Influenza and Other Respiratory Viruses
Weekly Report
California Influenza Surveillance Program

Highlights (Week 52: December 22, 2019 – December 28, 2019)

Statewide Activity

- **Deaths:** 54* since Sept. 29, 2019
- **Outbreaks:** 16 since Sept. 29, 2019
- **Laboratory:** 27.7% flu positive
- **Outpatient ILI:** Above expected levels
- **Hospitalizations:** Above expected levels

*Influenza-coded deaths from death certificates
Click on images and links for more information

Regions with Elevated Activity

Key messages:

- Flu activity continues to increase in California.
- Flu B viruses are predominating in California, but A (H1)pdm09 viruses are also circulating.
- Haven’t gotten a flu shot yet? Get immunized now to protect you and your loved ones.
- Everyone >6 months of age needs a flu shot.
- Take action to stop the spread of flu: wash hands often, cover coughs and sneezes, and stay home when sick.

Note: This report includes data from many sources of influenza surveillance and it should be viewed as a preliminary “snapshot” of influenza activity for each surveillance week. Because data are preliminary, the information may be updated in later reports as additional data are received. These data should not be considered population-based or representative of all California public health jurisdictions.
A. Outpatient, Inpatient, and Death Data

1. Influenza Sentinel Providers

Sentinel providers (physicians, nurse practitioners, and physician assistants) situated throughout California report on a weekly basis the number of patients seen with influenza-like illness (ILI) and the total number of patients seen for any reason. ILI is defined as any illness with fever (≥100°F or 37.8°C) AND cough and/or sore throat (in the absence of a known cause other than influenza).

A total of 76 enrolled sentinel providers have reported data for Week 52. Based on available data, the percentage of visits for ILI during Week 52 was 4.7% compared to Week 51 (3.5%) and is above expected levels for this time of year (Figure 1).

Figure 1. Percentage of Influenza-like Illness Visits Among Patients Seen by California Sentinel Providers, 2015–2020

The seasonal baseline was calculated using a regression model applied to data from the previous five years. Two standard deviations above the seasonal baseline is the point at which the observed percentage of ILI is significantly higher than would be expected at that time of year.

2. Kaiser Permanente Hospitalization Data

Inpatients at Kaiser Permanente facilities with an admission diagnosis including the keywords “flu,” “influenza,” “pneumonia,” or variants of the keywords are defined as pneumonia and influenza (P&I)-related admissions. The number of P&I admissions is divided by the total number of hospital admissions occurring in the same time period to estimate the percentage of P&I admissions. Admissions for pregnancy, labor and delivery, birth, and outpatient procedures are excluded from the denominator.

The percentage of admissions for pneumonia and influenza (P&I) in Kaiser Permanente facilities in northern California during Week 52 was 6.7% compared to Week 51 (5.2%) and is above expected levels for this time of the year (Figure 2).
Figure 2. Percentage of P&I Admissions in Kaiser Permanente Northern California Hospitals, 2015–2020

The seasonal baseline was calculated using a regression model applied to data from the previous five years. Two standard deviations above the seasonal baseline and is the point at which the observed percentage of pneumonia and influenza hospitalizations in Kaiser Permanente hospitals in Northern California is significantly higher than would be expected at that time of the year.

The majority of admissions for pneumonia and influenza did not result in intensive care unit (ICU) admission or death; however, 292 ICU admissions and 155 deaths have occurred among persons with P&I admission diagnoses (Figure 3a). The majority of P&I admissions occurred among persons ≥65 years of age across all severity categories, especially among deaths (Figure 3b). Please note that pneumonia and influenza admissions serve as a proxy for influenza activity, but do not necessarily represent laboratory-confirmed influenza infections.
Figure 3. Number (a) and age group distribution (b) of non-ICU, ICU, and deaths associated with P&I Admissions in Kaiser Permanente Northern California hospitals, 2019–2020 season to date
3. Influenza-Associated Hospitalizations, California Emerging Infections Program

The California Emerging Infections Program (CEIP), Influenza Surveillance Network (FluSurv-NET) conducts population-based surveillance for laboratory-confirmed influenza-associated hospitalizations among patients of all ages in Alameda, Contra Costa, and San Francisco counties.

The incidence of influenza-associated hospitalizations per 100,000 population was higher in week 50 (1.52) compared to Week 49 (1.46) (Figure 4). Data for the most recent two weeks are not presented because results are still being collected and are likely to change.

![Figure 4. Incidence of Influenza Hospitalizations in CEIP Counties, 2017–2020](image)

4. Influenza Mortality Surveillance from Death Certificates

Deaths occurring in California among residents who had influenza noted in any cause of death field on the death certificate (text or coded) are defined as influenza-coded deaths. The percentage of influenza-coded deaths is calculated by dividing the number of influenza-coded deaths by the total number of all cause deaths during the same period. Influenza-coded deaths are not necessarily laboratory-confirmed and are an underestimate of all influenza-associated deaths.
During Week 52, 8 new influenza-coded deaths were identified. To date during the 2019–2020 influenza season, 54 influenza-coded deaths have been identified (Figure 5). The percentage of deaths coded as influenza during Week 52 was 0.5% compared to 0.1% during Week 51 (Figure 6).

**Figure 5. Number of Influenza-coded Deaths Identified from Death Certificates by Week of Death, 2019–2020 Season**

Note: Coding of deaths can be delayed by several weeks. Influenza-coded deaths will be included once enough information is available to identify them.
To date, more deaths have occurred among persons ≥65 years of age (64.8%) than among persons <65 years of age during the 2019–2020 influenza season; however, the percentage of deaths occurring among persons <65 years of age (35.2%) is consistent with other seasons during which influenza viruses other than influenza A (H3N2) have circulated in greater numbers, such as the 2015–2016 and 2018–2019 seasons (Figure 7).
Figure 7. Age Distribution of Influenza-coded Deaths Occurring in California among California Residents, 2015–2016 Season through 2019–2020 Seasons

<table>
<thead>
<tr>
<th>Influenza Season</th>
<th>&lt;18 years*</th>
<th>18-49 years</th>
<th>50-64 years</th>
<th>65+ years</th>
</tr>
</thead>
<tbody>
<tr>
<td>2015-2016</td>
<td>182</td>
<td>472</td>
<td>1342</td>
<td>35</td>
</tr>
<tr>
<td>2016-2017</td>
<td>75</td>
<td>64</td>
<td>219</td>
<td>13</td>
</tr>
<tr>
<td>2017-2018</td>
<td>50</td>
<td>33</td>
<td>83</td>
<td>2</td>
</tr>
<tr>
<td>2018-2019†</td>
<td>19</td>
<td>33</td>
<td>22</td>
<td>4</td>
</tr>
<tr>
<td>2019-2020</td>
<td>4</td>
<td>4</td>
<td>17</td>
<td>2</td>
</tr>
</tbody>
</table>

* Methods used to identify pediatric influenza-coded deaths on death certificates differ from those used to identify the influenza-associated pediatric deaths presented below.
† One death during the 2018–2019 influenza season has unknown age and is not included in the figure.
2016–2017 influenza season: October 2, 2016–September 30, 2017; influenza A (H3N2) predominant season
2017–2018 influenza season: October 1, 2017–September 29, 2018; influenza A (H3N2) predominant season
2018–2019 influenza season: September 30, 2018–September 28, 2019; mixed influenza A (H1N1)pdm09 and influenza A (H3N2) season
2019–2020 influenza season: September 29, 2019–September 26, 2020; influenza B (Victoria) viruses predominating, influenza A (H1N1)pdm09 viruses are also circulating

5. Laboratory-Confirmed Severe Influenza-associated Pediatric Deaths

Influenza-associated deaths in children <18 years of age are nationally notifiable. The weekly influenza report includes confirmed deaths formally reported to CDPH through December 28, 2019 (Week 52).

No laboratory-confirmed influenza-associated fatalities in a child <18 years of age were reported to CDPH during Week 52. To date, CDPH has received four reports of laboratory-confirmed influenza-associated deaths among children <18 years of age during the 2019–2020 influenza season.
B. Laboratory Update – Influenza

1. Respiratory Laboratory Network (RLN) and Clinical Sentinel Laboratory Surveillance Results

Laboratory surveillance for influenza and other respiratory viruses involves the use of data from clinical sentinel laboratories (hospital, academic, and private laboratories) and public health laboratories in the Respiratory Laboratory Network located throughout California. These laboratories report the number of laboratory-confirmed influenza and other respiratory virus detections and isolations on a weekly basis.

The overall percentage of influenza detections in clinical sentinel laboratories in Week 52 (27.7%) was lower than Week 51 (29.7%) (Figure 8). Additional details, including influenza typing and subtyping information from public health laboratories can be found in Figures 8 and 9 and Tables 1 and 2.

Neither the RLN nor CDPH-VRDL has identified any influenza viruses by polymerase chain reaction (PCR) that are suggestive of a novel influenza virus.

Figure 8. Percentage of Influenza Detections at Clinical Sentinel Laboratories, 2015–2020
Figure 9. Number of Influenza Detections by Type and Subtype Detected in the Respiratory Laboratory Network, 2019–2020

![Graph showing number of positive specimens by week and type/subtype for A(H1N1)pdm09, A(H3N2), A Not Subtyped, B/Victoria, B/Yamagata, B Not Lineage Typed.]

Table 1. Respiratory Specimens Testing Positive for Influenza — Clinical Sentinel Laboratories, Current Week and Season to Date

<table>
<thead>
<tr>
<th></th>
<th>Current Week Number</th>
<th>Current Week Percent</th>
<th>Season to Date Number</th>
<th>Season to Date Percent</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Number of Specimens Tested</strong></td>
<td>2,988</td>
<td></td>
<td>39,745</td>
<td></td>
</tr>
<tr>
<td><strong>Influenza Positive</strong></td>
<td>829</td>
<td>27.7</td>
<td>5,592</td>
<td>14.1</td>
</tr>
<tr>
<td><strong>A</strong></td>
<td>357</td>
<td>43.1*</td>
<td>1,989</td>
<td>35.6*</td>
</tr>
<tr>
<td><strong>B</strong></td>
<td>472</td>
<td>56.9*</td>
<td>3,603</td>
<td>64.4*</td>
</tr>
</tbody>
</table>

* Percent of specimens positive for influenza
Table 2. Respiratory Specimens Testing Positive for Influenza by Influenza Type and Subtype — Respiratory Laboratory Network, Current Week and Season to Date

<table>
<thead>
<tr>
<th>Influenza Positive</th>
<th>Current Week Number</th>
<th>Current Week Percent</th>
<th>Season to Date Number</th>
<th>Season to Date Percent</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>159</td>
<td>46.1*</td>
<td>804</td>
<td>47.3*</td>
</tr>
<tr>
<td>A (H1)pdm09</td>
<td>119</td>
<td>74.8†</td>
<td>620</td>
<td>77.1†</td>
</tr>
<tr>
<td>A (H3)</td>
<td>13</td>
<td>8.2†</td>
<td>147</td>
<td>18.3†</td>
</tr>
<tr>
<td>A, not subtyped</td>
<td>27</td>
<td>17.0†</td>
<td>37</td>
<td>4.6†</td>
</tr>
<tr>
<td>B</td>
<td>186</td>
<td>53.9*</td>
<td>895</td>
<td>52.7*</td>
</tr>
<tr>
<td>B Victoria</td>
<td>99</td>
<td>53.2‡</td>
<td>580</td>
<td>64.8‡</td>
</tr>
<tr>
<td>B Yamagata</td>
<td>2</td>
<td>1.1‡</td>
<td>24</td>
<td>2.7‡</td>
</tr>
<tr>
<td>B, not lineage typed</td>
<td>85</td>
<td>45.7‡</td>
<td>291</td>
<td>32.5‡</td>
</tr>
</tbody>
</table>

* Percent of specimens positive for influenza
† Percent of influenza A positives
‡ Percent of influenza B positives

2. Influenza Virus Characterization

Close monitoring of influenza viruses is required to better assess the potential impact on public health. CDC characterizes influenza viruses through one or more tests including genetic characterization by analyzing viral genome sequences, and antigenic characterization by hemagglutination inhibition (HI) assays and/or neutralization based Focus Reduction assays (FRA). These data are used to compare how similar, or well-inhibited, currently circulating influenza viruses are to the reference viruses. Antigenic and genetic characterization of circulating influenza viruses gives an indication of the influenza vaccines ability to induce an immune response against the wide array of influenza viruses that are co-circulating every season. However, annual vaccine effectiveness estimates are needed to determine how much protection was provided to the population by vaccination.

Twenty influenza A (H3N2) viruses from California have been genetically characterized to date during the 2019–2020 influenza season, of which three were antigenically characterized. All 20 influenza A (H3N2) viruses were members of the 3c.2a1 subclade. Of the three influenza A (H3N2) viruses antigenically characterized, two (66.7%) were well-inhibited by A/Kansas/14/2017-like (H3N2) antisera and one (33.3%) was poorly inhibited by A/Kansas/14/2017-like (H3N2) antisera (Table 2a).

Twenty-two influenza A (H1N1)pdm09 viruses from California have been genetically characterized to date during the 2019–2020 influenza season, of which four were antigenically characterized. All 22 influenza A (H1N1)pdm09 viruses were members of the 6B.1A clade. Of the four influenza A (H1N1)pdm09 viruses antigenically characterized, four (100%) were well-inhibited by A/Brisbane/02/2018-like (H1N1) antisera (Table 2b).
Thirty-six influenza B (Victoria) viruses from California have been genetically characterized to date during the 2019–2020 influenza season, of which seven were antigenically characterized. Four influenza B (Victoria) viruses were members of the V1A.1 subclade and 32 were members of the V1A.3 subclade. Of the seven influenza B (Victoria) viruses antigenically characterized, four (57.1%) were well-inhibited by B/Colorado/06/2017-like (Victoria) antisera and three (42.9%) were poorly inhibited by B/Colorado/06/2017-like (Victoria) antisera (Table 2c).

Two influenza B (Yamagata) viruses from California have been genetically characterized to date during the 2019–2020 influenza season, of which one was antigenically characterized. Both influenza B (Yamagata) viruses were members of the Y3 clade. The antigenically characterized virus was well inhibited by B/Phuket/3073/2013-like (Yamagata) antisera (Table 2d).

Table 2 a–d. Influenza virus antigenic characterization and genomic sequencing by influenza A subtype and influenza B lineage type — California, 2019–2020 influenza season

a. Influenza A (H3N2) viruses

<table>
<thead>
<tr>
<th>Antigenic Characterization</th>
<th>3C.2a1</th>
<th>3C.3a*</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Well inhibited by A/Kansas/14/2017*</td>
<td>2</td>
<td>0</td>
<td>2</td>
</tr>
<tr>
<td>Poorly inhibited by A/Kansas/14/2017*</td>
<td>1</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Not antigenically characterized</td>
<td>17</td>
<td>0</td>
<td>17</td>
</tr>
<tr>
<td><strong>Total A (H3N2)</strong></td>
<td>20</td>
<td>0</td>
<td>20</td>
</tr>
</tbody>
</table>

* A/Kansas/14/2017 is the influenza A (H3N2) component included in the 2019–2020 quadrivalent and trivalent influenza vaccines and is a member of the 3c.3a clade

b. Influenza A (H1N1)pdm09 viruses

<table>
<thead>
<tr>
<th>Antigenic Characterization</th>
<th>6B.1</th>
<th>6B.1A*</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Well inhibited by A/Brisbane/02/2018*</td>
<td>0</td>
<td>4</td>
<td>4</td>
</tr>
<tr>
<td>Poorly inhibited by A/Brisbane/02/2018*</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Not antigenically characterized</td>
<td>0</td>
<td>18</td>
<td>18</td>
</tr>
<tr>
<td><strong>Total A (H1N1)pdm09</strong></td>
<td>0</td>
<td>22</td>
<td>22</td>
</tr>
</tbody>
</table>

* A/Brisbane/02/2018 is the influenza A (H1N1)pdm09 component included in the 2019–2020 quadrivalent and trivalent influenza vaccines and is a member of the 6B.1A clade

c. Influenza B (Victoria) viruses

<table>
<thead>
<tr>
<th>Antigenic Characterization</th>
<th>V1A.1*</th>
<th>V1A.3</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Well inhibited by B/Colorado/06/2017*</td>
<td>0</td>
<td>4</td>
<td>4</td>
</tr>
<tr>
<td>Poorly inhibited by B/Colorado/06/2017*</td>
<td>0</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>Not antigenically characterized</td>
<td>4</td>
<td>25</td>
<td>29</td>
</tr>
<tr>
<td><strong>Total B (Victoria)</strong></td>
<td>4</td>
<td>32</td>
<td>36</td>
</tr>
</tbody>
</table>

* B/Colorado/06/2017-like is the influenza B (Victoria) lineage component included in the 2019–2020 quadrivalent and trivalent influenza vaccines and is a member of the V1A.1 subclade
d. Influenza B (Yamagata) viruses

<table>
<thead>
<tr>
<th>Antigenic Characterization</th>
<th>Y3*</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Well inhibited by B/Phuket/3073/2013*</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Poorly inhibited by B/Phuket/3073/2013*</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Not antigenically characterized</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td><strong>Total B (Yamagata)</strong></td>
<td>2</td>
<td>2</td>
</tr>
</tbody>
</table>

* B/Phuket/3073/2013 is the influenza B (Yamagata) component included in the 2019–2020 quadrivalent influenza vaccine and is a member of the Y3 clade

C. Influenza-Associated Outbreaks

Four laboratory-confirmed influenza outbreaks were reported to CDPH during Week 52. To date, 16 laboratory-confirmed influenza outbreaks have been reported to CDPH for the 2019–2020 season.

Figure 10. Number of Laboratory-Confirmed Influenza-Associated Outbreaks by Week of First Onset, 2018–2020

D. California Border Region Influenza Surveillance Network Data

The border influenza surveillance network is comprised of outpatient provider sentinel sites whose geographical coverage extends approximately 100 kilometers (60 miles)
north of the California-Baja California border and includes Imperial and San Diego Counties, as well as some parts of Riverside County.

1. Syndromic Surveillance Update
A total of 7 border region sentinel providers reported data during Week 52. The total number of patients screened by all sentinel sites for ILI during Week 52 was 7,553. Outpatient ILI activity was 3.6% in Week 52. ILI activity for the California border region during Week 52 was lower when compared to activity for the same week during the 2017–2018 season and higher when compared to activity for the same week during the 2018–2019 season (Figure 11). All influenza syndromic data summarized for the border region represent a subset of CDC influenza sentinel providers in California.

Figure 11. Percentage of Influenza-like Illness Visits among Patients Seen by Sentinel Providers — California Border Region, 2017–2020

2. Virologic Surveillance Update
During Week 52, 444 respiratory specimens were tested from border region sentinel clinical laboratories; of these, 114 (25.7%) tested positive for influenza (49 [43.0%] influenza A and 65 [57.0%] influenza B). Cumulatively this season, a total of 5,283 respiratory specimens were tested from border region sentinel clinical laboratories; of these, 871 (16.5%) tested positive for influenza (165 [18.9%] influenza A; 706 [81.1%] influenza B).

During Week 52, 83 influenza positive specimens were detected at border region RLN laboratories; of which, 29 (34.9%) were influenza A and 54 (65.1%) were influenza B. Of
the 29 specimens that tested positive for influenza A at RLN laboratories, 26 (89.7%) were subtyped as A (H1)pdm09 and three (10.3%) were subtyped as A (H3). Of the 54 specimens that tested positive for influenza B at RLN laboratories, one (1.9%) was lineage typed as B (Yamagata) and 53 (98.1%) were lineage typed as B (Victoria). Cumulatively this season, a total of 229 influenza positive specimens have been detected at border region RLN laboratories; of which, 90 (39.3%) were influenza A and 139 (60.7%) were influenza B. Of the 90 specimens that tested positive for influenza A at RLN laboratories, 67 (74.4%) were subtyped as A (H1)pdm09 and 23 (25.6%) were subtyped as A (H3). Of the 139 specimens that tested positive for influenza B, five (3.6%) were lineage typed as B (Yamagata) and 134 (96.4%) were lineage typed as B (Victoria).

Laboratory data summarized in Figure 12 include data from border region influenza clinical sentinel laboratories (percentage of specimens testing positive for influenza) as well as data from border region RLN laboratories (influenza type and subtype/lineage type).

Figure 12. Number of Influenza Detections by Type and Subtype Detected in Respiratory Laboratory Network Laboratories and the Percentage of Specimens Testing Positive at Clinical Sentinel Laboratories — California Border Region, 2019–2020
E. Other Respiratory Viruses

1. Laboratory-Confirmed Severe Respiratory Syncytial Virus Case Reports

Currently, as mandated under Section 2500 of the California Code of Regulations, deaths among children aged 0–4 years with laboratory-confirmed respiratory syncytial virus (RSV) are reportable to CDPH. The weekly influenza report includes confirmed deaths formally reported to CDPH through December 28, 2019 (Week 52).

No laboratory-confirmed RSV-associated deaths among children <5 years of age were reported to CDPH during Week 52. To date, CDPH has received no reports of laboratory-confirmed RSV-associated deaths among children <5 years of age during the 2019–2020 influenza season.

2. Other Respiratory Virus Laboratory Update

During Week 52, 2,708 specimens were tested for RSV and 373 (13.8%) were positive, which is higher compared to Week 51 (10.7%) (Figure 13). During Week 52, adenovirus, coronavirus, human metapneumovirus, parainfluenza, and rhinovirus/enterovirus activity increased (Figure 14).

Figure 13. Percentage of RSV Detections at Clinical Sentinel Laboratories, 2015–2020
Figure 14. Percentage of Other Respiratory Pathogen Detections at Clinical Sentinel Laboratories, 2019–2020
**Activity Levels:**

**No Activity:** No laboratory-confirmed cases of influenza and no reported increase in the number of cases of ILI.

**Sporadic:** Small numbers of laboratory-confirmed influenza cases or a single laboratory-confirmed influenza outbreak has been reported, but there is no increase in cases of ILI.

**Local:** Outbreaks of influenza or increases in ILI cases and recent laboratory-confirmed influenza in a single region of the state.

**Regional:** Outbreaks of influenza or increases in ILI and recent laboratory confirmed influenza in at least two but less than half the regions of the state with recent laboratory evidence of influenza in those regions.

**Widespread:** Outbreaks of influenza or increases in ILI cases and recent laboratory-confirmed influenza in at least half the regions of the state with recent laboratory evidence of influenza in the state.

**California Regions:**

**Northern:** Alpine, Amador, Butte, Colusa, Del Norte, El Dorado, Glenn, Humboldt, Lake, Lassen, Mendocino, Modoc, Nevada, Placer, Plumas, Sacramento, Shasta, Sierra, Siskiyou, Sutter, Tehama, Trinity, Yolo, and Yuba counties

**Bay Area:** Alameda, Contra Costa, Marin, Napa, Solano, San Francisco, San Mateo, Santa Clara, Santa Cruz, and Sonoma counties

**Central Valley:** Calaveras, Fresno, Inyo, Kings, Mono, Madera, Mariposa, Merced, Monterey, San Benito, San Joaquin, Stanislaus, Tulare, and Tuolumne counties

**Upper Southern:** Kern, Los Angeles, San Luis Obispo, Santa Barbara, and Ventura counties

**Lower Southern:** Imperial, Orange, Riverside, San Bernardino, and San Diego counties

For questions regarding influenza surveillance and reporting in California, please email InfluenzaSurveillance@cdph.ca.gov. This account is monitored daily by several epidemiologists.

To obtain additional information regarding influenza, please visit the CDPH influenza website (https://www.cdph.ca.gov/Programs/CID/DCDC/Pages/Immunization/Influenza.aspx).

A copy of the case report form for reporting any laboratory-confirmed influenza case that was either admitted to the ICU or died can be downloaded from the CDPH influenza website (https://www.cdph.ca.gov/Programs/CID/DCDC/Pages/Immunization/Influenza.aspx).

For information about national influenza activity, please visit the Centers for Disease Control and Prevention’s FluView (https://www.cdc.gov/flu/weekly/index.htm) and FluView Interactive (https://www.cdc.gov/flu/weekly/fluviewinteractive.htm) websites.