



MORBIDITY AND MORTALITY WEEKLY REPORT

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World AIDS Day — December 1, 1996

"One World, One Hope" is the theme designated by the Joint United Nations Programme on HIV/AIDS (UNAIDS) for this year's World AIDS Day, December 1, 1996. Worldwide, 190 countries observe World AIDS Day to focus attention on the human immunodeficiency virus (HIV)/acquired immunodeficiency syndrome (AIDS) pandemic, which has resulted in an estimated 27.9 million HIV infections and 5.8 million deaths, including 1.3 million deaths in children (1,2). In the United States, activities for World AIDS Day are coordinated by the American Association for World Health in collaboration with UNAIDS, the Pan American Heath Organization, and the U.S. Department of Health and Human Services.

Additional information about HIV infection, AIDS, and World AIDS Day is available from the CDC National AIDS Hotline, telephone (800) 342-2437, (919) 361-8400, (800) 344-7432 (Spanish), and (800) 243-7889 (TTY/TDD); the CDC National AIDS Clearinghouse, telephone (800) 458-5231 or (301) 217-0023; and the CDC Home Page on the World Wide Web, http://www.cdc.gov/nchstp/hiv_aids/worldaid/worldaid.htm.

References

- Joint United Nations Programme on HIV/AIDS. The HIV/AIDS situation in mid 1996: global and regional highlights. Fact Sheet. Geneva, Switzerland: World Health Organization, July 1996
- 2. American Association for World Health. Resource Booklet. World AIDS Day—1 December 1996, One World One Hope. Washington DC: American Association for World Health, 1996.

AIDS Among Children — United States, 1996

As of September 30, 1996, a total of 566,002 acquired immunodeficiency syndrome (AIDS) cases, including 7472 cases among children aged <13 years (1%), had been reported to CDC by state and territorial health departments. Most children reported with AIDS acquired human immunodeficiency virus (HIV) infection perinatally from their mothers (1). During 1988–1993, an estimated 6000–7000 children were born each year to HIV-infected women; an estimated 1000–2000 of these children were infected annually (2). In 1994, results of clinical trials demonstrating effective therapy for reducing perinatal HIV transmission indicated a two-thirds decrease in such transmis-

sion associated with zidovudine (ZDV) therapy for HIV-infected pregnant women and their newborns. The Public Health Service (PHS) issued recommendations in 1994 for ZDV treatment to reduce perinatal HIV transmission, and in 1995 for routine HIV counseling and voluntary testing for all pregnant women in the United States (3,4). This report summarizes the epidemiology of AIDS in children in the United States reported cumulatively from 1982 through September 1996, presents rates for 1995 (the most recent year for which census estimates are available), and describes a recent decrease in the rate of perinatally acquired AIDS.*

AIDS Among Children

Of the 7472 children reported with AIDS, 58% were non-Hispanic black, 23% were Hispanic, 18% were non-Hispanic white, and 1% were of other racial/ethnic groups. During 1995, the rates of reported AIDS cases per 100,000 children were 6.4 for non-Hispanic blacks, 2.3 for Hispanics, 0.4 for non-Hispanic whites, 0.4 for American Indians/Alaskan Natives, and 0.3 for Asians/Pacific Islanders. Among all U.S. children with AIDS, 6750 (90%) acquired HIV perinatally, 370 (5%) through receipt of contaminated blood transfusions, and 231 (3%) through receipt of contaminated blood products for coagulation disorders; 121 (2%) had no reported risk factor. Among children with perinatally acquired AIDS, the median age at diagnosis was 18 months. Approximately 80% of all children with AIDS had AIDS diagnosed before age 5 years. The highest numbers of cases were reported from New York (1901), Florida (1199), New Jersey (661), California (524), Puerto Rico (347), and Texas (296); combined, these cases accounted for 66% of all AIDS cases reported among children.

Risk exposures for HIV infection among the mothers of the 6750 children with perinatally acquired AIDS included injecting-drug use (IDU) (41%), sexual contact with a partner with or at risk for HIV/AIDS (34%), and receipt of contaminated blood or blood products (2%); for 13%, no risk was specified.

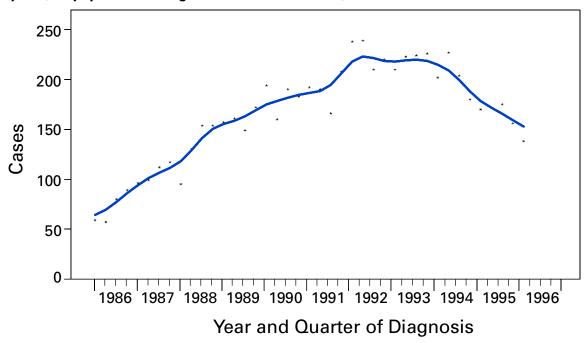
Trends in Perinatally Acquired AIDS

To examine trends in the incidence of AIDS among children born to HIV-infected mothers, the number of perinatally acquired AIDS cases diagnosed each quarter from 1986 through March 1996 was estimated using standard statistical adjustments that account for delays in reporting cases to CDC and estimates of behavioral risk among persons reported without a risk (1). The estimated number of children with perinatally acquired AIDS peaked at 905 during 1992, followed by a decline in incidence (Figure 1).

From 1992 through 1995, the estimated annual number of perinatally acquired AIDS cases declined 27%, from 905 to 663. During this time, the estimated annual number of cases declined 39% among non-Hispanic white, 26% among non-Hispanic black, and 25% among Hispanic children. The proportionate decrease in the number of children with perinatally acquired AIDS from the six areas reporting the highest number of cases was greater than the decrease for all remaining areas and for all areas combined (Table 1).

^{*}Single copies of this report will be available until November 22, 1997, from the CDC National AIDS Clearinghouse, P.O. Box 6003, Rockville, MD 20849-6003; telephone (800) 458-5231 or (301) 217-0023.

FIGURE 1. Number of perinatally acquired AIDS cases among children aged <13 years,* by quarter of diagnosis — United States, 1986–March 1996



^{*}Estimates were based on cases reported through September 1996, adjusted for reporting delays and unreported risk but not for incomplete reporting of diagnosed AIDS cases. Points represent estimated quarterly incidence, and the line represents "smoothed" incidence.

TABLE 1. Estimated number of children diagnosed with perinatally acquired AIDS*, by area of residence, year of diagnosis, and percentage change, 1992 to 1995 — United States and territories

Area of		N	lo.		% Change
residence	1992	1993	1994	1995	1992 to 1995
Top six areas [†]	583	562	509	398	-32%
All others	322	306	291	265	-18%
Total	905	868	800	663	-27%

^{*}Cases diagnosed through 1995 using data reported to CDC through September 1996, adjusted for reporting delays and unreported risk. Estimates are not adjusted for incomplete reporting of diagnosed AIDS cases.

HIV Infection Among Children

To enhance the usefulness of surveillance systems to characterize affected populations and to improve the targeting of resources for prevention and care, 28 states require confidential reporting of children with HIV infection without a diagnosis of AIDS as well as those with AIDS (1). Through September 1996, these states reported

Six areas reporting the highest number of cases: California, Florida, New Jersey, New York, Puerto Rico, and Texas.

TABLE 2. Number of children aged <13 years reported with HIV infection* and AIDS — United States and territories, 1995[†]

Area	HIV	AIDS	Area	HIV	AIDS
Alabama	8	4	Nebraska	2	1
Alaska		0	Nevada	1	4
Arizona	6	1	New Hampshire	_	0
Arkansas	8	3	New Jersey	48	61
California		89	New Mexico	_	0
Colorado	4	1	New York	_	166
Connecticut	12	18	North Carolina	25	12
Delaware		1	North Dakota	0	0
District of Columbia		13	Ohio	14	14
Florida		111	Oklahoma	3	0
Georgia		28	Oregon	_	2
Hawaii	_	0	Pennsylvania	_	19
Idaho	0	0	Puerto Rico	_	46
Illinois	_	26	Rhode Island	_	0
Indiana	4	3	South Carolina	24	7
Iowa		0	South Dakota	1	0
Kansas		2	Tennessee	12	10
Kentucky	_	1	Texas	51	31
Louisiana	23	12	Utah	0	0
Maine	_	1	Vermont	_	0
Maryland	_	37	Virginia	10	19
Massachusetts	_	19	Virgin Islands	_	5
Michigan	19	9	Washington	_	3
Minnesota	3	3	West Virginia	0	2
Mississippi	7	8	Wisconsin	7	0
Missouri	10	5	Wyoming	0	0
Montana	_	0	Total	302	797

^{*}Twenty-eight states reported children with HIV infection without a diagnosis of AIDS in addition to children with AIDS.

29% (2155) of all children with AIDS and 1447 children with HIV infection. During 1995, these states reported 228 AIDS cases among children and 302 children with documented HIV infection who had not developed AIDS (Table 2). During 1995, these states received 1464 additional reports of children who were born to HIV-infected mothers but who require follow-up with providers to determine their HIV-infection status. Among the six reporting areas with the highest cumulative number of children with AIDS, only New Jersey and Texas require reports of HIV infection among children. Reported by state, territorial, and local health departments. Div of HIV/AIDS Prevention, National Center for HIV, STD, and TB Prevention, CDC.

Editorial Note: The findings in this report document a decline in the incidence of perinatally acquired AIDS before and after the release of PHS recommendations for HIV counseling and voluntary testing for pregnant women and for ZDV therapy to prevent perinatal transmission (3,4). The recommendations were issued to promote the adoption of these HIV-prevention strategies as standard medical practice in the United States. Because the number of HIV-infected women who gave birth each year was stable during 1989–1994 (5), this decline suggests that the decrease in perinatal HIV

[†]Data reported to CDC through September 1996.

transmission rates probably reflected the effect of perinatal ZDV therapy. Increasing proportions of women may be accepting voluntary prenatal HIV testing and using ZDV to prevent perinatal transmission (6,7).

Because the incidence of perinatally acquired AIDS declined slightly before the PHS recommendations on ZDV therapy were issued in 1994, other factors may have contributed to the decrease in perinatally acquired AIDS cases during this period. For example, the proportion of HIV-infected childbearing women who received ZDV therapy before and during pregnancy for treatment of their HIV disease was increasing (8). Among children, increased use of prophylaxis to prevent AIDS opportunistic infections may have delayed the development of these conditions. However, the incidence of *Pneumocystis carinii* pneumonia, the most common AIDS-defining condition among children, has not decreased substantially among young children (9,10).

AIDS surveillance conducted in all reporting areas provides a standardized means to monitor AIDS incidence in children as a measure of the effectiveness of perinatal prevention efforts. To further characterize implementation of counseling, testing, and treatment for HIV-infected mothers and their children, CDC and other federal agencies are initiating facility-based program evaluations in selected high-incidence areas. These studies also will examine factors that may contribute to a change in perinatal HIV transmission rates (e.g., changing obstetrical practices and womens' attitudes toward and adherence to ZDV and other preventive therapy). In states that conduct confidential HIV reporting for children, timely assessment of HIV-prevention measures in mother-infant pairs (e.g., prenatal care and prenatal and neonatal ZDV therapy) will measure changes in perinatal HIV transmission rates statewide and permit refinement and redirection of prevention efforts. The Council of State and Territorial Epidemiologists has recommended that all states implement HIV infection reporting for children and consider reporting of all children of indeterminate HIV status who were born to infected mothers.

In the United States, HIV and AIDS disproportionately affect non-Hispanic black and Hispanic women and their children. This disparity probably reflects socioeconomic factors, access to and use of medical services, or differences in behaviors associated with HIV transmission risks among women. Health-care providers in the public and private sectors should implement comprehensive integrated-service delivery programs to ensure that all women have access to HIV counseling and voluntary testing and to services for related health needs (e.g., antiretroviral therapy, substance-abuse treatment, and social and support services).

The ZDV regimen recommended in the United States is not an affordable prevention strategy in many countries where HIV prevalence rates among women are highest. Worldwide, an estimated 8.8 million women and 800,000 children have HIV/AIDS; most of these persons reside in sub-Saharan Africa where resources for health services infrastructure are limited (World Health Organization, unpublished data, 1996). CDC and other organizations are collaborating with ministries of health in Africa and Asia to evaluate the effectiveness of shorter and simplified ZDV regimens, other antiretroviral medications, and other interventions for reducing perinatal HIV transmission. However, because ZDV treatment or other potential interventions are not universally effective in preventing perinatal transmission, primary prevention of HIV infection among children will continue to require preventing new HIV infections among women in the United States and other countries.

References

- 1. CDC. HIV/AIDS surveillance report. Atlanta: US Department of Health and Human Services, Public Health Service, 1996:3–4, 30–3. (Vol 8, no. 1).
- 2. Davis SF, Byers RH, Jr, Lindegren ML, Caldwell MB, Karon JM, Gwinn M. Prevalence and incidence of vertically acquired HIV infection in the United States. JAMA 1995;274:952–5.
- CDC. Recommendations of the U.S. Public Health Service Task Force on the use of zidovudine to reduce perinatal transmission of human immunodeficiency virus. MMWR 1994;43(no. RR-11).
- 4. CDC. U.S. Public Health Service recommendations for human immunodeficiency virus counseling and voluntary testing for pregnant women. MMWR 1995;44(no. RR-7).
- Davis SF, Steinberg S, Jean-Simon M, Rosen D, Gwinn M. HIV prevalence among U.S. childbearing women, 1989–1994 [Abstract]. Vancouver, British Columbia: XI International Conference on AIDS, 1996.
- 6. Lindsay MK, Peterson HB, Feng TI, Slade BA, Willis S, Klein L. Routine antepartum human immunodeficiency virus infection screening in an inner-city population. Obstet Gynecol 1989;74:289–94.
- 7. Thomas P, Singh T, Lindegren ML, Saletan S, Brooks A, Forlenza S. Patterns of zidovudine (ZDV) use in pregnant HIV-infected women in New York City (NYC) [Abstract]. Vancouver, British Columbia: XI International Conference on AIDS, 1996.
- 8. Simonds RJ, Nesheim S, Matheson P, et al. Declining mother-to-child HIV transmission following perinatal zidovudine recommendations, United States [Abstract]. Vancouver, British Columbia: XI International Conference on AIDS, 1996.
- Lindegren ML, Byers R, Fleming P, et al. A decline in the incidence of perinatally acquired (PA) AIDS in the United States [Abstract]. Vancouver, British Columbia: XI International Conference on AIDS, 1996.
- 10. CDC. 1995 Revised guidelines for prophylaxis against *Pneumocystis carinii* pneumonia for children infected with or perinatally exposed to human immunodeficiency virus. MMWR 1995;44(no. RR-4).

Serogroup Y Meningococcal Disease — Illinois, Connecticut, and Selected Areas, United States, 1989–1996

Neisseria meningitidis is a leading cause of bacterial meningitis and sepsis in the United States. N. meningitidis is classified into serogroups based on the antigenic characteristics of its capsular polysaccharide. During 1989–1991 in the United States, serogroups B and C accounted for most (91%) of invasive meningococcal disease while serogroup Y caused <5% (1); however, during 1992–1995, serogroup Y accounted for an increasing proportion of meningococcal disease. This report describes the epidemiology of serogroup Y meningococcal disease (SYMD) during 1991–1996 in Illinois and Connecticut, which conducted enhanced surveillance through active reviews of clinical records, and in areas participating in active laboratory-based surveillance during 1989–1995. The findings indicate a substantial increase in the proportion of meningococcal disease caused by N. meningitidis serogroup Y since 1989.

Illinois

In Illinois (1990 population: 11,430,602), 589 cases of invasive disease attributed to *N. meningitidis* were reported from January 1991 through March 1996, representing an annual incidence ranging from 0.9 to 1.0 per 100,000 population. Serogrouping was conducted for 371 (83%) of 447 culture-confirmed cases. The proportion of SYMD increased from 6% in 1991 to 29% in 1995; the proportion of disease attributed to serogroups B and C decreased from 85% to 59%.

Meningococcal Disease — Continued

From January 1991 through March 1996, the Chicago Department of Public Health received 145 reports of suspected meningococcal disease among persons residing in Chicago (1990 population: 2,783,726), and N. meningitidis was isolated from a normally sterile site in 133 (92%) case-patients. The overall annual incidence of cultureconfirmed disease ranged from 0.7 to 1.3 cases per 100,000 population. Of the 105 culture-confirmed isolates for which serogroup was known, 42 (40%) were serogroup Y; 29 (28%), serogroup B; 27 (26%), serogroup C; and two (2%), serogroup W-135. Among case-patients with known serogroups, the proportion of SYMD increased from 6% in 1991 to 71% in 1995. In comparison, the proportion of serogroups B and C decreased from 94% to 25%. Of 42 patients in Chicago with culture-confirmed SYMD, 22 (52%) were female; two (5%) died. The median age of patients with SYMD was 16 years compared with 2 years for patients with disease caused by non-serogroup Y meningococci. Although patients with SYMD were more likely to present with purulent sputum (six [14%] versus one [2%]) and chest pain (eight [19%] versus one [2%]), they were not more likely to have an infiltrate on chest radiograph (seven [23%] versus 10 [20%]). Case-fatality rates were similar among patients with SYMD (two [5%] of 42), compared with case-patients with disease caused by other known serogroups (six [10%] of 63).

Connecticut

From January 1991 through June 1996, a total of 190 culture-confirmed cases of invasive *N. meningitidis* infection among residents of Connecticut (1990 population: 3,287,116) were reported to the Connecticut Department of Public Health. The overall annual incidence of culture-confirmed cases ranged from 0.7 to 1.4 per 100,000 population. Of the 144 isolates for which serogroup was known, 69 (48%) were serogroup C; 38 (26%), serogroup Y; 35 (24%), serogroup B; and two (1%), serogroup W-135. The proportion of SYMD increased from 1991 (6%) to 1995 (35%). Of the 33 case-patients with SYMD identified since 1994, 18 (55%) were female; two (6%) died. The median age for patients with SYMD was 29 years, compared with 13 years for patients with disease caused by non-SYMD.

Active Laboratory-Based Surveillance

During 1989–1995, active laboratory-based surveillance was conducted in three counties in the San Francisco metropolitan area, eight counties in the Atlanta metropolitan area, and four counties in Tennessee, and during 1992–1995, in Maryland, representing an aggregate population of approximately 12 million. A case was defined as *N. meningitidis* isolated from a normally sterile site in a resident of a surveillance area. In the three active surveillance areas for which continuous data were available, the rate of SYMD per 100,000 persons increased from 1989 (0) to 1995 (0.4). Among the case-patients for whom serogroup was known, the proportion of SYMD increased from 1989 (0) to 1995 (32.5%). During the same period, the overall rate of meningococcal disease remained stable at 1.0–1.4.

During 1992–1995, in the four active surveillance areas, SYMD patients were older than patients with non-serogroup Y (median age: 21.8 years versus 14.2 years). Pneumonia was four times more likely to be diagnosed in persons with SYMD (12%) than in persons with other serogroups (3%), even after adjusting for age.

Meningococcal Disease — Continued

Laboratory Investigation

Multilocus enzyme electrophoresis (MEE) (2) using 24 enzymes was used to characterize genetic relatedness of serogroup Y isolates systematically sampled from 1995 Illinois surveillance (n=40), 1995 Connecticut surveillance (n=17), 1992–1995 U.S. active laboratory-based surveillance (n=40), 1972–1975 U.S. active laboratory-based surveillance (when SYMD accounted for 18% of the isolates submitted to CDC) (n=27) (3), and 1970–1974 surveillance of U.S. military personnel (n=12). Two major enzyme type complexes could be distinguished by a difference in peptidase mobility. One group of enzyme types accounted for 54% (33 of 97) of the isolates tested during 1992–1995, one of the 1972–1975 surveillance strains, and none of the strains from U.S. military personnel. The other group accounted for 34% (33 of 97) of 1992–1995 isolates and 62% (24 of 39) of the 1972–1975 isolates.

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The finding in this report that patients with SYMD in Chicago, Connecticut, and the active laboratory-based surveillance areas were older than patients with disease caused by non-serogroup Y meningococci is consistent with cases reported to CDC through NETSS. One possible explanation for this and the increase in SYMD is waning population immunity against SYMD. However, the increase in SYMD also may reflect, in part, the emergence of a distinct clone that differs in peptidase motility, as characterized by MEE. Although the association between epidemic meningococcal disease and clonality has been clearly established, the possible relation between shifts in endemic disease serogroup distribution and emergence of particular clones requires further assessment.

The clinical illness associated with SYMD differs from that of the other serogroups; in particular, findings from the active laboratory-based surveillance system indicated

Meningococcal Disease — Continued

that pneumonia was more common among patients with SYMD, consistent with studies in some military populations (5,6) in which serogroup Y was more likely than other serogroups to be associated with pneumonia and other forms of nonmeningitic disease. Meningococcal pneumonia may not be diagnosed because isolation of the organism from the sputum cannot distinguish persons who are meningococcal carriers from those with pneumonia caused by this organism, and because physicians may not consider *N. meningitidis* as a possible cause of pneumonia. As a result, infections that occur in the absence of meningitis or bacteremia may be underreported in current surveillance.

The current meningococcal vaccine (Connaught Laboratories, Swiftwater, Pennsylvania), which contains the purified polysaccharide capsules of serogroups A, C, W-135 and Y, has been effective in controlling serogroup C outbreaks and may be useful in controlling an SYMD outbreak. However, this vaccine has not been used to control endemic disease because its immunogenicity is low in young children and immunity is of limited duration. Conjugated vaccines for serogroup C, which are similar to those now available for preventing *Haemophilus influenzae* type b, are being evaluated in safety and immunogenicity trials (10). Because of the increased proportion of SYMD, manufacturers should consider developing a serogroup Y conjugate component for controlling endemic meningococcal disease.

References

- Jackson LA, Wenger JD. Laboratory-based surveillance for meningococcal disease in selected areas, United States, 1989–1991. In: CDC surveillance summaries (June). MMWR 1993;42(no. SS-2):21–30.
- Selander SK, Caugant DA, Ochman H, Musser JM, Gilmour MN, Whittam TS. Methods of multilocus enzyme electrophoresis for bacterial population genetics and systematics. Appl Environ Microbiol 1986:51:873–84.
- 3. Anonymous. Analysis of endemic meningococcal disease by serogroup and evaluation of chemoprophylaxis. J Infect Dis 1976;134:201–4.
- 4. Galaid El, Cherubin CE, Marr JS, Schaefler S, Barone J, Lee W. Meningococcal disease in New York City, 1973 to 1978: recognition of groups Y and W-135 as frequent pathogens. JAMA 1980;244:2167–71.
- 5. Koppes GM, Ellenbogen C, Gebhart RJ. Group Y meningococcal disease in United States Air Force recruits. Amer J Med 1977;62:661–6.
- 6. Smilack JD. Group Y meningococcal disease: twelve cases at an army training center. Ann Intern Med 1974;81:740–5.
- 7. Risko JA, Hodges GR. *Neisseria meningitidis* serogroup Y: incidence and description of clinical illness. Amer J Med Sci 1974;267:345–52.
- 8. Schlech WF III, Ward JI, Band JD, Hightower A, Fraser DW, Broome CV. Bacterial meningitis in the United States, 1978 through 1981: the National Bacterial Meningitis Surveillance Study. JAMA 1985;253:1749–54.
- 9. CDC. National Electronic Telecommunications System for Surveillance—United States, 1990–1991. MMWR 1991;40:502–3.
- Lieberman JM, Chiu SS, Wong VK, et al. Safety and immunogenicity of a serogroups A/C Neisseria meningitidis oligosaccharide-protein conjugate vaccine in young children: a randomized controlled trial. JAMA 1996;275:1499–503.

Update: Influenza Activity — United States, 1996–97 Season

In collaboration with the World Health Organization (WHO), its collaborating laboratories, and state and local health departments, CDC conducts surveillance to monitor influenza activity and to detect antigenic changes in the circulating strains of influenza viruses. This report summarizes influenza surveillance in the United States from September through early November 1996, which indicates that influenza activity is at typical levels for this time of year.

From September 4 through November 9, influenza A virus isolates were reported from 10 states (Alaska, California, Colorado, Iowa, Maryland, Montana, New York, North Carolina, Washington, and Wisconsin), and influenza B isolates were reported from seven states (Alaska, Illinois, Kentucky, Missouri, Ohio, Texas, and Wisconsin) (Figure 1). Most isolates were associated with sporadic cases. Of the five influenza type A isolates confirmed at CDC, all were identified as influenza type A(H3N2) and, when further characterized, were closely related to the influenza type A(H3N2) strain included in the 1996–97 influenza vaccine. Of the seven states reporting influenza B, Alaska and Illinois reported isolates obtained from patients who probably became infected while traveling outside the United States (Hong Kong and China, respectively).

For the week ending November 9, most state and territorial epidemiologists reported no influenza activity or sporadic* activity; Alaska and Montana reported regional activity.

Reported by: Participating state and territorial epidemiologists and state public health laboratory directors. World Health Organization collaborating laboratories. Epidemiology Div, Public Health Laboratory Svcs Communicable Diseases Surveillance Center, United Kingdom. Influenza Br and WHO Collaborating Center for Surveillance, Epidemiology, and Control of Influenza, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases, CDC.

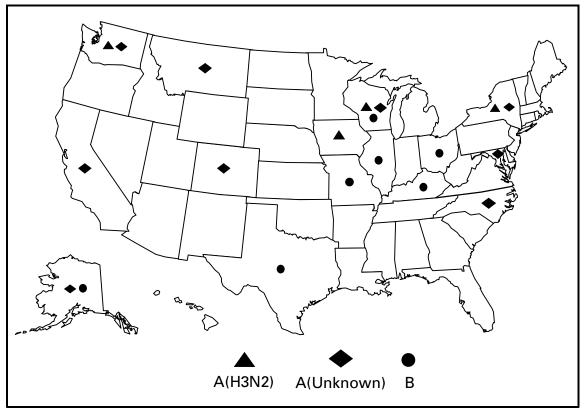
Editorial Note: Levels of activity described in this report are typical for September and October. Although the timing and intensity of influenza activity vary by season, sporadic influenza activity can begin in September, and isolated outbreaks can occur during October and November; widespread influenza activity usually does not begin before December. Although the optimal time for vaccination programs is October through mid-November, health-care providers should continue to offer vaccine to high-risk persons after mid-November and even after influenza activity has been documented in a community. Influenza vaccine contains influenza type A(H1N1), type A(H3N2), and type B strains representing the influenza virus strains that are expected to circulate during the 1996–97 influenza season. The 1996–97 vaccine contains A/Texas/36/91-like (H1N1), A/Wuhan/359/95-like (H3N2), and B/Beijing/184/93-like antigens. For both A/Wuhan/359/95-like and B/Beijing/184-like antigens, U.S. manufacturers used the antigenically equivalent strains A/Nanchang/933/95(H3N2) and B/Harbin/07/94 because of their growth properties.

When influenza vaccine is administered after local outbreaks of influenza type A have been reported, short-term prophylaxis with amantadine or rimantadine can be considered. These drugs can be used as treatment or prophylaxis for influenza type A

^{*}Levels of activity are 1) no activity; 2) sporadic—sporadically occurring influenza-like illness (ILI) or culture-confirmed influenza with no outbreaks detected; 3) regional—outbreaks of ILI or culture-confirmed influenza in counties with a combined population of <50% of the state's total population; and 4) widespread—outbreaks of ILI or culture-confirmed influenza in counties with a combined population of ≥50% of the state's total population.

Influenza Activity — Continued

FIGURE 1. Laboratory-confirmed influenza, by state and type of isolate — United States, September 4–November 9, 1996



infection, but they are not effective against influenza type B. Because early virologic surveillance has indicated circulation of influenza type A and type B viruses, use of viral culture and rapid antigen-detection testing throughout the season is particularly important (1).

Throughout the influenza season, surveillance data collected by CDC will be updated weekly and made available through the CDC voice information system (telephone ([404] 332-4551) and fax information system ([404] 332-4565 and requesting document number 361100). Information about local influenza activity is available from county and state health departments.

Reference

1. CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 1996;45(no. RR-5).

Salmonellosis Associated with a Thanksgiving Dinner — Nevada, 1995

On November 28, 1995, the county coroner's office notified the Clark County Health District in Las Vegas, Nevada, about a death suspected to have resulted from a food-borne disease. This report summarizes the investigation of the outbreak of gastroenteritis among persons who attended a Thanksgiving dinner. The investigation documented *Salmonella* serotype Enteritidis (SE) infection associated with eating improperly prepared turkey and stuffing containing eggs and emphasizes the need to use a meat thermometer to ensure complete cooking of turkey and stuffing.

During November 25–28, 1995, all six persons who attended a Thanksgiving dinner at a private home on November 23 and a seventh person who on November 25 ate food remaining from the dinner had onset of abdominal cramps, vomiting, and diarrhea. Two persons were hospitalized because of dehydration; a third person was found comatose at home and died from severe dehydration and sepsis. Stool cultures obtained from three persons, including the decedent, yielded SE phage type 13a. Turkey and stuffing were the only foods eaten by all seven ill persons. No leftover food was available for culture.

The Clark County Health District interviewed the ill persons (including the cook) to obtain details about the preparation and cooking of the turkey and stuffing. On November 22, a 13-pound frozen turkey was thawed for 6 hours in a sink filled with cold water. After thawing, the packet of giblets (heart, liver, and gizzard) was removed, and the turkey was stored in a refrigerator overnight. However, on November 23, parts of the turkey were noted to be frozen. The turkey was filled with a stuffing made from bread, the giblets, and three raw eggs, and then placed for 1 hour in an oven set at 350 F (177 C). The setting was lowered to 300 F (149 C) while the turkey cooked for an estimated additional 4 hours. The turkey was removed from the oven when the exterior had browned. A meat thermometer was not used. The stuffing was removed immediately and was served with the turkey. After the outbreak, health officials tested the oven set at 300 F (149 C) and found the temperature to be 350 F (176 C).

Reported by: O Ravenholt, MD, CA Schmutz, LC Empey, DJ Maxson, PL Klouse, AJ Bryant, Clark County Health District, Las Vegas; R Todd, DrPH, State Epidemiologist, Nevada State Health Div. Foodborne and Diarrheal Diseases Br, Div of Bacterial and Mycotic Diseases, National Center for Infectious Diseases, CDC.

Editorial Note: An estimated 2–4 million cases of salmonellosis occur each year in the United States, resulting in at least 500 deaths (1). Approximately 40,000 of these infections are culture-confirmed, serotyped, and reported to CDC through the National *Salmonella* Surveillance System. In 1995, SE was the most common serotype reported, accounting for 25% of the 40,720 serotyped culture-confirmed cases.

Salmonellosis is frequently associated with eating undercooked eggs and poultry. Undercooked eggs are a particularly common source of SE infections. During 1988–1992, among foodborne disease outbreaks of salmonellosis reported to CDC in which a single food item was implicated, consumption of turkey and eggs accounted for 4% and 14% of cases, respectively. In addition, eggs or foods containing eggs as a principal ingredient caused 64% of the SE outbreaks (2).

Factors probably associated with the outbreak described in this report included inadequate thawing, use of raw eggs in the stuffing, and undercooking; in addition, the browned color of the turkey may have caused the cook to believe that the turkey and

Salmonellosis — Continued

stuffing were thoroughly cooked. Although the original source of the *Salmonella* is unknown, the raw eggs used in the stuffing probably contained SE, and these eggs probably were incompletely cooked; undercooking may occur more commonly in turkeys that contain stuffing (J. Carpenter, Ph.D., University of Georgia, personal communication, 1996).

Each year, an estimated 45 million turkeys are eaten in the United States at Thanksgiving (J. DeYoung, National Turkey Federation, personnel communication, 1996). Salmonella infection may result from eating improperly cooked turkey and stuffing (3,4). This risk for infection can be reduced by cooking stuffing outside the turkey. Guidelines prepared by the U.S. Department of Agriculture (USDA) for persons who choose to cook stuffing inside the turkey recommend preparing the stuffing immediately before it is placed inside the turkey, stuffing the turkey loosely, inserting a meat thermometer into the center of the stuffing, and ensuring that the thermometer attains a temperature of at least 165 F (74 C). Additional recommendations for safely preparing and cooking a turkey include thawing the turkey completely before cooking, cooking in an oven set no lower than 325 F (163 C), and using a meat thermometer to ensure that the innermost part of the thigh attains a temperature of 180 F (82 C). Although the set temperature and cooking time can be used as guides to determine whether food is completely cooked, inaccuracies in the actual temperature and incomplete thawing before cooking can lead to undercooking. Use of a meat thermometer provides a more accurate determination of thorough cooking. Further advice on cooking turkeys and stuffing is available from USDA's Meat and Poultry Hotline, telephone (800) 535-4555.

References

- 1. Cohen ML, Tauxe RV. Drug-resistant *Salmonella* in the United States: an epidemiologic perspective. Science 1986;234:964–9.
- 2. Bean NH, Goulding JS, Loa C, Angulo FJ. Surveillance for foodborne-disease outbreaks—United States, 1988–1992. In: CDC surveillance summaries (October). MMWR 1996;45(no. SS-5).
- 3. CDC. Foodborne nosocomial outbreak of *Salmonella reading*—Connecticut. MMWR 1991;40: 804–6.
- 4. CDC. Restaurant outbreak of salmonellosis due to undercooked turkey—Washington. MMWR 1978;27:514,519.

Unintentional Administration of Varicella Virus Vaccine — United States, 1996

Since June 1995, seven separate cases of unintentional administration of varicella virus vaccine (Varivax®*) to pregnant women have been reported in the United States to the Varivax® Pregnancy Registry[†]. All seven women had household exposure to varicella, and varicella zoster immune globulin (VZIG) prophylaxis was indicated.

^{*}Use of trade names and commercial sources is for identification only and does not imply endorsement by the Public Health Service or the U.S. Department of Health and Human Services.

[†]This registry is maintained jointly by Merck and Company and by CDC; Merck and Company is responsible for daily management and operation of the registry. The registry was established to determine the risk for congenital varicella syndrome or other birth defects following vaccination with Varivax® 3 months before or at any time during pregnancy.

Varicella Vaccine — Continued

However, Varivax[®] was administered unintentionally instead of VZIG to these women. One of the women received five times the recommended dose of vaccine. All had negative histories for varicella, and the status of their immunity to varicella before receiving the vaccine was not reported to the registry. Gestational age at vaccination ranged from 6 to 31 weeks; four of the seven pregnancies were <20 weeks' gestation. Two of these women have since delivered healthy infants; pregnancy outcomes are pending for five women.

Reported by: JM Manson, PhD, RG Sharrar, MD, Merck Research Laboratories, Worldwide Product Safety and Epidemiology Div, West Point, Pennsylvania. Vaccine Safety and Development Activity, Child Vaccine Preventable Diseases Br, Epidemiology and Surveillance Div, National Immunization Program, CDC.

Editorial Note: The use of Varivax[®] is contraindicated during pregnancy (1) because its effects on the fetus are unknown and because infection with wild varicella zoster virus during the first half of pregnancy may result in congenital varicella syndrome (2). The Advisory Committee on Immunization Practices recommends that VZIG be used for postexposure prophylaxis in susceptible persons at high risk for varicella complications, including women exposed to varicella at any stage of pregnancy (1). The risk for congenital varicella syndrome after natural infection with wild varicella zoster virus is 1%–2%; because the virulence of the attenuated virus used in the vaccine is less than that of the wild-type virus, the risk to the fetus, if any, should be lower (1).

Two potential reasons these incidents occurred are 1) use of the wrong vial by mistake and 2) a lack of understanding of the appropriate indications for the use of these two products. These cases underscore the need for health-care providers and pharmacists to carefully check product labels before administering any drug and to read the package inserts for any drug if they are uncertain of the appropriate indications for its use. VZIG is shipped as a liquid in 2-mL or 10-mL vials and must be stored at 36 F–46 F (2 C–8 C). In contrast, Varivax[®] is shipped as a lyophilized powder for suspension in 0.7-mL vials, must be reconstituted with diluent before use, and must be stored at 5 F (–15 C).

Before a vaccine or any drug is administered to a woman of childbearing age, a health-care provider should be careful to obtain a history of pregnancy or intended pregnancy from the patient. Health-care providers are strongly encouraged to enroll any women who were unintentionally vaccinated with varicella virus vaccine 3 months before or at any time during pregnancy in the Varivax[®] Pregnancy Registry, telephone (800) 986-8999.

References

- 1. CDC. Prevention of varicella: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 1996;45(no. RR-11).
- 2. Enders G, Miller E, Cradock-Watson J, Bolley I, Ridehalgh M. Consequences of varicella and herpes zoster in pregnancy: prospective study of 1739 cases. Lancet 1994;343:1548–51.

FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending November 16, 1996, with historical data — United States

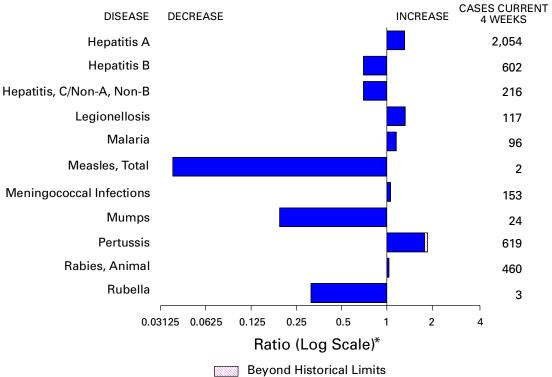


TABLE I. Summary — provisional cases of selected notifiable diseases, United States, cumulative, week ending November 16, 1996 (46th Week)

	Cum. 1996		Cum. 1996
Anthrax Brucellosis Cholera Congenital rubella syndrome Cryptosporidiosis* Diphtheria Encephalitis: California* eastern equine* St. Louis* western equine* Hansen Disease Hantavirus pulmonary syndrome*† HIV infection, pediatric*	82 3 1 2,034 1 105 2 - - 95 19 227	Plague Poliomyelitis, paralytic¶ Psittacosis Rabies, human Rocky Mountain spotted fever (RMSF) Streptococcal toxic-shock syndrome* Syphilis, congenital** Tetanus Toxic-shock syndrome Trichinosis Typhoid fever Yellow fever	5 37 1 657 13 225 25 112 17 318 1

^{*}Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

^{-:} no reported cases *Not notifiable in all states.

^{*}Not notifiable in all states.

† Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases (NCID).

§ Updated monthly to the Division of HIV/AIDS Prevention, National Center for HIV, STD, and TB Prevention (NCHSTP), last update October 29, 1996.

¶ Three suspected cases of polio with onset in 1996 has been reported to date.

**Updated quarterly from reports to the Division of STD Prevention, NCHSTP.

††This fatal case of yellow fever is the first occurrence of this disease reported in the United States since 1924. The infection is presumed to have been acquired in Brazil.

is presumed to have been acquired in Brazil.

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending November 16, 1996, and November 18, 1995 (46th Week)

	AIDS*		Chlamydia	Esche coli O NETSS [†]	richia 157:H7 PHLIS [§]	Gono	rrhea		atitis A,NB	Legionellosis		
Reporting Area	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1996	Cum. 1996	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	
UNITED STATES	56,760	60,827	335,242	2,480	1,366	265,688	345,981	2,928	3,564	882	1,022	
NEW ENGLAND	2,334	2,943	14,818	325	79	6,246	6,845	105	110	66	31	
Maine N.H.	39 72	82 77	844 397	22 39	38	53 80	83 99	8	12	2 5	6 2	
Vt.	18	28	U	34	31	42	58	36	13	4	-	
Mass. R.I.	1,134 159	1,336 205	6,210 1,652	146 15	10 -	1,950 441	2,410 475	55 6	78 7	28 27	19 4	
Conn.	912	1,215	5,715	69	-	3,680	3,720	-	-	N	N	
MID. ATLANTIC	15,871 2,180	16,428 1,973	40,726 N	209 141	43 16	32,138	38,010	276 218	434 222	205 69	180 50	
Upstate N.Y. N.Y. City	8,653	8,417	18,756	13	-	5,771 10,373	8,246 15,233	1	1	10	5	
N.J. Pa.	3,102 1,936	3,977 2,061	6,324 15,646	55 N	5 22	4,812 11,182	3,468 11,063	- 57	172 39	13 113	30 95	
E.N. CENTRAL	4,442	4,504	71,178	551	401	49,915	69,873	397	306	256	308	
Ohio	940	942	15,667	163	97	11,323	21,433	32	14	96	137	
Ind. III.	497 1,988	467 1,871	8,863 21,032	82 210	48 126	5,751 15,615	8,231 18,322	8 63	12 77	41 9	72 33	
Mich.	782	919	17,705	96	70	13,379	16,042	294	203	86	30	
Wis.	235	305	7,911	N	60	3,847	5,845	-	-	24	36	
W.N. CENTRAL Minn.	1,324 260	1,439 345	24,510 2,702	552 248	339 220	10,968 U	17,542 2,638	114 4	80 4	54 8	73 6	
lowa	76	94	3,801	121	88	1,004	1,431	48	13	10	20	
Mo. N. Dak.	673 11	642 5	10,635 2	64 16	- 15	7,190 -	9,921 29	36 -	19 5	17 -	16 3	
S. Dak.	11	17	1,315	24	-	166	199	-	1	2	3	
Nebr. Kans.	87 206	93 243	2,084 3,971	49 30	4 12	786 1,822	973 2,351	7 19	23 15	12 5	17 8	
S. ATLANTIC	14,203	15,365	49,123	131	64	85,788	96,375	232	221	139	157	
Del. Md.	248 2,008	277 2,287	1,148 6,016	1 N	2 8	1,287 12,681	2,024 12,134	1 3	- 7	11 27	2 25	
D.C.	1,120	896	0,010 N	-	-	3,865	4,165	- -	-	8	5	
Va. W. Va.	965 101	1,204 94	10,367 1	N N	32 3	8,192 473	9,479 598	16 9	18 44	23 1	21 4	
N.C.	744	898	-	43	12	16,722	21,174	45	55	12	31	
S.C. Ga.	717 2,058	815 1,999	- 11,051	12 30	7	10,161 16,391	11,079 17,698	28 U	19 15	6 3	30 14	
Fla.	6,242	6,895	20,540	33	-	16,016	18,024	130	63	48	25	
E.S. CENTRAL	1,931	1,919	27,783	69	59	30,875	35,958	504	890	44	52	
Ky. Tenn.	345 708	245 763	5,935 11,901	13 32	8 48	3,777 10,567	4,196 12,294	27 366	29 859	8 19	10 24	
Ala.	512	520	7,492	13	3	11,991	14,814	7	2	4	6	
Miss. W.S. CENTRAL	366 5,722	391 5,173	U 33,252	11 71	13	4,540 25.705	4,654 48,303	104 413	U 312	13 19	12 21	
Ark.	229	241	33,232	13	4	25,705	5,121	14	7	2	6	
La. Okla.	1,264 227	902 236	6,532 6,606	6 12	4 1	7,232 4,326	9,644 5,192	188 69	169 50	2 5	3 4	
Tex.	4,002	3,794	20,114	40	4	11,375	28,346	142	86	10	8	
MOUNTAIN	1,644	1,888	14,784	203	98	6,037	8,385	513	424	46	105	
Mont. Idaho	34 35	20 41	1,361	25 37	13	34 93	61 127	18 95	14 46	1 -	4 2	
Wyo.	5	17	507	11	9	33	48	168	178	7	12	
Colo. N. Mex.	437 139	572 148	U 3,521	73 11	40	1,077 840	2,513 942	56 66	61 44	8 2	38 4	
Ariz.	486	551	6,164	N	24	3,058	3,313	70	48	19	9	
Utah Nev.	161 347	113 426	1,398 1,833	31 15	12	260 642	249 1,132	22 18	11 22	3 6	16 20	
PACIFIC	9,288	11,168	59,068	369	270	18,016	24,690	374	787	53	95	
Wash. Oreg.	587 412	780 409	8,085 4,649	138 86	123 59	1,809 552	2,443 732	50 7	202 36	6 1	20	
Calif.	8,103	9,712	44,031	141	78	14,931	20,406	129	471	40	70	
Alaska Hawaii	28 158	62 205	1,093 1,210	4 N	2 8	388 336	601 508	3 185	2 76	1 5	- 5	
Guam	4	-	168	N	-	31	90	103	6	2	1	
P.R.	2,026	2,159	N	17	U	342	521	76	203	-	-	
V.I. Amer. Samoa	17 -	30	N -	N N	U U	-	32	-	-	-	-	
C.N.M.I.	1	-	N	N	U	11	51	-	5	-	-	

U: Unavailable

-: no reported cases

C.N.M.I.: Commonwealth of Northern Mariana Islands

^{*}Updated monthly to the Division of HIV/AIDS Prevention, National Center for HIV, STD, and TB Prevention, last update October 29, 1996.

†National Electronic Telecommunications System for Surveillance.

§Public Health Laboratory Information System.

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending November 16, 1996, and November 18, 1995 (46th Week)

	Lyı Dise		Mal	aria	Mening Dise			hilis Secondary)	Tubero	ulosis	Rabies	Animal
Reporting Area	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995
UNITED STATES	12,669	9,932	1,330	1,181	2,803	2,647	9,591	14,574	16,535		6,048	6,952
NEW ENGLAND	3,782	1,901	64	45	126	133	165	324	372	444	653	1,370
Maine N.H.	52 45	25 24	8	7 2	13 7	10 22	1	2 1	19 14	11 17	102 51	46 139
Vt. Mass.	15 316	9 135	7 21	1 15	4 54	11 42	- 70	- 61	- 185	4 246	127 97	166 391
R.I.	471	297	7	4	13	6	3	4	27	45	35	303
Conn. MID. ATLANTIC	2,883 7,692	1,411 6,503	18 359	16 333	35 256	42 319	91 425	256 733	126 2,841	121 3,826	241 1,314	325 1,785
Upstate N.Y.	3,948	3,294	75 194	61	80 33	90 48	68 120	76 337	381	479	972	1,067
N.Y. City N.J.	292 1,831	408 1,611	60	182 65	58	71	126	139	1,340 652	2,101 683	124	309
Pa.	1,621	1,190	30 115	25 149	85	110	111	181	468	563	218	409
E.N. CENTRAL Ohio	72 45	415 28	115 13	11	387 141	369 107	1,366 498	2,495 813	1,765 276	1,770 249	89 13	98 12
Ind. III.	24 3	18 18	13 35	17 73	54 108	52 95	174 372	305 933	155 910	161 918	8 23	14 15
Mich. Wis.	Ū	5 346	39 15	26 22	42 42	67 48	166 156	262 182	324 100	362 80	31 14	39 18
W.N. CENTRAL	184	204	47	25	220	167	321	670	419	511	472	343
Minn. Iowa	97 19	117 13	21 3	5 3	25 47	26 29	51 20	41 43	92 55	124 56	27 219	27 119
Mo. N. Dak.	27 1	46	10 1	8 1	91 4	64 1	207	548	176 6	201 4	18 65	30 27
S. Dak.	-	-	-	2	10	7			17	22	105	93
Nebr. Kans.	5 35	6 22	3 9	3 3	20 23	17 23	11 32	12 26	21 52	20 84	5 33	5 42
S. ATLANTIC	650	623	274	233	558	452	3,348	3,652	3,113	3,271	2,506	1,980
Del. Md.	105 377	49 395	4 75	1 62	2 65	6 36	35 569	15 443	30 262	53 350	68 559	83 393
D.C. Va.	3 47	3 52	7 49	16 53	10 55	8 60	121 352	97 532	121 282	91 255	10 549	11 406
W. Va. N.C.	11 63	22 67	5 27	4 15	14 68	8 72	3 972	10 1,014	50 447	64 393	94 635	109 428
S.C.	6	16	12	1	57	55	353	528	291	282	83	117
Ga. Fla.	1 37	14 5	27 68	36 45	127 160	99 108	607 336	679 334	562 1,068	617 1,166	272 236	257 176
E.S. CENTRAL	71	67	34	24	209	185	2,138	3,000	1,123	1,267	200	263
Ky. Tenn.	25 20	13 28	7 14	3 10	27 58	42 73	138 743	161 801	210 346	284 388	39 82	26 91
Ala. Miss.	7 19	9 17	6 7	8 3	76 48	38 32	485 772	588 1,450	370 197	353 242	76 3	137 9
W.S. CENTRAL	110	105	47	48	303	318	1,225	2,965	2,179	2,783	371	557
Ark. La.	24 6	8 8	6	2 5	33 55	31 50	131 454	457 924	170 175	217 307	28 15	46 42
Okla. Tex.	22 58	45 44	41	1 40	36 179	39 198	163 477	174 1,410	154 1,680	326 1,933	30 298	28 441
MOUNTAIN	7	12	54	57	157	186	124	187	547	610	137	171
Mont. Idaho	- 1	-	7	3 1	6 22	3 11	4	4	14 7	10 14	21	43 3
Wyo. Colo.	2	3	7 22	25	3 36	8 45	2 23	1 98	6 75	4 76	27 41	26 9
N. Mex.	1	1	2	6	25	33	1	6	73	70	6	6
Ariz. Utah	1	1 1	7 5	11 6	38 15	55 15	79 2	43 4	212 51	298 38	31 4	55 15
Nev.	2	6	4	5	12	16	13	31	109	100	7	14
PACIFIC Wash.	101 17	102 10	336 21	267 21	587 93	518 83	479 6	548 15	4,176 219	4,235 247	306 6	385 15
Oreg. Calif.	19 64	17 75	19 284	19 214	106 374	95 324	11 460	21 510	137 3,590	128 3,628	3 289	3 360
Alaska Hawaii	1	-	3	3 10	8	12	- 2	2	63 167	68 164	8	7
Guam	-	-	-	10	1	2	3	8	35	99	-	-
P.R. V.I.	-	-	-	1 2	4	23	124	261	63	162	40	37
Amer. Samoa	-	-	-	-	-	-	-	-	-	5	-	-
C.N.M.I.	-	-	-	1	-	-	1	9	-	36	-	-

U: Unavailable

-: no reported cases

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending November 16, 1996, and November 18, 1995 (46th Week)

	H. influ	-		Hepatitis (vi	al), by type			Measles	s (Rubeola)		
	inva Cum.	sive Cum.	Cum.	A Cum.	Cum.	Cum.	Ind	igenous Cum.	lm	ported [†] Cum.	
Reporting Area	1996*	1995	1996	1995	1996	1995	1996	1996	1996	1996	
UNITED STATES	868	998	25,289	26,728	8,695	8,792	-	415	-	49	
NEW ENGLAND Maine	28	38 3	365 21	283 28	178 2	203 12	-	11	-	4	
N.H.	9	10	22	12	20	20	-	-	-	-	
Vt. Mass.	1 16	2 12	10 176	5 123	11 60	5 80	-	1 9	-	1 3	
R.I. Conn.	2	5 6	20 116	32 83	9 76	8 78	-	- 1	-	-	
MID. ATLANTIC	128	151	1,672	1,706	1,293	1,262	-	23	-	5	
Upstate N.Y. N.Y. City	15 34	38 34	400 525	426 801	305 523	336 376	-	9	-	3	
N.J. Pa.	51 28	26 53	311 436	272 207	227 238	336 214	-	3 11	-	2	
E.N. CENTRAL	146	169	2,124	2,889	877	992	-	6	-	7	
Ohio Ind.	83 15	88 20	689 323	1,617 169	114 134	97 206	-	2	-	3	
III. Mich.	32 8	42 17	520 433	594 335	227 337	257 362	-	2	-	1 3	
Wis.	8	2	159	174	65	70	-	2	-	-	
W.N. CENTRAL Minn.	41 25	76 42	2,309 115	1,740 173	456 57	574 57	- U	20 16	Ū	2 2	
lowa	6	3	325	76	72	43	-	-	-	-	
Mo. N. Dak.	7 -	24 -	1,166 117	1,200 22	246 2	391 4	-	3 -	-	-	
S. Dak. Nebr.	1 1	1 3	42 194	72 49	5 44	2 31	-	-	-	-	
Kans.	1	3	350	148	30	46	-	1	-	-	
S. ATLANTIC Del.	171 2	195 -	1,280 18	1,031 9	1,324 7	1,158 8	-	5 1	-	9 -	
Md. D.C.	54 6	63	218 36	195 25	265 31	228 21	U -	- 1	U	2	
Va. W. Va.	9 10	28 8	165 14	190 24	128 30	99 50	-	-	-	3	
N.C.	24	28	157	96	280	273	-	3	-	1	
S.C. Ga.	5 39	2 60	49 150	42 53	88 32	49 62	-	-	-	2	
Fla. E.S. CENTRAL	22 26	6	473	397 1,896	463 751	368 748	-	2	-	1	
Ky.	4	11 5	1,135 41	41	55	61	-	-	-	-	
Tenn. Ala.	12 9	5	733 178	1,591 78	442 69	587 100	-	2	-	-	
Miss.	1	1	183	186	185	U	U	-	U	-	
W.S. CENTRAL Ark.	37 -	57 6	5,249 461	3,979 538	1,160 73	1,240 61	-	26 -	-	2	
La. Okla.	4 29	1 21	173 2,188	134 1,100	136 59	209 152	-	-	-	-	
Tex.	4	29	2,427	2,207	892	818	-	26	-	2	
MOUNTAIN Mont.	89 -	108	3,960 108	3,816 147	1,022 15	760 21	-	153 -	-	5 -	
ldaho Wyo.	1 35	4 8	224 33	291 101	85 44	91 26	-	1 1	-	-	
Colo. N. Mex.	14 10	16 14	413 328	468 738	120	119 277	-	4 17	-	3	
Ariz.	13	26	1,559	1,163	376 222	109	-	8	-	-	
Utah Nev.	8 8	11 29	939 356	636 272	85 75	62 55	-	117 5	-	2	
PACIFIC	202	193	7,195	9,388	1,634	1,855	-	169	-	15	
Wash. Oreg.	4 26	9 26	650 754	787 2,488	93 84	177 107	Ū	51 10	Ū	-	
Calif. Alaska	167 2	153 1	5,670 39	5,912 44	1,427 18	1,546 11	-	38 63	-	8 -	
Hawaii	3	4	82	157	12	14	-	7	-	7	
Guam P.R.	1	3	2 123	8 97	350	5 582	-	- 7	-	-	
V.I. Amer. Samoa	-	-	-	8 6	-	15 -	U	-	U U	-	
C.N.M.I.	10	11	1	24	5	22	ŭ	-	ŭ	-	

U: Unavailable

^{-:} no reported cases

 $^{^{*}\}text{Of 209}$ cases among children aged <5 years, serotype was reported for 48 and of those, 17 were type b.

[†]For imported measles, cases include only those resulting from importation from other countries.

TABLE III. (Cont'd.) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending November 16, 1996, and November 18, 1995 (46th Week)

	I Manadaa (Dada		IOVEI	indei id	o, 1999	(+011	I VVCCK		1		
	Measles (Rub			Mump	s		Pertussi	s		Rubella	a
Reporting Area	Cum. 1996	Cum. 1995	1996	Cum. 1996	Cum. 1995	1996	Cum. 1996	Cum. 1995	1996	Cum. 1996	Cum. 1995
UNITED STATES	464	292	7	571	765	208	5,026	3,995	-	201	116
NEW ENGLAND	15	10	-	2	11	51	1,088	579	-	27	48
Maine N.H.	-	-	-	-	4 1	10	20 127	42 45	-	-	1
Vt.	2	-	-	-	-	5	140	69	-	2	-
Mass. R.I.	12	3 5	-	2	2 1	36	742 30	391 4	-	21	8 -
Conn.	1	2	-	-	3	-	29	28	-	4	39
MID. ATLANTIC Upstate N.Y.	28	12 1	-	78 25	111 25	27 27	447 275	368 192	-	12 5	15 4
N.Y. City	12	5	-	17	16	-	38	49	-	4	8
N.J. Pa.	3 13	6	-	2 34	18 52	-	16 118	18 109	-	2 1	3
E.N. CENTRAL	13	15	_	93	155	16	549	500	_	3	4
Ohio	5	2	-	41	51	4	246	147	-	-	-
Ind. III.	3	2	-	9 20	9 45	-	102 149	55 106	-	1	-
Mich. Wis.	3 2	5 6	-	22 1	50	3	47 5	65 127	-	2	4
W.N. CENTRAL	22	2		18	43	2	362	251	_	_	1
Minn.	18	-	Ū	6	6	U	288	125	Ū	-	-
lowa Mo.	3	1	-	2 7	10 22	2	20 36	11 61	-	-	-
N. Dak.	-	-	-	2	1	-	1	8	-	-	-
S. Dak. Nebr.	-	-	-	-	4	-	4 9	12 12	-	-	-
Kans.	1	1	-	1	-	-	4	22	-	-	1
S. ATLANTIC Del.	14 1	19	2	96	116	18	565 15	327 10	-	93	10
Md.	2	1	U	26	34	U	200	43	U	-	1
D.C. Va.	1 3	-	-	1 15	25	- 15	4 95	6 25	-	2 2	-
W. Va.	-	-	-	-	-	-	2	-	-	-	-
N.C. S.C.	4	-	1	20 7	16 11	-	100 41	110 26	-	78 1	1 -
Ga. Fla.	2 1	4 14	- 1	3 24	8 22	3	17 91	24 83	-	10	- 8
E.S. CENTRAL	2	-	-	21	12	2	144	268	_	2	1
Ky.	-	-	-	-	-	-	90	25	-	-	-
Tenn. Ala.	2	-	-	3 3	4 4	1 1	21 24	206 35	-	2	1 -
Miss.	-	-	U	15	4	U	9	2	N	N	N
W.S. CENTRAL Ark.	28	34 2	1	33 2	51 7	-	115 12	287 38	-	3	7
La.	-	18	-	13	13	-	9	19	-	1	-
Okla. Tex.	28	- 14	1	1 17	31	-	17 77	31 199	-	2	- 7
MOUNTAIN	158	70	1	22	30	2	388	586	_	6	4
Mont.	-	-	-	-	1	1	34	4	-	-	-
Wyo.	1	2	1	1	3	1	101 7	103 1	-	2	-
Colo. N. Mex.	7 17	26 31	- N	3 N	2 N	-	98 61	104 131	-	2	-
Ariz.	8	10	-	1	2	-	27	153	-	1	3
Utah Nev.	119 5	1	-	2 15	11 11	-	22 38	27 63	-	1	1
PACIFIC	184	130	3	208	236	90	1,368	829	_	55	26
Wash.	51	19	Ū	19	13	90	642	303 55	Ū	2	1
Oreg. Calif.	10 46	1 108	3	157	201	U -	34 660	412	-	1 49	20
Alaska Hawaii	63 14	2	-	3 29	12 10	-	4 28	1 58	-	3	- 5
Guam	-	-	-	29 5	4	-	1	2	-	-	1
P.R.	7	3	-	1	2		1	2	-	-	-
V.I. Amer. Samoa	-	-	U U	-	3	U U	-	-	U U	-	-
C.N.M.I.	-	-	ŭ	-	1	ŭ	-	-	ŭ	-	

U: Unavailable

-: no reported cases

TABLE IV. Deaths in 121 U.S. cities,* week ending November 16, 1996 (46th Week)

	,	All Cau	ises, By	/ Age (Y	ears)		P&l [†]			All Cau	ises, By	Age (Y	ears)		P&I [†]
Reporting Area	All Ages	>65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	>65	45-64	25-44	1-24	<1	Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn. Cambridge, Mass. Fall River, Mass. Hartford, Conn. Lowell, Mass. Lynn, Mass. New Bedford, Mass. New Haven, Conn. Providence, R.I. Somerville, Mass. Springfield, Mass. Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J.		478 98 39 14 15 22 12 44 29 51 4 45 25 51 1,627 29 61 23	28 4 3 2 6 4 4 4 4 8 13 1 5 5 8 476 10 6 10 9	31 6 1 - 2 3 2 - 3 4 3 1 - 1 5 5 2 - 1 5 5 1 5 1 5 1 5 1 5 1 5 1 5 1 5 1 5	9 1 1 - 3 - - 2 1 - - 1 - - 49 3 1 1 1 1 3	13 4 - - 3 - - 5 - - 1 - 4 4 4	51 19 - - 2 4 1 1 10 1 9 133 7	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla. Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, Fla. Tampa, Fla. Washington, D.C. Wilmington, Del. E.S. CENTRAL Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala.	1,216 173 219 73 130 105 49 53 59 71 142 132 10 678 110 44 84 79 186 36 U	764 109 133 46 80 57 31 35 43 50 103 67 77 31 62 53 135 20	235 37 444 15 31 21 11 10 6 8 22 30 - 115 18 30 5 5 U	138 14 31 11 13 16 3 7 7 8 12 16 - 41 4 2 2 5 13 4 U	48 77 1 56 2 1 2 5 2 10 2 7 4 2 5 3 7 1 U	31 64 - 1 52 - 1 - 3 9 - 6 22 - 1	75 58 6 4 - 5 4 1 3 15 4 - 5 4 3 5 9 8 15 1 U
Erie, Pa.§ Jersey City, N.J. New York City, N.Y. Newark, N.J. Paterson, N.J. Philadelphia, Pa. Pittsburgh, Pa.§ Reading, Pa. Rochester, N.Y. Schenectady, N.Y. Scranton, Pa.§ Syracuse, N.Y. Trenton, N.J. Utica, N.Y. Yonkers, N.Y. E.N. CENTRAL Akron, Ohio Canton, Ohio	51 52 1,248 20 399 53 10 119 27 25 95 39 21 31 2,031 49	44 32 811 U 15 260 37 8 97 23 21 69 31 1,422 38 33	2 12 287 U 3 74 9 1 7 7 3 4 22 7 4 6 3 357 8	4 5 110 U 1 47 4 - 5 1 - 3 1 2 2 139 3	1 2 18 U 1 8 2 1 6 - - 1 42 1	1 21 U 9 1 - 4 - - - 70	3 2 45 U - 25 4 3 11 2 3 7 7 1 4 124	Montgomery, Ara. Nashville, Tenn. W.S. CENTRAL Austin, Tex. Baton Rouge, La. Corpus Christi, Tex. Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La. San Antonio, Tex. Shreveport, La. Tulsa, Okla. MOUNTAIN Albuquerque, N.M. Colo. Springs, Colo Denver, Colo.	139 1,454 56 45 47 210 72 123 356 73 111 181 67 113 878	912 40 26 37 118 45 79 223 52 60 118 38 76 594 67 27 61	305 9 12 6 49 14 27 75 8 22 44 19 20 154 24 7	11 144 4 3 1 27 7 9 39 5 19 13 2 15 85 10 2 14	5 48 2 1 6 5 3 1 1 4 9 2 5 - 2 7 - 1 5	1 45 1 3 3 10 1 5 8 4 1 4 3 2 17 3 2	13 66 3 1 1 2 3 2 30 5 - 13 3 3 3 67 2 11 12
Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Gary, Ind. Grand Rapids, Micl Indianapolis, Ind. Madison, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans. Kansas City, Kons. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	163 U 128 39 41 35 97 66 750 58 35 47 75 22	229 139 144 91 138 37 41 U 33 113 30 75 52 537 44 26 36 51 114 65 78 42 66	31 35 34 18 47 20 10 9 25 10 19 7 7 4 10 9 12 8 7 7 15 5 23 18 18 18 18 19 19 19 19 19 19 19 19 19 19 19 19 19	45 100 17 37 1 U 2 11 U 9 1 1 1 6 2 53 3 1 1 6 6 2 10 6 6 12 4 8	13 32 4 - 4 U2 6 U1 33 17 11 11 51 32 3	133 4 7 7 8 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	19 16 2 18 8 7 2 5 0 0 0 7 5 1 4 6 4 2 4 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, Utah Tucson, Ariz. PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawaii Long Beach, Calif. Los Angeles, Calif. Pasadena, Calif. Portland, Oreg. Sacramento, Calif. San Diego, Calif. San Diego, Calif. San Jose, Calif. San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash.	163 33 176 103 103 146 1,723 11 94 19 431 20 133 192 123	109 25 113 17 65 110 1,240 7 75 14 322 52 303 15 90 138 82 89 91 23 23 100 65 100 100 100 100 100 100 100 100 100 10	38 4 32 3 15 17 285 3 8 4 5 9 76 1 30 29 20 22 32 6 21 10 9	14 2 20 10 13 129 1 4 5 37 3 12 16 10 3 12 2 15 3 1	32 22 3 10 4 33 3 1 1 1 9 1 1 2 6 2 3 1 3 3 1 3 3 1 3 3 1 3 3 1 3 3 3 3 1 3	7 - 3 2 2 35 - 4 - 1 2 6 6 - 7 5 5 2 1 1 3 3 1 2 2 281	12 8 15 143 11 2 4 24 1 8 14 16 17 5 5 4 7 57

U: Unavailable -: no reported cases

*Mortality data in this table are voluntarily reported from 121 cities in the United States, most of which have populations of 100,000 or more. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included.

†Pneumonia and influenza.

Because of changes in reporting methods in these 3 Pennsylvania cities, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks.

Total includes unknown ages.

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