



Weekly U.S. Influenza Surveillance Report

Updated May 17, 2024



A Weekly Influenza Surveillance Report Prepared by the Influenza Division

Key Updates for Week 19, ending May 11, 2024

Seasonal influenza activity is low nationally.

Viruses

Clinical Lab

2.4% (Trend **♣**) positive for influenza this week

(/flu/weekly/index.htm#ClinicalLaboratories)

Public Health Lab

Influenza A(H1N1)pdm09, A(H3N2), and B viruses were all co-circulating this week.

(/flu/weekly/index.htm#PublicHealthLaborator

Virus Characterization

Genetic and antigenic characterization and antiviral susceptibility are summarized in this report.

(/flu/weekly/index.htm#VirusCharacterization)

Illness

Outpatient Respiratory Illness

2.0% (Trend →)

of visits to a health care provider this week were for respiratory illness

(below baseline).

(/flu/weekly/index.htm#ILINet)

Outpatient Respiratory Illness: Map

This week no jurisdictions experienced moderate, high, or very high activity.

(/flu/weekly/index.htm#ORIAM)

FluSurv-NET

80.6 per 100,000

cumulative hospitalization rate.

(/flu/weekly/index.htm#FluSurvNet)

NCHS Mortality

0.1% (Trend →)

of deaths attributed to influenza this week.

(/flu/weekly/index.htm#NCHSMortality)

Pediatric Deaths

3

influenza-associated deaths were reported this week for a total of 167 deaths this season.

(/flu/weekly/index.htm#PedMortality)

All data are preliminary and may change as more reports are received.

Directional arrows indicate changes between the current week and the previous week. Additional information (/flu/weekly/index.htm#Trends) on the arrows can be found at the bottom of this page.

A description of the CDC influenza surveillance system, including methodology and detailed descriptions of each data component is available on the surveillance methods (/flu/weekly/overview.htm) page.

Additional information on the current and previous influenza seasons for each surveillance component are available on FluView Interactive (/flu/weekly/fluviewinteractive.htm).

Key Points

- Seasonal influenza activity is low nationally.
- Nationally, outpatient respiratory illness remained stable and is below baseline for the sixth week in a row. All 10 HHS regions are below their region-specific baselines.
- During Week 19, of the 94 viruses reported by public health laboratories, 57 (60.6%) were influenza A and 37 (39.4%) were influenza B. Of the 43 influenza A viruses subtyped during Week 19, 17 (39.5%) were influenza A(H1N1)pdm09 and 26 (60.5%) were A(H3N2).
- Three influenza-associated pediatric deaths were reported to CDC during Week 19, bringing the season total to 167 pediatric deaths.
- CDC estimates that there have been at least 35 million illnesses, 390,000 hospitalizations, and 24,000 deaths from flu so far this season.
- CDC recommends that everyone 6 months and older get an annual flu vaccine as long as flu activity continues.²
- There also are prescription flu antiviral drugs that can treat flu illness; those should be started as early as possible and are especially important for higher risk patients.³
- Seasonal flu viruses are among several viruses contributing to respiratory disease activity. CDC is providing updated, integrated information (https://www.cdc.gov/respiratory-viruses/index.html) about COVID-19, flu, and RSV activity on a weekly basis.

U.S. Virologic Surveillance (https://www.cdc.gov/flu/weekly/overview.htm#LabSurveillance)

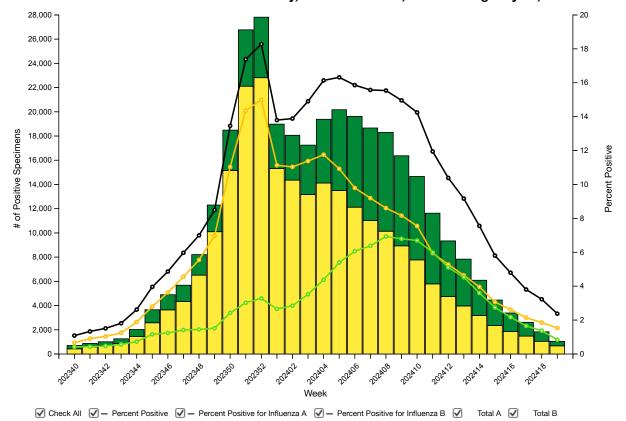
Nationally, the percentage of respiratory specimens testing positive for influenza in clinical laboratories decreased (change of >0.5 percentage points) compared to the previous week. Regions 1, 2, 3, 5, 6, 7, and 8 decreased in percent positivity while regions 4 and 9 remained stable and Region 10 increased slightly during Week 19 compared to Week 18. The regions with the highest percent positivity were regions 10 (4.4%), 9 (4.0%), and 7 (3.1%). Nationally, influenza A(H1N1)pdm09, A(H3N2), and B/Victoria viruses are all co-circulating. However, the distribution of circulating viruses varies by region. For regional and state level data and age group distribution, please visit FluView Interactive (https://gis.cdc.gov/grasp/fluview/fluportaldashboard.html). Viruses known to be associated with recent receipt of live attenuated influenza vaccine (LAIV) or found upon further testing to be a vaccine virus are not included, as they are not circulating influenza viruses.

Clinical Laboratories

The results of tests performed by clinical laboratories nationwide are summarized below. Data from clinical laboratories (the percentage of specimens tested that are positive for influenza virus) are used to monitor whether influenza activity is increasing or decreasing.

	Week 19	Data Cumulative since October 1, 2023 (Week 40)
No. of specimens tested	44,071	3,208,299
No. of positive specimens (%)	1,048 (2.4%)	343,365 (10.7%)
Positive specimens by type		
Influenza A	676 (64.5%)	236,845 (69.0%)
Influenza B	372 (35.5%)	106,509 (31.0%)

Influenza Positive Tests Reported to CDC by Clinical Laboratories, National Summary, 2023-24 Season, week ending May 11, 2024



View Additional Graphs and Data (http://gis.cdc.gov/grasp/fluview/fluportaldashboard.html) | Download Chart Data | Download PowerPoint Presentation

Public Health Laboratories

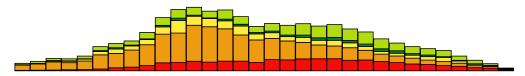
The results of tests performed by public health laboratories nationwide are summarized below. Data from public health laboratories are used to monitor the proportion of circulating influenza viruses that belong to each influenza subtype/lineage.

	Week 19	Data Cumulative since October 1, 2023 (Week 40)
No. of specimens tested	796	107,016
No. of positive specimens	94	35,385
Positive specimens by type/subtype		
Influenza A	57 (60.6%)	26,958 (76.2%)
Subtyping Performed	43 (75.4%)	22,327 (82.8%)

(H1N1)pdm09	17 (39.5%)	15,333 (68.7%)
H3N2	26 (60.5%)	6,994 (31.3%)
H3N2v	0 (0.0%)	0 (0.0%)
Subtyping not performed	14 (24.6%)	4,631 (17.2%)
Influenza B	37 (39.4%)	8,427 (23.8%)
Lineage testing performed	13 (35.1%)	6,994 (83.0%)
Yamagata lineage	0 (0.0%)	0 (0.0%)
Victoria lineage	13 (100.0%)	6,994 (100.0%)
Lineage not performed	24 (64.9%)	1,433 (17.0%)



Influenza Positive Tests Reported to CDC by Public Health Laboratories, National Summary, 2023-24 Season, week ending May 11, 2024





t of Positive Specime

Additional virologic surveillance information for current and past seasons:

Surveillance Methods (/flu/weekly/overview.htm#LabSurveillance) | FluView Interactive: National, Regional, and State Data (http://gis.cdc.gov/grasp/fluview/fluportaldashboard.html) or Age Data (https://gis.cdc.gov/grasp/fluview/flu_by_age_virus.html)

Influenza Virus Characterization (/flu/weekly/overview.htm#VirusCharacterization)

CDC performs genetic (https://www.cdc.gov/flu/about/professionals/genetic-characterization.htm) and antigenic

(https://www.cdc.gov/flu/about/professionals/antigenic.htm) characterization of U.S. viruses submitted from state and local public health laboratories according to the Right Size Roadmap submission guidance. These data are used to compare how similar the currently circulating influenza viruses are to the reference viruses representing viruses contained in the current influenza vaccines. The data are also used to monitor evolutionary changes that continually occur in influenza viruses circulating in humans. CDC also tests susceptibility of circulating influenza viruses to antiviral medications including the neuraminidase inhibitors (oseltamivir, zanamivir, and peramivir) and the PA endonuclease inhibitor baloxavir.

CDC has genetically characterized 4,374 influenza viruses collected since October 1, 2023.

	Genetic Characterization					
Virus Subtype or Lineage	Total No. of Subtype/Lineage Tested	HA Clade	Number (% of subtype/lineage tested)	HA Subclade	Number (% of subtype/lineage tested)	
A/H1	1,673					
		6B.1A.5a	1,673 (100%)	2a	387 (23.1%)	
				2a.1	1,286 (76.9%)	
A/H3	1,466					
		3C.2a1b.2a	1,466 (100%)	2a.1b	1 (0.1%)	
				2a.3a	1 (0.1%)	
				2a.3a.1	1,463 (99.8%)	
				2b	1 (0.1%)	
B/Victoria	1,235					
		V1A	1,235 (100%)	3a.2	1,235 (100%)	
B/Yamagata	0					
		Y3	0	Y3	0 (0%)	

CDC antigenically characterizes (https://www.cdc.gov/flu/about/professionals/antigenic.htm) influenza viruses by hemagglutination inhibition (HI) (H1N1pdm09, H3N2, B/Victoria, and B/Yamagata viruses) or neutralization-based HINT (https://pubmed.ncbi.nlm.nih.gov/30804469/) (H3N2 viruses) using antisera that ferrets make after being infected with reference viruses representing the 2023-2024 Northern Hemisphere recommended cell or recombinant-based vaccine viruses. Antigenic differences between viruses are determined by comparing how well the antibodies made against the vaccine reference viruses recognize the circulating viruses that have been grown in cell culture. Ferret antisera are useful because antibodies raised against a particular virus can often recognize small changes in the surface proteins of other viruses. In HI assays, viruses with similar antigenic properties have antibody titer differences of less than or equal to 4-fold when compared to the reference (vaccine) virus. In HINT, viruses with similar antigenic properties have antibody neutralization titer differences of less than or equal to 8-fold. Viruses selected for antigenic characterization are a subset representing the genetic changes in the surface proteins seen in genetically characterized viruses.

Influenza A Viruses

- A (H1N1)pdm09: 387 A(H1N1)pdm09 viruses were antigenically characterized by HI, and all were well-recognized (reacting at titers that were within 4-fold of the homologous virus titer) by ferret antisera to cell-grown A/Wisconsin/67/2022-like reference viruses representing the A(H1N1)pdm09 component for the cell- and recombinant-based influenza vaccines.
- A (H3N2): 371 A(H3N2) viruses were antigenically characterized by HI or HINT, and 364 (98.1%) were well-recognized (reacting at titers that were within 4-fold of the homologous virus titer in HI or reacting at titers that were less than or equal to 8-fold of the homologous virus in HINT) by ferret antisera to cell-grown A/Darwin/6/2021-like reference viruses representing the A(H3N2) component for the cell- and recombinant-based influenza vaccines.

Influenza B Viruses

- B/Victoria: 264 influenza B/Victoria-lineage virus were antigenically characterized by HI, and all were well-recognized (reacting at titers that were within 4-fold of the homologous virus titer) by ferret antisera to cell-grown B/Austria/1359417/2021-like reference viruses representing the B/Victoria component for the cell- and recombinant-based influenza vaccines.
- B/Yamagata: No influenza B/Yamagata-lineage viruses were available for antigenic characterization.

Assessment of Virus Susceptibility to Antiviral Medications

CDC assesses susceptibility of influenza viruses to the antiviral medications including the neuraminidase inhibitors (oseltamivir, zanamivir, and peramivir) and the PA endonuclease inhibitor baloxavir using next generation sequence analysis supplemented by laboratory assays. Information about antiviral susceptibility test methods can be found at U.S. Influenza Surveillance: Purpose and (https://www.cdc.gov/flu/weekly/overview.htm) Methods | CDC (https://www.cdc.gov/flu/weekly/overview.htm).

Viruses collected in the U.S. since October 1, 2023, were tested for antiviral susceptibility as follows:

Antiviral Medication		Total Viruses	A/H1	A/H3	B/Victoria	
Neuraminidase Inhibitors	Oseltamivir	Viruses Tested	4,279	1,648	1,438	1,193
		Reduced Inhibition	1 (0.02%)	1 (0.06%)	0 (0.00%)	0 (0.00%)
		Highly Reduced Inhibition	2 (0.05%)	2 (0.1%)	0 (0.00%)	0 (0.00%)
	Peramivir	Viruses Tested	4,279	1,648	1,438	1,193
Zanamivi		Reduced Inhibition	0 (0.00%)	0 (0.00%)	0 (0.00%)	0 (0.00%)
		Highly Reduced Inhibition	2 (0.05%)	2 (0.1%)	0 (0.00%)	0 (0.00%)
	Zanamivir	Viruses Tested	4,279	1,648	1,438	1,193
		Reduced Inhibition	0 (0.00%)	0 (0.00%)	0 (0.00%)	0 (0.00%)
	Highly Reduced Inhibition	0 (0.00%)	0 (0.00%)	0 (0.00%)	0 (0.00%)	
PA Cap- Dependent	Т	Viruses Tested	4,155	1,594	1,399	1,162
Endonuclease Inhibitor		Decreased Susceptibility	1 (0.02%)	0 (0.0%)	1 (0.1%)	0 (0.0%)

Two A(H1N1)pdm09 viruses had NA-H275Y amino acid substitution conferring highly reduced inhibition by oseltamivir and peramivir. One (H1N1)pdm09 virus had NA-I223V and NA-S247N amino acid substitutions and showed reduced inhibition by oseltamivir. One A(H3N2) virus had PA-I38T amino acid substitution and showed reduced susceptibility to baloxavir.

High levels of resistance to the adamantanes (amantadine and rimantadine) persist among influenza A(H1N1)pdm09 and influenza A(H3N2) viruses (the adamantanes are not effective against influenza B viruses). Therefore, use of these antivirals for treatment and prevention of influenza A virus infection is not recommended and data from adamantane resistance testing are not presented.

Outpatient Respiratory Illness Surveillance (https://www.cdc.gov/flu/weekly/overview.htm#ILINet)

The U.S. Outpatient Influenza-like Illness Surveillance Network (ILINet) monitors outpatient visits for respiratory illness referred to as influenza-like illness [ILI (fever plus cough or sore throat)], not laboratory-confirmed influenza and will therefore capture respiratory illness visits due to infection with pathogens that can present with similar symptoms, including influenza viruses, SARS-CoV-2, and RSV. It is important to evaluate syndromic surveillance data, including that from ILINet, in the context of other sources of surveillance data to obtain a more complete and accurate picture of influenza, SARS-CoV-2, and other respiratory virus activity. CDC is providing integrated information about COVID-19, influenza, and RSV activity on a website (https://www.cdc.gov/respiratory-viruses/index.html) that is updated weekly. Information about other respiratory virus activity can be found on CDC's National Respiratory and Enteric Virus Surveillance System (NREVSS) website (https://www.cdc.gov/surveillance/nrevss/index.html).

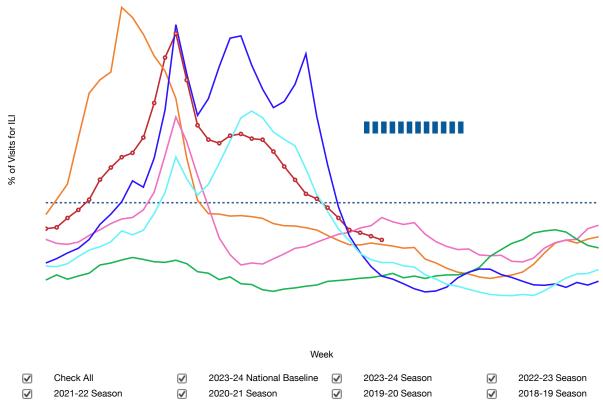
Outpatient Respiratory Illness Visits

Nationally, during Week 19, 2.0% of patient visits reported through ILINet were due to respiratory illness that included fever plus a cough or sore throat, also referred to as ILI. This has remained stable (change of \leq 0.1 percentage points) since Week 18 and is below the national baseline. The percentage of visits for ILI decreased in regions 1 and 5 and remained stable in all other regions in Week 19 compared to Week 18. All 10 regions are below their respective baselines. Multiple respiratory viruses are co-circulating, and the relative contribution of influenza virus infection to ILI varies by location.

Season:

2023-24 and 5 previous seasons ▼

Percentage of Outpatient Visits for Respiratory Illness Reported by The U.S. Outpatient Influenza-like Illness Surveillance Network (ILINet), Weekly National Summary, 2023-24 Season and Selected Previous Seasons

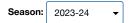


View National and Regional Level Graphs and Data (https://gis.cdc.gov/grasp/fluview/fluportaldashboard.html) | Download Chart Data | Download PowerPoint Presentation

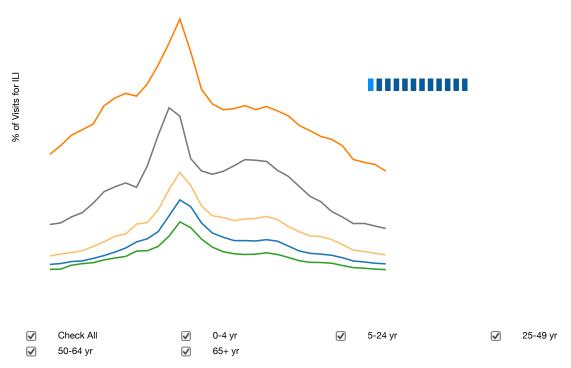
Outpatient Respiratory Illness Visits by Age Group

About 70% of ILINet participants provide both the number of patient visits for respiratory illness and the total number of patient visits for the week broken out by age group. Data from this subset of providers are used to calculate the percentages of patient visits for respiratory illness by age group.

The percentage of visits for respiratory illness reported in ILINet decreased in the 0-4 years and 5-24 years age groups and remained stable in all other age groups in Week 19 compared to Week 18.



Percent of Outpatient Visits for Respiratory Illness by Age Group Reported by the U.S. Outpatient Influenza-like Illness Surveillance Network (ILINet) Weekly National Summary, 2023-24 Influenza Season through the Week Ending May 11, 2024



View FluView Interactive (https://gis.cdc.gov/grasp/fluview/flu_by_age_virus.html) | Download Chart Data | Download PowerPoint Presentation

Outpatient Respiratory Illness Activity Map

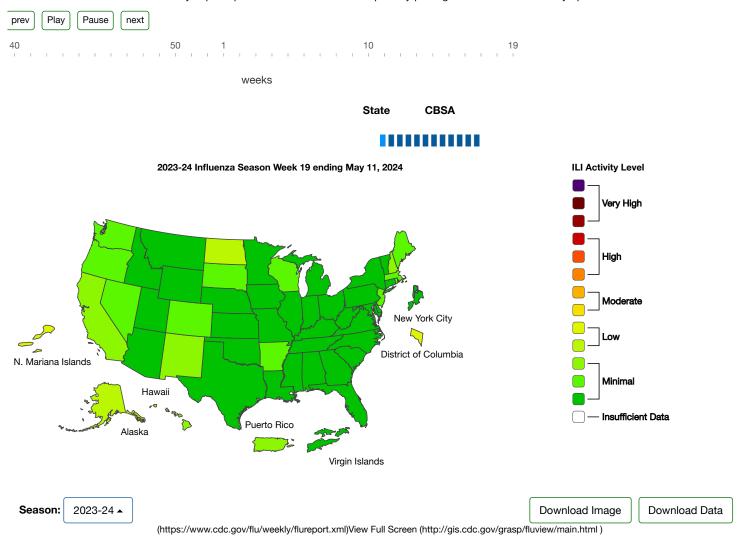
Data collected in ILINet are used to produce a measure of ILI activity* (/flu/weekly/overview.htm#ILINet) by state/jurisdiction and Core Based Statistical Areas (CBSA).

Number of Jurisdictions		Number of CBSAs	
Week 19	Week 18	Week 19	Week 18
(Week ending	(Week ending	(Week ending	(Week ending

Activity Level	May 11, 2024)	May 4, 2024)	May 11, 2024)	May 4, 2024)
Very High	0	0	0	0
High	0	0	1	1
Moderate	0	0	1	6
Low	4	4	36	33
Minimal	51	51	655	657
Insufficient Data	0	0	236	232

A Weekly Influenza Surveillance Report Prepared by the Influenza Division Outpatient Respiratory Illness Activity Map Determined by Data Reported to ILINet

This system monitors visits for respiratory illness that includes fever plus a cough or sore throat, also referred to as ILI, not laboratory confirmed influenza and may capture patient visits due to other respiratory pathogens that cause similar symptoms.



Additional information about medically attended visits for ILI for current and past seasons:

Surveillance Methods (/flu/weekly/overview.htm#ILINet) | FluView Interactive: National, Regional, and State Data (http://gis.cdc.gov/grasp/fluview/fluportaldashboard.html) or ILI Activity Map (https://gis.cdc.gov/grasp/fluview/main.html)

Hospitalization Surveillance (http://www.cdc.gov/flu/weekly/overview.htm#HospitalizationSurv)

^{*}Data collected in ILINet may disproportionally represent certain populations within a jurisdiction or CBSA, and therefore, may not accurately depict the full picture of influenza activity for the entire jurisdiction or CBSA. Differences in the data presented here by CDC and independently by some health departments likely represent differing levels of data completeness with data presented by the health department likely being the more complete.

FluSurv-NET

The Influenza Hospitalization Surveillance Network (FluSurv-NET) conducts population-based surveillance for laboratory-confirmed influenza-related hospitalizations in select counties in 14 states and represents approximately 9% of the U.S. population. FluSurv-NET hospitalization data are preliminary. As data are received each week, prior case counts and rates are updated accordingly.

A total of 24,643 laboratory-confirmed influenza-associated hospitalizations were reported by FluSurv-NET sites between October 1, 2023, and May 11, 2024. The weekly hospitalization rate observed in Week 19 was 0.2 per 100,000 population. The weekly hospitalization rate observed during Week 52 is tied with the 2014-2015 season for the second highest peak weekly rate observed during all seasons going back to 2010-2011 and only lower than the 2017-2018 season. The overall cumulative hospitalization rate was 80.6 per 100,000 population.

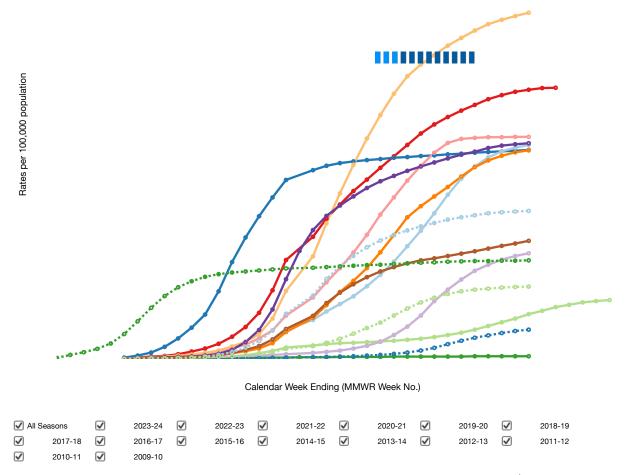
When examining rates by age, the highest cumulative hospitalization rate per 100,000 population was among adults aged 65 years and older (220.7) followed by adults aged 50-64 years (95.6) and children aged 0-4 years (82.9). When examining age-adjusted rates by race and ethnicity, the highest rate of hospitalization per 100,000 population was among non-Hispanic Black persons (146.7), followed by non-Hispanic American Indian or Alaska Native persons (111.9), Hispanic persons (76.6), non-Hispanic White persons (62.3), and non-Hispanic Asian/Pacific Islander persons (43.6).

Among 24,643 hospitalizations, 20,857 (84.6%) were associated with influenza A virus, 3,609 (14.7%) with influenza B virus, 48 (0.2%) with influenza A virus and influenza B virus co-infection, and 128 (0.5%) with influenza virus for which the type was not determined. Among those with influenza A subtype information, 3,563 (69.3%) were A(H1N1) pdm09 and 1,578 (30.7%) were A(H3N2).

Among 3,845 hospitalized adults with information on underlying medical conditions, 95.4% had at least one reported underlying medical condition, the most commonly reported were hypertension, cardiovascular disease, obesity, and metabolic disease. Among 1,582 hospitalized women of childbearing age (15-49 years) with information on pregnancy status, 22.9% were pregnant. Among 1,085 hospitalized children with information on underlying medical conditions, 67.7% had at least one reported underlying medical condition; the most commonly reported was asthma, followed by obesity and neurologic disease.

Laboratory-Confirmed Influenza Hospitalizations Preliminary cumulative rates as of May 11, 2024

FluSurv-NET :: Entire Network :: Overall Age Group :: Cumulative Rate



View National and Regional Level Graphs and Data (https://gis.cdc.gov/grasp/fluview/FluHospRates.html) | Download Chart Data | Download PowerPoint Presentation

In these figures, cumulative and weekly rates for all seasons prior to the 2023-2024 season reflect end-of-season rates. For the 2023-2024 season, rates for recent hospitals admissions are subject to reporting delays. As hospitalization data are reviewed each week, prior case counts and rates are updated accordingly.

Additional FluSurv-NET hospitalization surveillance information for current and past seasons and additional age groups:

Surveillance Methods (https://www.cdc.gov/flu/weekly/overview.htm#FluSurvNet) | FluView Interactive: Rates by Age, Sex, and Race/Ethnicity (http://gis.cdc.gov/GRASP/Fluview/FluHospRates.html) or Data on Patient Characteristics (http://gis.cdc.gov/grasp/fluview/FluHospChars.html) | RESP-NET Interactive (https://www.cdc.gov/surveillance/resp-net/dashboard.html)

National Healthcare Safety Network (NHSN) Hospitalization Surveillance

Effective May 1, 2024, hospitals are no longer required to report hospital admissions, hospital capacity, or hospital occupancy data to HHS through NHSN. Voluntarily reported NHSN hospital data can found at Weekly United States Hospitalization Metrics by Jurisdiction (https://data.cdc.gov/Public-Health-Surveillance/Weekly-United-States-Hospitalization-Metrics-by-Ju/aemt-mg7g/about_data).

Additional NHSN Hospitalization Surveillance information:

Surveillance Methods (https://www.cdc.gov/flu/weekly/overview.htm#NHSN) | Additional Data (https://data.cdc.gov/Public-Health-Surveillance/Weekly-United-States-Hospitalization-Metrics-by-Ju/aemt-mg7g/about_data) | FluView Interactive (http://gis.cdc.gov/grasp/fluview/FluView12.html)

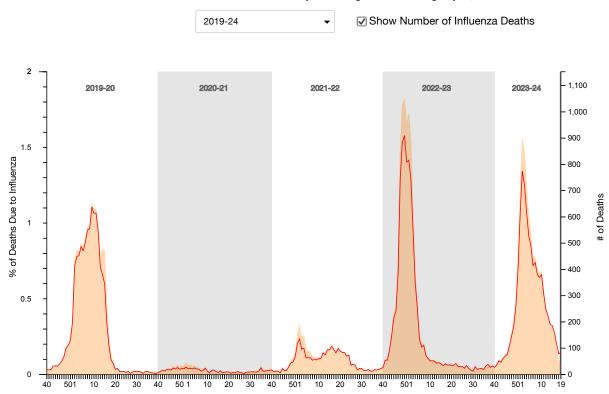
Mortality Surveillance (https://www.cdc.gov/flu/weekly/overview.htm#MortalitySurveillance)

National Center for Health Statistics (NCHS) Mortality Surveillance

Based on NCHS mortality surveillance data available on May 16, 2024, 0.1% of the deaths that occurred during the week ending May 11, 2024 (Week 19), were due to influenza. This percentage remained stable (< 0.1 percentage point change) compared to Week 18. The data presented are preliminary and may change as more data are received and processed.

Pneumonia and Influenza Mortality from the National Center for Health Statics Mortality Surveillance System

National Summary data through the week ending May 11, 2024



MMWR Week

Influenza Coded Deaths

View Regional and State Level Data (https://gis.cdc.gov/grasp/fluview/Mortality.html) | Download Chart Data | Download PowerPoint Presentation

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% of Deaths due to Influenza

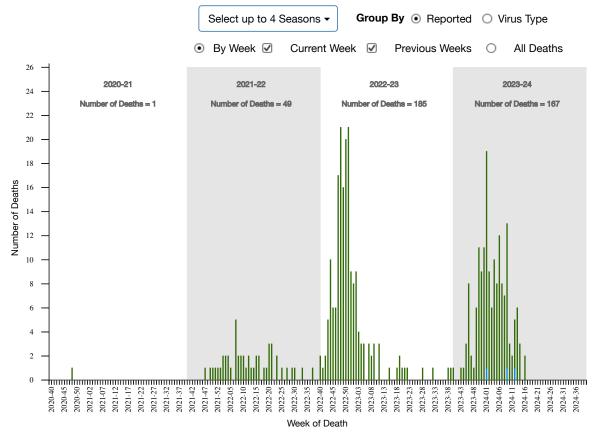
Additional pneumonia, influenza and COVID-19 mortality surveillance information for current and past seasons:

 $Surveillance \ Methods \ (https://www.cdc.gov/flu/weekly/overview.htm\#NCHSMortality) \ | \ FluView \ Interactive \ (https://gis.cdc.gov/grasp/fluview/mortality.html)$

Influenza-Associated Pediatric Mortality

Three influenza-associated pediatric deaths were reported to CDC during Week 19, bringing the total pediatric deaths for this season to 167. The deaths occurred during weeks 1, 9, and 12 of 2024 (the weeks ending January 6, March 2, and March 23, 2024). Two deaths were associated with influenza A viruses and one death was associated with an influenza B/Victoria virus. One of the influenza A viruses had subtyping performed and was an A(H3N2) virus.

Influenza-Associated Pediatric Deaths by Week of Death, 2020-21 season to 2023-24 season



View FluView Interactive (https://gis.cdc.gov/grasp/fluview/PedFluDeath.html) | Download Chart Data | Download PowerPoint Presentation

Additional pediatric mortality surveillance information for current and past seasons:

Surveillance Methods (https://www.cdc.gov/flu/weekly/overview.htm#PediatricMortality) | FluView Interactive (https://gis.cdc.gov/GRASP/Fluview/PedFluDeath.html)

Trend Indicators

Increasing: Decreasing: Stable:

Indicators Status by System

Clinical Labs: Up or down arrows indicate a change of greater than or equal to 0.5 percentage points in the percent of specimens positive for influenza compared to the previous week.

Outpatient Respiratory Illness (ILINet): Up or down arrows indicate a change of greater than 0.1 percentage points in the percent of visits due to respiratory illness (ILI) compared to the previous week.

NHSN Hospitalizations: Up or down arrows indicate change of greater than or equal to 5% of the number of patients admitted with laboratory-confirmed influenza compared to the previous week.

NCHS Mortality: Up or down arrows indicate change of greater than 0.1 percentage points of the percent of deaths due to influenza compared to the previous week.

Reference Footnotes

¹U.S. Influenza Surveillance: Purpose and Methods (2023 Oct). Centers for Disease Control and Prevention. https://www.cdc.gov/flu/weekly/overview.htm#ILINet (https://www.cdc.gov/flu/weekly/overview.htm#ILINet).

²Grohskopf LA, Blanton LH, Ferdinands JM, Chung JR, Broder KR, Talbot HK. Prevention and Control of Seasonal Influenza with Vaccines: Recommendations of the Advisory Committee on Immunization Practices — United States, 2023–24 Influenza Season. MMWR Recomm Rep 2023;72(No. RR-2):1–25.

DOI: http://dx.doi.org/10.15585/mmwr.rr7202a1 [2] (http://dx.doi.org/10.15585/mmwr.rr7202a1)

³Influenza Antiviral Medications: Summary for Clinicians (2023 Sept). Centers for Disease Control and Prevention. https://www.cdc.gov/flu/professionals/antivirals/summary-clinicians.htm (https://www.cdc.gov/flu/professionals/antivirals/summary-clinicians.htm).

Additional National and International Influenza Surveillance Information

FluView Interactive: FluView includes enhanced web-based interactive applications that can provide dynamic visuals of the influenza data collected and analyzed by CDC. These FluView Interactive applications (http://www.cdc.gov/flu/weekly/fluviewinteractive.htm) allow people to create customized, visual interpretations of influenza data, as well as make comparisons across flu seasons, regions, age groups and a variety of other demographics.

National Institute for Occupational Safety and Health: Monthly surveillance data on the prevalence of health-related workplace absenteeism among full-time workers in the United States are available from NIOSH (https://www.cdc.gov/niosh/topics/absences/default.html).

U.S. State and local influenza surveillance: Select a jurisdiction below to access the latest local influenza information.

Alabama (http://adph.org/influenza/)	Alaska (http://dhss.alaska.gov/dph/Epi/id/Pages/influenza/fluinf
Colorado (https://www.mass.gov/info-details/influenza-reporting)	Connecticut (https://portal.ct.gov/DPH/Epidemiology- Emerging-Infections/Influenza-Surveillance-and-Statist
Georgia (https://dph.georgia.gov/flu-activity-georgia)	Hawaii

	(http://health.hawaii.gov/docd/resources/reports/influen: reports/)
lowa (https://idph.iowa.gov/influenza/reports)	Kansas (http://www.kdheks.gov/flu/surveillance.htm)
Maryland (https://phpa.health.maryland.gov/influenza/fluwatch/)	Massachusetts (https://www.mass.gov/infodetails/influenza-reporting)
Missouri	Montana
(http://health.mo.gov/living/healthcondiseases/communicable/influenza/reports.php)	(https://dphhs.mt.gov/publichealth/cdepi/diseases/Pan- Respiratory/index)
New Jersey (http://www.nj.gov/health/cd/topics/flu.shtml)	New Mexico (https://nmhealth.org/about/erd/ideb/isp
Ohio (http://www.flu.ohio.gov)	Oklahoma (https://oklahoma.gov/health/health-education/acute-disease-service/disease-information/influenza-home-page.html)
South Carolina	South Dakota (https://doh.sd.gov/health-data-
(http://www.scdhec.gov/Health/DiseasesandConditions/InfectiousDiseases/Flu/FluData/)	reports/data-dashboards/influenza-dashboard/)
Vermont (http://www.healthvermont.gov/immunizations-infectious-	Virginia
disease/influenza/flu-activity-and-surveillance)	(http://www.vdh.virginia.gov/epidemiology/influenza-fluvirginia/influenza-surveillance/)
Wyoming (https://health.wyo.gov/publichealth/infectious-disease-epidemiology-	New York City
unit/disease/influenza/)	(http://www1.nyc.gov/site/doh/providers/health-topics/f
	alerts.page)

World Health Organization:

Additional influenza surveillance information from participating WHO member nations is available through FluNet (https://www.who.int/tools/flunet) and the Global Epidemiology Reports. (https://www.who.int/teams/global-influenza-programme/surveillance-and-monitoring/influenza-surveillance-outputs)

WHO Collaborating Centers for Influenza:

Australia (http://www.influenzacentre.org/Surveillance_Samples_Received.html), China (http://www.chinaivdc.cn/cnic/), Japan (http://idsc.nih.go.jp/index.html), the United Kingdom (https://www.crick.ac.uk/research/worldwide-influenza-centre), and the United States (http://www.cdc.gov/flu/) (CDC in Atlanta, Georgia)

Europe:

The most up-to-date influenza information from Europe is available from WHO/Europe and the European Centre for Disease Prevention and Control (http://www.flunewseurope.org/).

Public Health Agency of Canada:

The most up-to-date influenza information from Canada is available in Canada's weekly FluWatch report (http://www.phac-aspc.gc.ca/fluwatch/).

Public Health England:

The most up-to-date influenza information from the United Kingdom is available from Public Health England (http://www.hpa.org.uk/Topics/InfectiousDiseases/InfectionsAZ/SeasonalInfluenza/).

Any links provided to non-Federal organizations are provided solely as a service to our users. These links do not constitute an endorsement of these organizations or their programs by CDC or the Federal Government, and none should be inferred. CDC is not responsible for the content of the individual organization web pages found at these links.

A description of the CDC influenza surveillance system, including methodology and detailed descriptions of each data component is available on the surveillance methods (http://www.cdc.gov/flu/weekly/overview.htm) page.

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Source: Centers for Disease Control and Prevention (https://www.cdc.gov/), National Center for Immunization and Respiratory Diseases (NCIRD) (https://www.cdc.gov/ncird/index.html)

Was this page helpful?

Yes

Partly

No