www.who.int/csr/disease/avian_influenza/en/

World Organization for Animal Health: www.oie.int/eng/en_index.htm

Contact Us:

Tel: (604) 707-2510

Fax: (604) 707-2516

Email: InfluenzaFieldEpi@bccdc.ca

Communicable Disease Prevention and Control Services (CDPACS)

BC Centre for Disease Control

655 West 12th Ave, Vancouver BC V5Z 4R4

Online: www.bccdc.ca/health-professionals/data-reports/influenza-surveillance-reports

Influenza-Like Illness (ILI) Outbreak Summary Report Form

Please complete and email to 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.

A	<u>Reporting Information</u> 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	<u>First Notification</u>
	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
Total		
With ILI		
Hospitalized		
Died		

C	<u>Update AND Outbreak Declared Over</u>
	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
Total		
With ILI		
Hospitalized		
Died		

D	<u>Laboratory Information</u>
	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