



## **Morbidity and Mortality Weekly Report**

Weekly

November 15, 2002 / Vol. 51 / No. 45

## HIV Testing Among Pregnant Women — United States and Canada, 1998–2001

Since 1994, the availability of increasingly effective antiretroviral drugs for both the prevention of perinatal human immunodeficiency virus (HIV) transmission and maternal treatment has resulted in a greater emphasis on prenatal HIV testing and substantial increases in prenatal testing rates. In 2000, preliminary data indicated that 766 (93%) of 824 HIV-infected women in 25 states knew their HIV status before delivery (CDC, unpublished data, 2002). However, an estimated 280-370 perinatal HIV transmissions continue to occur in the United States each year (1). The primary strategy to prevent perinatal HIV transmission is to maximize prenatal HIV testing of pregnant women. States and Canadian provinces have implemented three different prenatal HIV-testing approaches. To assess their effectiveness, CDC reviewed prenatal HIV-antibody testing rates associated with these approaches. Medical record data suggest that the "opt-in" voluntary testing approach is associated with lower testing rates than either the "opt-out" voluntary testing approach or the mandatory newborn HIV testing approach.

Under the opt-in approach, women typically are provided pre-HIV test counseling and must consent specifically to an HIV-antibody test. Under the opt-out approach, women are notified that an HIV test will be included in a standard battery of prenatal tests and procedures and that they may refuse testing (2). Under mandatory newborn HIV testing, newborns are tested for HIV, with or without the mother's consent, if the mother's HIV status is unknown at delivery.

Three methods were used to estimate prenatal testing rates among all women who delivered, regardless of whether they received prenatal care. First, eight U.S. areas that participated during 1998–1999 in CDC's Active Bacterial Core Surveillance/Emerging Infections Program (ABC) Network assessed HIV testing during prenatal care and ≤2 days before delivery by reviewing a stratified random sample of labor and delivery records and prenatal records forwarded to birthing hospitals

(3); in collaboration with CDC, network staff received a sample of records from all birthing hospitals in the surveillance areas and weighted testing rates to represent all liveborn infants in those areas. Second, public health investigators in each of the five Canadian provinces tallied the number of HIV tests among pregnant women that were submitted to provincial laboratories and divided the total by an estimate of all live and stillborn births in each province during the same year. Third, CDC analyzed weighted data collected in 1999 by interviewers in nine states for CDC's Pregnancy Risk Assessment Monitoring System (PRAMS) (an ongoing, population-based survey conducted in 32 states and New York City among women who have given birth during the preceding 2-6 months [4]), who had asked women if they had been tested for HIV during pregnancy. Data on state prenatal HIVtesting policies were obtained from the American College of Obstetricians and Gynecologists (5).

HIV-testing rates varied depending on which approach to testing was used. Rates for states using the opt-in approach to prenatal HIV testing included in the ABC Network ranged from 25% to 69% (Table 1), testing rates in Canada ranged from 54% to 83% (Table 2), and rates derived from PRAMS data ranged from 61% to 81% (Table 3). Two U.S. states (Arkansas and Tennessee) and two Canadian provinces (Alberta, and Newfoundland and Labrador) reported using

#### **INSIDE**

- 1016 Influenza Outbreak Madagascar, July–August 2002
- 1019 Influenza and Pneumococcal Vaccination Levels Among Persons Aged ≥65 Years — United States, 2001
- 1024 Use of Anthrax Vaccine in Response to Terrorism: Supplemental Recommendations of the Advisory Committee on Immunization Practices
- 1026 West Nile Virus Activity United States, November 7–13, 2002

The MMWR series of publications is published by the Epidemiology Program Office, Centers for Disease Control and Prevention (CDC), U.S. Department of Health and Human Services, Atlanta, GA 30333.

## SUGGESTED CITATION

Centers for Disease Control and Prevention. [Article Title]. MMWR 2002;51:[inclusive page numbers].

#### **Centers for Disease Control and Prevention**

Julie L. Gerberding, M.D., M.P.H. *Director* 

David W. Fleming, M.D.

Deputy Director for Science and Public Health

Dixie E. Snider, Jr., M.D., M.P.H.

Associate Director for Science

## **Epidemiology Program Office**

Stephen B. Thacker, M.D., M.Sc. *Director* 

## Office of Scientific and Health Communications

John W. Ward, M.D. Director Editor, MMWR Series

David C. Johnson

Acting Managing Editor, MMWR (Weekly)

Jude C. Rutledge Teresa F. Rutledge Jeffrey D. Sokolow, M.A. Writers/Editors, MMWR (Weekly)

Lynda G. Cupell Malbea A. Heilman Beverly J. Holland Visual Information Specialists

Quang M. Doan Erica R. Shaver Information Technology Specialists

## Division of Public Health Surveillance and Informatics

#### Notifiable Disease Morbidity and 122 Cities Mortality Data

Robert F. Fagan Deborah A. Adams Felicia J. Connor Lateka Dammond Patsy A. Hall Pearl C. Sharp an opt-out prenatal HIV-testing policy. ABC Network data indicated that Tennessee had a testing rate of 85% (Table 1). Canada's population-based data indicated a 98% testing rate in Alberta and a 94% testing rate in Newfoundland and Labrador (Table 2). PRAMS interview data indicated a 71% testing rate in Arkansas (Table 3), compared with a 57% testing rate early in 1997 before the law was implemented (Arkansas Department of Health, personal communication, 2002). Two states (New York and Connecticut) require HIV testing of newborns whose mothers were not tested during pregnancy. In New York, an ABC Network review of medical records in seven counties in the Rochester area indicated that the proportion of pregnant women who received a prenatal HIV test increased from 52% of 438 charts during January 1998-July 1999 to 83% of 112 charts during August-December 1999 after New York required that newborn HIV testing results be made available within 48 hours of specimen collection (Table 1). PRAMS data for 1999 indicated that the proportion of women statewide who reported having received an HIV test during pregnancy increased from 69% of 758 women during January-July to 93% of 502 during August–December (Table 3). In separate, statewide analyses of prenatal testing reported on newborn metabolic screening forms from all live-born infants, New York reported prenatal HIV-testing rates of 89% in 2000 and 93% in 2001 (New York State Department of Health, personal communication, 2002). In Connecticut, an ABC Network review of 668 charts indicated a testing rate of 31% during January 1998-September 1999, compared with 81% of 93 charts reviewed during October-December 1999 after enactment of the mandatory newborn testing law (Table 1).

Reported by: A Roome, PhD, J Hadler MD, Connecticut Dept of Public Health. G Birkhead, MD, AIDS Institute, New York State Dept of Health. S King, MD, The Hospital for Sick Children, Toronto; C Archibald, MD, Health Canada. S Schrag, DPhil, Active Bacterial Core Surveillance/Emerging Infections Program Network, Div of Bacterial and Mycotic Diseases, National Center for Infectious Diseases; A Lansky, PhD, Pregnancy Risk Assessment Monitoring System, Div of Reproductive Health, National Center for Chronic Disease Prevention and Health Promotion; S Sansom, PhD, M Fowler, MD, I Onorato, MD, J Anderson, PhD, Div of HIV/AIDS Prevention, National Center for HIV, STD, and TB Prevention, CDC.

**Editorial Note:** Prenatal HIV testing affords the best opportunity for the prevention of perinatal HIV transmission. On the basis of clinical trial data, perinatal HIV-transmission rates among HIV-infected women who begin antiretroviral treatment during pregnancy are as low as  $\leq 2\%$  (6), compared with 12%–13% early transmission rates among women who do not begin preventive treatment until labor and delivery or after birth (7) and 25% among women who receive no preventive treatment (8).

TABLE 1. Number of medical charts reviewed and percentage of charts with a documented prenatal HIV test for pregnant women, by testing approach and area — Active Bacterial Core Surveillance/Emerging Infections Program Network, eight states, 1998–1999

State	Testing approach	No. charts reviewed	% with HIV test*	(95% CI <sup>†</sup> )
Tennessee (five counties)	Opt-out <sup>§</sup>	623	85%	(82.1%-88.5%)
New York (seven counties in the Rochester area)	Mandatory newborn testing without expedited testing requirement Mandatory newborn testing; results returned within 48 hours th	* 438 112	52% 83%	(47.3%–57.1%) (75.0%–91.5%)
Connecticut	Opt-in <sup>§§</sup> Mandatory newborn testing; results within 48 hours <sup>¶¶</sup>	668 93	31% 81%	(27.0%–34.3%) (72.3%–88.7%)
Maryland	Opt-in	665	69%	(65.4%-72.8%)
Georgia (20 counties in the Atlanta area)	Opt-in	866	66%	(61.8%–69.6%)
Minnesota (seven counties in the Minneapolis/St. Paul area)	Opt-in	605	62%	(57.5%–65.8%)
California (three counties in the San Francisco area)	Opt-in	575	39%	(34.5%–42.4%)
Oregon (three counties in the Portland area)	Opt-in	498	25%	(21.5%–29.1%)

\* Percentages are weighted to reflect all live-born infants and account for sample weights and design.

Confidence interval.

\*\* Policy in effect until August 1999.

<sup>††</sup> Policy in effect beginning August 1999.

<sup>11</sup> Policy in effect beginning October 1999.

TABLE 2. Number of women delivering and percentage receiving prenatal HIV testing, by testing approach, year, and province — Canada, 1999–2001

		Testing		
Province	Year	approach	No.	(%)*
Alberta	2000	Opt-out <sup>†</sup>	37,963	(98)
Newfoundland and Labrador	2001	Opt-out	4,770	(94)
Quebec	1999	Opt-in <sup>§</sup>	73,781	(83)
British Columbia	1999	Opt-in	41,739	(80)
Ontario	2001	Opt-in	129,758	(54)

<sup>\*</sup> Canadian prenatal human immunodeficiency virus (HIV) testing rates are \_based on all live-born infants in each province for the year.

Pregnant women are required to consent specifically to an HIV test.

Among the three prenatal HIV testing approaches assessed in this report, opt-out voluntary testing and the mandatory testing of newborns appear to be associated with the highest testing rates. On the basis of the chart-review methodology, prenatal testing rates were higher in Tennessee, which uses the opt-out approach, than rates in states using the opt-in approach and similar to rates achieved with mandatory newborn testing in New York during the same time period. A similar trend was observed among Canadian provinces. In New York and Connecticut, mandatory HIV testing of newborns was associated with increases in prenatal testing rates. On the

basis of PRAMS data, three of seven states using the opt-in approach achieved lower prenatal HIV-testing rates than states using the opt-out or mandatory newborn testing approaches.

Increases in prenatal HIV-testing rates were noted in states that shifted from an opt-in approach to either an opt-out or mandatory newborn testing approach and were probably associated with a greater likelihood that woman were offered HIV testing during prenatal care. Data from the Perinatal Guidelines Project indicated that the majority of women will accept HIV testing if it is recommended by their health-care provider (9). Perinatal HIV experts and professional organizations have advocated streamlining prenatal HIV pre-test counseling and consent procedures to reduce barriers to the offer of testing by health-care providers (1,2,10).

The findings in this report are subject to at least seven limitations. First, testing results for each strategy are for all women, and the proportion of HIV-positive women who accepted testing under each strategy is not known. Second, among women who did not receive prenatal testing, the proportion of women who were not tested because they did not seek prenatal care is unknown. Third, among women who did not receive prenatal testing, the proportion of women who were tested at labor and delivery or whose infants were tested at birth is not known. Fourth, maternal self-reported data from

<sup>§</sup> Pregnant women are informed that a human immunodeficiency virus (HIV) test is being conducted as a standard part of prenatal care and that they may refuse it.

Infants are tested for HIV antibodies if the mother was not tested during prenatal care or at delivery. Mother's consent is not required. Neither Connecticut nor New York have data on numbers of newborn infants tested under these laws.

<sup>§§</sup> Pregnant women are required to consent specifically to an HIV test.

<sup>†</sup>Pregnant women are informed that an HIV test is being conducted as a standard part of prenatal care and that they may refuse it.

TABLE 3. Percentage of women who responded that they had, had not, or did not know if they had received an HIV test during their most recent pregnancy, by testing approach and state — Pregnancy Risk Assessment Monitoring Survey, United States, 1999

			F	Percentage	е
State	Testing approach	No.	Yes	No	Don't know
Florida	Opt-in*	1,990	81%	13%	6%
New York <sup>†</sup>	Mandatory newborn testing (1/99–7/99)	758	69%	28%	3%
	Mandatory newborn testing; results within 48 hours of delivery (8/99–12/99)	502	93%	6%	1%
North Carolina	Opt-in	1,770	75%	20%	5%
Illinois	Opt-in	1,994	72%	17%	10%
Colorado	Opt-in	2,039	72%	21%	8%
Arkansas	Opt-out <sup>§</sup>	1,892	71%	13%	16%
West Virginia	Opt-in	1,327	67%	22%	11%
Oklahoma	Opt-in	1,980	62%	25%	13%
Ohio	Opt-in	1,589	61%	25%	4%

<sup>\*</sup> Pregnant women are required to consent specifically to a human immunodeficiency virus (HIV) test.

PRAMS collected 2–6 months after delivery might be subject to recall bias. Fifth, PRAMS data do not indicate whether a prenatal-care provider was aware of the woman's HIV status. Sixth, among the women interviewed in PRAMS, up to 16% (in Arkansas) indicated they did not know if they had been tested. Finally, chart abstraction can document only prenatal HIV testing recorded in maternal medical records; without such documentation, clinicians might not be aware of the need to offer effective perinatal interventions to infected women and their HIV-exposed infants.

This report emphasizes the need for better data to assess perinatal HIV testing rates in the United States. Ongoing, randomized reviews of prenatal, labor/delivery, and pediatric charts, with a sampling framework ensuring that the sample is representative of the population of women delivering, might provide the most valid approach to assessing a state's progress on perinatal HIV testing and prevention. CDC is working with states with high HIV prevalence rates among women of childbearing age and high numbers of pediatric AIDS cases to ensure standardized monitoring of prenatal testing rates. The data suggest that jurisdictions that use an opt-in approach and that have low prenatal HIV-testing rates should reevaluate their approach.

## References

- 1. CDC. Revised recommendations for HIV screening of pregnant women. MMWR 2001;50(No. RR-19).
- Institute of Medicine. Reducing the Odds: Preventing Perinatal Transmission of HIV in the United States. Washington, DC: National Academy Press, 1998.
- Schrag SJ, Zell ER, Lynfield R, et al. A population-based comparison of strategies to prevent early-onset group B streptococcal disease in neonates. N Engl J Med 2002;347:233–9.
- CDC. Prevalence of selected maternal behaviors and experiences, Pregnancy Risk Assessment Monitoring System (PRAMS). In: CDC surveillance summaries (April 26). MMWR 2002;51(No. SS-2).

- American College of Obstetricians and Gynecologists. Survey of state laws on HIV and pregnant women, 1999–2000. Moore KG, ed. Washington, DC: American College of Obstetricians and Gynecologists, 2000.
- Dorenbaum A, Cunningham CK, Gelber RD, et al. Two-dose intrapartum/newborn nevirapine and standard antiretroviral therapy to reduce perinatal HIV transmission: a randomized trial. JAMA 2002;288:189–98
- 7. Guay LA, Musoke P, Fleming T, et al. Intrapartum and neonatal single-dose nevirapine compared with zidovudine for prevention of mother-to-child transmission of HIV-1 in Kampala, Uganda: HIVNET 012 randomised trial. Lancet 1999;354:795–802.
- 8. Connor EM, Sperling RS, Gelber R, et al. Reduction of maternal-infant transmission of human immunodeficiency virus type 1 with zidovudine treatment. N Engl J Med 1994; 331:1173–80.
- 9. Fernandez MI, Wilson TE, Ethier KA, et al. Acceptance of HIV testing during prenatal care. Public Health Rep 2000;115:460–8.
- American College of Obstetricians and Gynecologists. Joint statement of ACOG/AAP on human immunodeficiency virus screening. ACOG statement of policy. Washington, DC: American College of Obstetricians and Gynecologists, 1999.

## Influenza Outbreak — Madagascar, July-August 2002

In mid-July 2002, Madagascar health authorities were notified of a substantial number of deaths attributed to acute respiratory illness (ARI) in the village of Sahafata (population: 2,160), located in the rural highlands of Fianarantsoa Province, southeastern Madagascar (Figure 1). This region is approximately 450 km (280 miles) south of the capital Antananarivo. The Madagascar Ministry of Health (MOH) and the Institut Pasteur, Madagascar (IPM) initiated an investigation, which found an attack rate of 70% for ARI, with 27 deaths in Sahafata. Pharyngeal swab specimens were collected from ill persons for viral culture. Of the four influenza A viruses that were isolated at IPM, two were identified

<sup>†</sup> Excludes New York City.

<sup>§</sup>Pregnant women are informed that an HIV test is being conducted as a standard part of prenatal care and that they may refuse it.

FIGURE 1. A remote village in Madagascar's Fianarantsoa Province, one of many areas reporting an outbreak of influenza-like illness during July-August 2002



Photo/Tim Uyeki

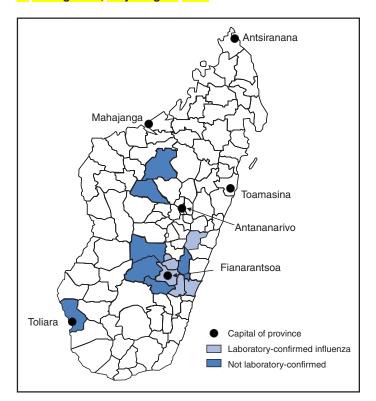
as type A (H3N2) viruses. In late July, health authorities investigated a similar outbreak in Ikongo District, Fianarantsoa Province. In August, MOH requested assistance from the World Health Organization (WHO) and CDC in investigating the outbreak. In response, an international team of experts from CDC; Institut de Veille Sanitaire, France; Institut Pasteur, France; and WHO was mobilized from the Global Outbreak Alert and Response Network; the team arrived in Madagascar on August 14. This report summarizes the preliminary epidemiologic and virologic findings, which suggest that the outbreak was attributable to influenza A (H3N2) viruses. Further surveillance and research about the epidemiology of influenza in Madagascar is planned.

Nationwide surveillance for influenza-like illness (ILI) cases implemented by MOH suggested that the outbreak peaked during the week of August 22. As of September 19, the outbreak appeared to be over, with 30,304 cumulative cases and 754 deaths reported from 13 of 111 health districts and four of six provinces (Figure 2); approximately 85% of cases were reported from Fianarantsoa Province. The majority of illnesses occurred in rural areas, and 95% of deaths occurred away from health facilities and could not be investigated. No standardized case definition was used, and the degree of overreporting or underreporting of ILI cases is uncertain.

Field investigations were conducted in three highland districts of Fianarantsoa Province in which high numbers of cases and deaths had been reported. The investigations' objectives were to confirm the etiology of the outbreak and to make recommendations based on the epidemiologic findings. An analysis of ARI data from 1999–2002 collected at health centers indicated that ARI cases in highland districts peaked each

FIGURE 2. Districts reporting cases of influenza-like illness

Madagascar, July-August 2002



year during the winter months of May-September. The peaks in ARI cases coincided with peaks of mortality from all causes and from respiratory conditions such as pneumonia during 1999–2002. In Ikongo District (estimated 2002 population: 161,494) of Fianarantsoa Province, the numbers of ARI cases evaluated at health centers and deaths from all causes that occurred during July-August were substantially higher than those that occurred during identical periods in previous years. However, the ratio of deaths to ARI cases appeared to be similar to proportions recorded during previous years. In three communes of Ikongo District (estimated 2002 population: 58,037), 54% of the reported deaths attributed to ARI that occurred during July-August were among children aged <5 years, but the highest mortality rate was among persons aged ≥60 years. A survey of a remote village (population: 750) in Ikongo District indicated an ARI attack rate of 67% and an estimated case-fatality ratio of 2%. In contrast, no unusually high morbidity or mortality was reported among the population of Fianarantsoa Province's capital city or in Antananarivo (estimated 2002 population: 1.25 million), where morbidity and virologic surveillance for influenza is conducted all year by IPM.

During July 19-August 22, a total of 152 respiratory specimens were collected for viral isolation from ill persons in three areas of Fianarantsoa Province (Sahafata, Ikongo, and

Manandriana) where outbreaks occurred. The international team also used rapid influenza-antigen tests to test specimens in the field. Influenza A viruses were isolated from specimens collected from ill persons in each area that was investigated; 27 influenza isolates were characterized antigenically at IPM and confirmed by the WHO Collaborating Centre for Reference and Research on Influenza, London, United Kingdom; all isolates were A/Panama/2007/99-like (H3N2) viruses. The A (H3N2) component of both the 2002 Southern Hemisphere and 2002–03 Northern Hemisphere influenza vaccines are well matched to the outbreak strain.

Reported by: L Rasoazanamiarina, MD, A Lamina, MD, Ministry of Health; M Andrianarivelo, MD, G Razafitrimo, Institut Pasteur; A Ndikuyeze, MD, B Andriamahefazafy, MD, World Health Organization, Antananarivo, Madagascar. C Paquet, MD, International Health Dept, I Bonmarin, MD, Infectious Diseases Dept, Institut de Veille Sanitaire, Saint-Maurice, France. J Manuguerra, PhD, Molecular Genetics Unit for Respiratory Viruses, National Reference Center for Influenza, Institut Pasteur, Paris, France. B Koumare, PhD, World Health Organization Regional Office for Africa, Brazzaville, Congo. N Shindo, MD, K Stohr, PhD, K Ait-Ikhlef, Dept of Communicable Disease Surveillance and Response, World Health Organization, Geneva, Switzerland. T Uyeki, MD, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases, CDC.

Editorial Note: The epidemiologic and virologic data suggest that the large outbreak described in this report was attributable to influenza A/Panama/2007/99-like (H3N2) viruses, which have been in circulation worldwide for several years. Influenza outbreaks in remote regions have been reported rarely (1-4). Several factors might have contributed to the widespread ARI morbidity and unusually high mortality reported from rural highland regions during this outbreak. In remote villages, crowded living conditions during an unusually cold and wet winter might have facilitated personto-person transmission of influenza among highly susceptible populations. Fianarantsoa Province is one of the poorest regions of Madagascar; malnutrition is prevalent, and access to health care is poor. These factors might have been exacerbated further by civil unrest during December 2001-June 2002.

This outbreak illustrates several important lessons for controlling influenza outbreaks in developing countries and for global pandemic influenza planning. Because the outbreak occurred primarily in remote areas, awareness of the outbreak and response by health authorities were delayed. Although influenza surveillance is conducted in Antananarivo by IPM's WHO-recognized National Influenza Center, no data were available for the most affected areas. In Madagascar, as in many developing countries, efforts to assess and control the outbreak were complicated by at least seven factors: 1) malnutrition,

2) poor access to health care in remote areas, 3) difficulties in reaching rural populations, 4) limited communicable disease surveillance, 5) shortages of antibiotics to treat secondary bacterial complications, 6) the unavailability of influenza vaccine, and 7) lack of awareness about influenza. In addition, limited influenza surveillance has prevented an understanding of the epidemiology and impact of influenza in many developing countries, especially in Africa (5). In response to this outbreak, the team recommended expanding influenza surveillance, educating the public and health-care providers about influenza, improving access to health care in rural areas, and ensuring that adequate supplies of antibiotics are available at health-care centers to treat bacterial complications of influenza. Influenza vaccination was not recommended because the outbreak was already widespread in August, and the ability to distribute vaccine in remote areas was extremely limited. Members of the international team plan to return to Madagascar to assist MOH to better characterize the outbreak.

## **Acknowledgments**

This report is based on contributions by R Migliani, MD, M Ratsitorahina, PhD, P Grosjean, N Rasolofonirina, MD, L Rabarijaona, MD, Institut Pasteur, Antananarivo; D Rabdrianasolo, MD, C Ravaonjanahary, PhD, M Ratolojanahary, MD, Ministry of Health; J Rasamizanaaka, MD, H Ravokatsoa, MD, L Razafilahy, Dept of Public Health, Fianarantsoa Province; B Tanjaka, MD, Ikongo District; P Rakotoarisoa, MD, Manadriana District; E Raharilalao, MD, Anjoma, Nadihizana District, Madagascar. C Bouchier, PhD, V Lorin, Institut Pasteur, Paris, France. A Hay, PhD, World Health Organization Collaborating Center for Reference and Research on Influenza, London, United Kingdom. R Arthur, PhD, M Ryan, MD, World Health Organization Global Alert and Response Team, Communicable Disease Surveillance and Response Dept, Geneva, Switzerland. S Harper, MD, K Fukuda, MD, J LeDuc, PhD, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases, CDC.

## References

- 1. World Health Organization. Acute respiratory infection, Afghanistan. Wkly Epidemiol Rec 1999;74:65.
- Corwin AL, Simanjuntak CH, Ingkokkusumo G, et al. Impact of epidemic influenza A-like acute respiratory illness in a remote jungle highland population in Irian Jaya, Indonesia. Clin Infect Dis 1998;26:880–8.
- 3. Canil KA, Pratt RD, Sungu MS, et al. An outbreak of influenza A in the highlands of Papua New Guinea. Southeast Asian J Trop Med Public Health 1984;15:265–9.
- Tangkanakul W, Tharmaphornpilas P, Thawatsupha P, et al. An outbreak of influenza A virus in a hilltribe village of Mae Hong Song Province, Thailand, 1997. J Med Assoc Thai 2000;83:1005–10.
- 5. Schoub BD, McAnerney JM, Besselaar TG. Regional perspectives on influenza surveillance in Africa. Vaccine 2002;20:S45–S46.

## Influenza and Pneumococcal Vaccination Levels Among Persons Aged ≥65 Years — United States, 2001

Two vaccine-preventable diseases, influenza and pneumococcal disease, contribute to the mortality of older persons in the United States. Influenza caused an average of 20,000 deaths per year during influenza epidemics in the United States from 1969 to 1996; persons aged ≥65 years accounted for approximately 90% of these deaths (1). Pneumococcal disease caused approximately 3,400 deaths among persons aged ≥65 years in the United States in 1998 (2). National health objectives for 2010 include increasing influenza and pneumococcal vaccination levels to ≥90% among persons aged ≥65 years (objective nos. 14.29a and 14.29b, respectively) (3). To assess progress toward achieving these objectives, CDC analyzed data from the 2001 Behavioral Risk Factor Surveillance System (BRFSS). This report summarizes the results, which indicate that the estimated point prevalences of influenza and pneumococcal vaccination were <80% among persons aged ≥65 years in all reporting areas. Influenza vaccination levels during 2000-2001 decreased from 1998-1999 levels in 27 of 52 reporting areas; pneumococcal vaccination prevalence increased a median of 7 percentage points from 1999 to 2001. Continued efforts are needed to increase the proportion of older adults who receive influenza and pneumococcal vaccines; health-care providers should offer pneumococcal vaccine all year and should continue to offer influenza vaccine during December and throughout the influenza season, even after influenza activity has been documented in the community.

BRFSS is a state-based, random-digit-dialed telephone survey of the noninstitutionalized civilian U.S. population aged >18 years. The survey is conducted in all 50 states, the District of Columbia, and three U.S. territories. Questions about influenza vaccination ("During the past 12 months, have you had a flu shot?") and pneumococcal vaccination ("Have you ever had a pneumonia vaccination?") were asked in all reporting areas in odd-numbered years starting in 1993. The response rate (CASRO method) was >60% in 10 of the 54 reporting areas (median: 51.1%; range: 33.3%-81.5%). Response rates for persons aged ≥65 years were not available. In 2001, the sample included 39,910 respondents aged ≥65 years. Respondents who reported an unknown influenza (0.3%) or pneumococcal (2.6%) vaccination status were excluded from the analysis. Overall vaccination levels were estimated for the 50 states and the District of Columbia; data for Guam, Puerto Rico, and the Virgin Islands were reported in area-specific results only. Data were weighted by age, sex, and, in some areas, by race/ethnicity to reflect each area's estimated adult population. SUDAAN was used to calculate point estimates and 95% confidence intervals (CIs) and to conduct multivariable logistic regression to calculate odds ratios and test associations of vaccination status with age, race/ethnicity, sex, education level, geographic region, self-reported health, diabetes status, smoking status, and asthma history.

During 2001, a total of 64.9% (95% CI=64.0%–65.8%) of respondents aged ≥65 years reported having received an influenza vaccination during the preceding year (Table 1), compared with 66.9% (95% CI=66.0%–67.8%) in 1999 (4). Previous analyses have indicated percentage point increases of 7.7%, 7.4%, and 1.5% from 1993 to 1995, 1995 to 1997, and 1997 to 1999, respectively (4). Estimated influenza vaccination levels exceeded 60% in 48 of 54 reporting areas; in 34 of these areas, 95% CIs exceeded 60% (Table 2). Vaccination prevalence ranged from 36.8% (Puerto Rico) to 79.0% (Hawaii). Of the 52 areas for which data were available for both 1999 and 2001, the median percentage point difference from 1999 to 2001 was −0.9 (range: −9.6–6.5).

The proportion of respondents reporting having ever received pneumococcal vaccination increased 5.9 percentage points, from 54.1% (95% CI=53.2%–55.1%) in 1999 to 60.0% (95% CI=59.2%–60.8%) in 2001 (Table 1). Previous analyses indicated percentage point increases of 6.9%, 9.8%, and 8.7% from 1993 to 1995, 1995 to 1997, and 1997 to 1999, respectively (4). Of the 52 reporting areas for which data were available for both 1999 and 2001, the proportion of respondents reporting having ever received pneumococcal vaccination increased in 51 areas (Table 2). Estimated pneumococcal vaccination levels exceeded 60% in 32 reporting areas, and 95% CIs exceeded 60% in 18 of these areas. Vaccination prevalence ranged from 24.1% (Puerto Rico) to 70.9% (Oregon).

Receipt of one vaccine was associated with receipt of the other vaccine. A total of 10.5% of respondents reported pneumococcal vaccination only, and 15.4% reported recent influenza vaccination only; 49.3% reported both, and 24.7% reported having received neither.

The estimated percentages of non-Hispanic blacks and Hispanics having received influenza (non-Hispanic black=48.1% and Hispanic=55.2%) and pneumococcal vaccination (non-Hispanic black=39.4% and Hispanic=41.6%) were less than those for non-Hispanic whites having received influenza (67.1%) and pneumococcal vaccination (63.5%) (Table 1). After accounting for variations in age, sex, education level, self-reported health, diabetes status, geographic region, smoking status, and asthma history by logistic regression, the disparity in vaccination coverage between non-Hispanic whites and non-Hispanic blacks and Hispanics remained statistically significant (Table 3).

TABLE 1. Percentage of persons aged ≥65 years who reported receiving influenza vaccine during the preceding year or pneumococcal vaccine ever, by selected characteristics — Behavioral Risk Factor Surveillance System (BRFSS), United States, 2001

Characteristic  Age group (yrs) 65-74 ≥75  Race/Ethnicity White, non-Hispanic Black, non-Hispanic Hispanic Other <sup>†</sup> Sex Men Women  Region <sup>§</sup> New England Mid Atlantic Northeast Central	% 62.1 69.1 67.1 48.1 55.2 65.7	(95% CI*) (61.0–63.2) (67.8–70.2) (66.4–68.0) (44.6–51.6)	% point difference 1999 to 2001 -1.3 -3.4	% 55.9 66.1	(95% CI)	% point difference 1999 to 2001
65–74 ≥75  Race/Ethnicity White, non-Hispanic Black, non-Hispanic Hispanic Other <sup>†</sup> Sex Men Women  Region <sup>§</sup> New England Mid Atlantic Northeast Central	69.1 67.1 48.1 55.2	(67.8–70.2) (66.4–68.0)			,	
65–74 ≥75  Race/Ethnicity White, non-Hispanic Black, non-Hispanic Hispanic Other <sup>†</sup> Sex Men Women  Region <sup>§</sup> New England Mid Atlantic Northeast Central	69.1 67.1 48.1 55.2	(67.8–70.2) (66.4–68.0)			,	
Race/Ethnicity White, non-Hispanic Black, non-Hispanic Hispanic Other <sup>†</sup> Sex Men Women  Region <sup>§</sup> New England Mid Atlantic Northeast Central	67.1 48.1 55.2	(67.8–70.2) (66.4–68.0)	-3.4	66.1	(0.4.0. 07.1)	6.0
White, non-Hispanic Black, non-Hispanic Hispanic Other <sup>†</sup> Sex Men Women  Region <sup>§</sup> New England Mid Atlantic Northeast Central	48.1 55.2	` ,			(64.8–67.4)	5.2
White, non-Hispanic Black, non-Hispanic Hispanic Other <sup>†</sup> Sex Men Women  Region <sup>§</sup> New England Mid Atlantic Northeast Central	48.1 55.2	` ,				
Black, non-Hispanic Hispanic Other <sup>†</sup> Sex Men Women  Region <sup>§</sup> New England Mid Atlantic Northeast Central	48.1 55.2	` ,	-1.8	63.5	(62.6-64.4)	6.6
Hispanic Other <sup>†</sup> Sex Men Women  Region <sup>§</sup> New England Mid Atlantic Northeast Central	55.2		0.0	39.4	(36.0–43.0)	3.1
Other†  Sex  Men  Women  Region§  New England  Mid Atlantic  Northeast Central		(49.0–61.4)	-3.4	41.6	(35.8–47.4)	7.0
Sex Men Women Region§ New England Mid Atlantic Northeast Central		(58.4–73.0)	-2.5	45.1	(37.8–52.4)	-6.6
Men Women  Region§  New England Mid Atlantic Northeast Central		(5511 1515)			(5115 5=11)	
Women  Region§  New England Mid Atlantic  Northeast Central	66.6	(65.2–68.0)	-1.5	58.7	(57.2-60.2)	5.2
Region§ New England Mid Atlantic Northeast Central	63.7	(62.6–64.8)	-2.4	60.9	(59.8–62.0)	6.3
New England Mid Atlantic Northeast Central	00.7	(02.0 01.0)		00.0	(00.0 02.0)	0.0
Mid Atlantic Northeast Central	70.4	(60 0 70 0)	1.4	64.0	(60 A 65 6)	8.8
Northeast Central	63.3	(68.8–72.0)	-0.6	64.0 57.7	(62.4–65.6)	5.9
	63.9	(60.8–65.8) (61.8–65.8)	-0.6 -4.1	57.7 59.3	(55.2–60.4) (57.2–61.4)	5.6
Northwoot Control	69.7	(68.0–71.4)	-4.1 2.1	61.1	(59.2–62.8)	6.6
Northwest Central Southern Atlantic	60.8	,	2.1 –2.4	59.9	,	5.7
Southeast Central	64.0	(59.0–62.4)	-2.4 -1.6	56.7	(58.2–61.6)	3.6
Southwest Central	62.5	(61.8–66.2) (60.4–64.8)	-1.0 -5.9	58.0	(54.4–59.0) (55.8–60.2)	5.3
Mountain	67.9	(65.8–70.2)	-3.9 -3.2	66.0	(63.8–68.4)	8.1
Pacific	69.9	(66.6–73.2)	-3.2 -1.1	61.8	(58.2–65.2)	5.1
	03.3	(00.0-73.2)	-1.1	01.0	(30.2-03.2)	5.1
Education level	E0.0	(FC 0, CO 4)	0.0	E0 0	(E1 O EE 4)	6.4
<high school<="" td=""><td>58.3 64.1</td><td>(56.2–60.4)</td><td>−2.2 −1.8</td><td>53.3 60.2</td><td>(51.2–55.4)</td><td>6.4</td></high>	58.3 64.1	(56.2–60.4)	−2.2 −1.8	53.3 60.2	(51.2–55.4)	6.4
High school graduate	68.8	(62.8–65.4)	-1.6 -2.7		(58.8–61.6)	6.4 4.4
>High school	00.0	(67.6–70.0)	-2.7	63.1	(61.8–64.4)	4.4
Self-reported health		(54.0.00.0)	2.2	50.0	(40.0.50.0)	0.0
Excellent	57.6	(54.6–60.6)	-3.6	50.9	(48.0–53.8)	6.2
Very good	64.0	(62.4–65.6)	-3.4	57.0	(55.4–58.8)	2.6
Good	66.6	(65.2–68.0)	-0.8	61.9	(60.4–63.4)	6.7
Fair	66.0	(64.2–67.8)	-2.5	63.2	(61.4–65.2)	6.7
Poor	69.3	(67.0–71.8)	0.0	66.9	(64.4–69.4)	9.0
Diabetes						
Yes	70.6	(68.8–72.6)	-1.9	66.1	(64.0-68.2)	6.8
No	63.9	(63.0-64.8)	-2.1	58.9	(58.0–59.8)	5.6
Asthma**						
Yes	70.7	(67.8-73.6)	NA <sup>††</sup>	72.7	(69.6-75.6)	NA <sup>††</sup>
No	64.3	(63.4-65.2)	NA <sup>††</sup>	58.7	(57.8-59.6)	NA <sup>††</sup>
Smoking status						
Ever smoked	66.3	(65.2-67.4)	-1.5	62.6	(61.4-63.6)	6.8
Never smoked		'				
Total	63.6	(62.4-64.8)	-2.6	57.4	(56.2 - 58.8)	4.7

<sup>\*</sup> Confidence interval.

Numbers for other racial/ethnic groups were too small for meaningful analysis.

New England=Connecticut, Maine, Massachusetts, New Hampshire, Rhode Island, and Vermont; Mid Atlantic=New Jersey, New York, and Pennsylvania; Northeast Central=Illinois, Indiana, Michigan, Ohio, and Wisconsin; Northwest Central=Iowa, Kansas, Minnesota, Missouri, Nebraska, North Dakota, and South Dakota; Southern Atlantic=Delaware, District of Columbia, Florida, Georgia, Maryland, North Carolina, South Carolina, Virginia, and West Virginia; Southeast Central=Alabama, Kentucky, Mississippi, and Tennessee; Southwest Central=Arkansas, Louisiana, Oklahoma, and Texas; Mountain=Arizona, Colorado, Idaho, Montana, Nevada, New Mexico, Utah, and Wyoming; Pacific=Alaska, California, Hawaii, Oregon, and Washington.

Based on response to the question, "Have you ever been told by a doctor that you have diabetes?"

<sup>\*\*</sup> Based on response to the question, "Have you ever been told by a doctor, nurse, or other health professional that you had asthma?"

<sup>††</sup> Not available. Questions about asthma were not included on the core section of the 1999 BRFSS.

TABLE 2. Percentage of persons aged ≥65 years who reported receiving influenza vaccine during the preceding year or pneumococcal vaccine ever, by reporting area — Behavioral Risk Factor Surveillance System (BRFSS), United States, 2001

	Donavioral Filott I	Influenza	e System (BRFSS	,, Officed States		
		Influenza	% point		Pneumococcal	% point
			difference			difference
Reporting area	%	(95% CI*)	1999 to 2001	%	(95% CI)	1999 to 2001
Alabama	65.9	(61.6-70.2)	1.3	60.3	(55.8-64.8)	6.4
Alaska	62.8	(54.0-71.4)	3.0	65.3	(56.8 - 74.0)	21.6
Arizona	61.8	(56.8-66.8)	-9.5	65.6	(60.8-70.6)	12.2
Arkansas	63.2	(59.0-67.4)	-4.0	59.0	(54.6-63.4)	8.9
California	68.9	(64.6 - 73.4)	-3.3	59.6	(55.0-64.2)	2.6
Colorado	77.4	(72.0 - 82.6)	2.6	68.6	(62.6-74.6)	5.9
Connecticut	69.1	(66.2-71.8)	4.3	63.3	(60.4-66.2)	14.3
Delaware	67.6	(63.6-71.8)	-0.1	68.9	(64.8 - 73.2)	2.4
District of Columbia	55.5	(49.0-62.0)	-0.4	49.0	(42.4-55.6)	13.7
Florida	54.9	(51.6-58.2)	-8.4	58.1	(54.8-61.4)	4.5
Georgia	62.2	(58.0-66.6)	5.3	57.9	(53.4-62.4)	8.2
Guam	39.5	(25.6–53.4)	$NA^{\dagger}$	33.1	(19.4–46.6)	NA <sup>†</sup>
Hawaii	79.0	(75.4-82.4)	4.9	63.7	(59.2-68.2)	7.9
Idaho	65.1	(61.6–68.6)	-3.9	60.3	(56.6–64.0)	5.1
Illinois	62.2	(57.0–67.4)	-5.3	56.7	(51.2–62.0)	9.3
Indiana	65.7	(62.0–69.4)	-0.4	60.2	(56.4–64.2)	8.6
Iowa	72.8	(69.4–76.2)	3.2	65.9	(62.2–69.6)	4.6
Kansas	68.5	(65.2–71.8)	1.5	62.9	(59.4–66.4)	7.8
Kentucky	60.9	(57.4–64.4)	-7.4	55.1	(51.6–58.6)	3.1
Louisiana	56.1	(52.4–59.8)	-4.3	49.5	(45.8–53.2)	9.1
Maine	71.5	(67.2–75.8)	-2.2	65.0	(60.4–69.6)	7.7
Maryland	67.3	(63.0–71.6)	4.7	62.3	(57.8–66.8)	8.1
Massachusetts	70.6	(68.0–73.4)	1.3	63.5	(60.6–66.4)	6.8
Michigan	60.4	(56.4–64.6)	-9.6	56.6	(52.2–60.8)	-1.2
Minnesota	70.1	(66.6–73.6)	6.1	62.9	(59.2–66.6)	11.0
Mississippi	61.8	(57.4–66.2)	-1.0	55.7	(51.2–60.2)	5.3
Missouri	67.5	(63.2–71.6)	-0.9	56.0	(51.6–60.4)	3.2
Montana	73.1	(69.0–77.2)	0.2	67.9	(63.4–72.2)	6.7
Nebraska	70.1	(66.6–73.6)	0.9	61.2	(57.4–65.0)	6.3
Nevada	63.3	(57.2–69.4)	1.2	66.3	(60.2–72.6)	4.6
New Hampshire	69.4	(65.6–73.2)	4.3	62.7	(58.6–66.6)	2.3
New Jersey	64.5	(61.0–68.0)	-0.9	58.9	(55.2–62.6)	3.9
New Mexico	70.0	(66.4–73.6)	-0.9 1.2	62.7	(58.8–66.6)	9.5
New York		,				
	62.5	(58.0–67.0)	-1.3 1.0	55.9	(51.2–60.6)	5.9
North Carolina	66.1	(62.2–70.0)	1.9	65.8	(61.8–69.6)	7.2
North Dakota	70.0	(65.4–74.6)	2.8	64.2	(59.4–69.0)	9.1
Ohio	63.4	(59.0–67.8)	-5.4	59.3	(54.8–63.8)	4.4
Oklahoma	72.7	(69.2–76.2)	0.8	66.1	(62.4–69.8)	12.4
Oregon	71.7	(67.4–76.0)	6.5	70.9	(66.4–75.2)	14.6
Pennsylvania	63.8	(60.0–67.4)	0.7	59.5	(55.6–63.2)	7.2
Puerto Rico	36.8	(32.6–41.0)	-3.5	24.1	(20.2–28.0)	2.3
Rhode Island	72.6	(69.0–76.2)	-3.2	67.0	(63.2–70.8)	10.1
South Carolina	66.2	(61.8–70.6)	-3.8	57.9	(53.2–62.6)	1.8
South Dakota	74.1	(71.4–76.6)	0.4	59.2	(56.2–62.2)	8.8
Tennessee	65.6	(61.0–70.2)	0.1	55.4	(50.6–60.2)	1.1
Texas	61.8	(58.6–65.0)	-8.1	58.0	(54.6–61.4)	2.2
Utah	68.7	(63.2–74.0)	-6.5	67.3	(62.4–72.4)	6.0
Vermont	71.5	(68.0–75.2)	-1.9	67.3	(63.4–71.2)	10.8
Virgin Islands	38.7	(31.4–46.0)	NA <sup>†</sup>	30.7	(23.8–37.6)	NA <sup>†</sup>
Virginia	65.3	(60.6–70.0)	-0.4	60.1	(55.2–65.0)	4.9
Washington	72.5	(69.0-76.0)	3.6	66.8	(63.0-70.6)	10.9
West Virginia	61.7	(57.8–65.4)	-1.2	61.3	(57.6-65.2)	7.0
Wisconsin	70.4	(66.2-74.6)	5.5	65.6	(61.0-70.0)	11.9
Wyoming	69.6	(65.4–73.8)	-4.2	68.4	(64.0-72.8)	6.9
Total	64.9	(64.0-65.8)	-2.0	60.0	(59.2-60.8)	5.9

<sup>\*</sup>Confidence interval.

† Not available. Guam and Virgin Islands did not participate in the 1999 BRFSS.

TABLE 3. Odds ratios (ORs) and corresponding p values determined by logistic regression for persons aged ≥65 years who reported receiving influenza vaccine during the preceding year or pneumococcal vaccine ever, by selected characteristics — Behavioral Risk Factor Surveillance System, United States, 2001

		Influenza		Pneumococcal					
Characteristic	OR	(95% CI*)	p-value	OR	(95% CI)	p-value			
Age group (yrs)									
65–74 <sup>†</sup>	1.00			1.00					
<u>≥</u> 75	1.40	(1.29-1.50)	< 0.0001	1.52	(1.41-1.64)	< 0.0001			
Race/Ethnicity									
White, non-Hispanic <sup>†</sup>	1.00			1.00					
Black, non-Hispanic	0.50	(0.43 - 0.59)	< 0.0001	0.39	(0.33 - 0.45)	< 0.0001			
Hispanic	0.63	(0.50-0.79)	0.0001	0.42	(0.34–0.54)	< 0.0001			
Other§	0.86	(0.62–1.19)	0.3606	0.44	(0.32-0.59)	< 0.0001			
Sex									
Men <sup>†</sup>	1.00			1.00					
Women	0.89	(0.82-0.96)	0.0049	1.15	(1.06-1.24)	0.0006			
Region <sup>1</sup>		(			,				
New England <sup>†</sup>	1.00			1.00					
Mid Atlantic	0.80	(0.70-0.91)	0.0008	0.86	(0.75-0.98)	0.0230			
Northeast Central	0.78	(0.70–0.88)	0.0001	0.87	(0.77–0.98)	0.0214			
Northwest Central	0.98	(0.87–1.10)	0.7111	0.90	(0.80–1.00)	0.0462			
Southern Atlantic	0.71	(0.64–0.80)	<0.0001	0.96	(0.86–1.07)	0.4887			
Southeast Central	0.84	(0.74–0.95)	0.0067	0.82	(0.73–0.94)	0.0028			
Southwest Central	0.77	(0.68–0.88)	0.0001	0.90	(0.79–1.02)	0.0890			
Mountain	0.90	(0.79–1.02)	0.1106	1.16	(1.02–1.32)	0.0259			
Pacific	1.07	(0.91–1.26)	0.4300	1.08	(0.93–1.26)	0.3224			
Education level									
<high school<sup="">†</high>	1.00			1.00					
High school graduate	1.26	(1.14-1.40)	< 0.0001	1.30	(1.17-1.45)	< 0.0001			
>High school	1.58	(1.41–1.76)	< 0.0001	1.51	(1.36–1.68)	< 0.0001			
Self-reported health		,			,				
Excellent <sup>†</sup>	1.00			1.00					
Very good	1.32	(1.16–1.51)	< 0.0001	1.27	(1.12-1.44)	0.0002			
Good	1.51	(1.32–1.72)	<0.0001	1.61	(1.42–1.82)	< 0.0001			
Fair	1.52	(1.31–1.77)	< 0.0001	1.72	(1.49–1.97)	< 0.0001			
Poor	1.81	(1.52–2.16)	< 0.0001	2.02	(1.71–2.40)	< 0.0001			
Diabetes**		/			,				
Yes	1.40	(1.25–1.56)	<0.0001	1.38	(1.23–1.54)	<0.0001			
No <sup>†</sup>	1.00	(1.20 1.00)	30.0001	1.00	(1.20 1.04)	30.0001			
Asthma <sup>††</sup>	1.50								
Yes	1.40	(1.21–1.61)	<0.0001	1.86	(1.61–2.16)	<0.0001			
No <sup>†</sup>	1.00	(1.21-1.01)	<b>\0.0001</b>	1.00	(1.01-2.10)	<0.0001			
	1.00			1.00					
Smoking status	4.05	(0.07.4.44)	0.0000	4.00	(1.10, 1.00)	-0.0004			
Ever smoked	1.05	(0.97–1.14)	0.2003	1.22	(1.13–1.32)	<0.0001			
Never smoked <sup>†</sup>	1.00			1.00					

<sup>\*</sup> Confidence interval.

Reference level for characteristic.

Numbers for other racial/ethnic groups were too small for meaningful analysis.

New England=Connecticut, Maine, Massachusetts, New Hampshire, Rhode Island, and Vermont; Mid Atlantic=New Jersey, New York, and Pennsylvania; Northeast Central=Illinois, Indiana, Michigan, Ohio, and Wisconsin; Northwest Central=Iowa, Kansas, Minnesota, Missouri, Nebraska, North Dakota, and South Dakota; Southern Atlantic=Delaware, District of Columbia, Florida, Georgia, Maryland, North Carolina, South Carolina, Virginia, and West Virginia; Southeast Central=Alabama, Kentucky, Mississippi, and Tennessee; Southwest Central=Arkansas, Louisiana, Oklahoma, and Texas; Mountain=Arizona, Colorado, Idaho, Montana, Nevada, New Mexico, Utah, and Wyoming; Pacific=Alaska, California, Hawaii, Oregon, and Washington.

<sup>\*\*</sup> Based on response to the question, "Have you ever been told by a doctor that you have diabetes?"

<sup>††</sup> Based on response to the question, "Have you ever been told by a doctor, nurse, or other health professional that you had asthma?"

The association between vaccination status and additional variables was examined by multivariable logistic regression (Table 3). Persons aged ≥75 years were more likely to report influenza or pneumococcal vaccination than persons aged 65–74 years. Men were more likely than women to report influenza vaccination and less likely to report pneumococcal vaccination. Persons with diabetes or asthma were significantly more likely to report influenza and pneumococcal vaccination, compared with those who did not have diabetes or asthma. Coverage with both vaccines increased as education level increased and as self-reported health declined. Pneumococcal vaccination coverage was higher among smokers than among nonsmokers.

**Reported by:** A MacNeil, MPH, Association of Schools of Public Health, Atlanta, Georgia. JA Singleton, MS, JS Moran, MD, Epidemiology and Surveillance Div, National Immunization Program, CDC.

Editorial Note: The findings in this report indicate that the estimated prevalences of influenza and pneumococcal vaccinations were <80% among persons ≥65 years in all reporting areas. National influenza vaccination coverage for persons aged ≥65 years increased linearly during 1993–1997, leveled off by 1999, and decreased during 1999–2001. The 2001 coverage is slightly below coverage reported in 1997. The decrease in influenza vaccine coverage might be due, in part, to delays in influenza vaccine distribution during the 2000–01 influenza season and the less severe distribution delays during the 2001–02 season (5).

Pneumococcal vaccination coverage among persons aged ≥65 years increased linearly during 1993–2001 and was significantly above 60% in 18 states in 2001. The number of states with point prevalence estimates of ≥60% increased from eight in 1999 to 32 in 2001. However, coverage in all 54 reporting areas remained <90% and must increase substantially to meet the national health objective for 2010.

Previous reports have noted racial/ethnic disparities in adult vaccine coverage (4). In the 2001 BRFSS, non-Hispanic blacks and Hispanics had substantially lower coverage than non-Hispanic whites. After adjusting for known potential confounding factors measured by BRFSS (e.g., education level but not direct measures of access to care, which were not available), the odds of members of these populations receiving influenza or pneumococcal vaccine remained substantially lower. These gaps were greatest for pneumococcal vaccine. In comparison with influenza vaccine, which is recommended annually, a single dose of pneumococcal vaccine is needed for persons aged ≥65 years. Strategies for addressing these disparities will be investigated by CDC's Racial and Ethnic Adult Disparities Immunization Initiative (READII) through 2-year demonstration projects in Chicago, Illinois; Milwaukee,

Wisconsin; a rural area of Mississippi; Rochester, New York; and San Antonio, Texas. Local and state health departments in these areas will work with community partners, CDC, and other federal agencies to identify and implement effective ways to improve influenza and pneumococcal vaccination levels among older non-Hispanic blacks and Hispanics.

Health-care providers should assess the vaccination status of their patients and offer indicated vaccines. Annual influenza vaccination provides such an opportunity; persons reporting recent influenza vaccination were 2.5 times more likely to report having received pneumococcal vaccine than were persons who did not report recent influenza vaccination. Administration of influenza and pneumococcal vaccine simultaneously does not increase the incidence or severity of adverse reactions (6). Nevertheless, approximately one fourth of persons reporting recent influenza vaccination did not report having ever received pneumococcal vaccine.

The findings in this report are subject to at least three limitations. First, receipt of influenza or pneumococcal vaccination was based on self-report and not validated. The validity of self-reported pneumococcal vaccination is lower than that of influenza vaccination (7). Second, the BRFSS excludes persons without telephones or those with only cellular telephones. Third, the BRFSS response rate was >60% in 10 of the 54 reporting areas.

To assess possible selection bias resulting from the two latter limitations, comparisons were made between national estimates of vaccination coverage from BRFSS and the National Health Interview Survey (NHIS). NHIS data are collected through household, face-to-face interviews and usually have higher response rates (e.g., 72.1% in 2000). Estimated influenza vaccination levels for persons aged ≥65 years in 1997, 1999, and 2001 were 63.2%, 65.7%, and 63.0%, respectively, from NHIS and 65.5%, 66.9%, and 64.9%, respectively, from BRFSS. For the same years, estimated pneumococcal vaccination levels were 42.4%, 49.7%, and 53.8%, respectively, from NHIS and 45.4%, 54.1%, and 60.0%, respectively, from BRFSS. National BRFSS vaccination estimates show similar trends and subgroup differences as NHIS estimates but are consistently slightly higher than NHIS estimates. Previous analysis has documented that NHIS respondents living in households without telephones were less likely to report being vaccinated than those living in households with telephones (4), but this accounts for only a small portion of the differences observed between NHIS and BRFSS estimates.

The optimal time to administer influenza vaccination is during October–November. However, influenza vaccination should continue into December and later because many persons at high risk for influenza-related complications,

household members of these persons, health-care workers, and other persons who want to decrease their risk for influenza remain unvaccinated by the end of November (1). Current projections indicate that 93 million doses of influenza vaccine will be available during the 2002-03 influenza season, and several million doses remain available for purchase. To maximize coverage among target groups and overall use, physicians should offer influenza vaccine throughout the influenza season. Influenza activity peaked in January or later in 21 of the preceding 25 influenza seasons (1). During influenza season and all year, pneumococcal vaccination also should be offered to persons aged ≥65 years and others at high risk who have not been vaccinated or whose vaccination status is unknown. Physicians can improve coverage by using strategies such as improved record keeping, standing orders, reminder/recall systems, and offering vaccinations to hospitalized patients before discharge (8,9). Additional information about influenza and pneumococcal vaccination is available at http://www.cdc.gov/nip.

## **Acknowledgment**

This report is based on data contributed by state BRFSS coordinators.

#### References

- CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices. MMWR 2002;51(No. RR-3).
- Robinson KA, Baughman W, Rothrock G, et al. Epidemiology of invasive Streptococcus pneumoniae infections in the United States, 1995–1998: opportunities for prevention in the conjugate vaccine era. JAMA 2001;285:1729–35.
- U.S. Department of Health and Human Services. Healthy people 2010, 2nd ed. With understanding and improving health and objectives for improving health (2 vols.). Washington, DC: U.S. Department of Health and Human Services, 2000.
- CDC. Influenza and pneumococcal vaccination levels among adults aged ≥65 years—United States, 1999. MMWR 2001;50:532–7.
- Fukuda K, O'Mara D, Singleton J. Part 4: How the delayed distribution of influenza vaccine created shortages in 2000 and 2001. Pharmacy and Therapeutics 2002;27:235–42.
- CDC. General recommendation on immunization: recommendations of the Advisory Committee on Immunization Practices (ACIP) and the American Academy of Family Physicians (AAFP). MMWR 2002;51(No. RR-2).
- MacDonald R, Baken L, Nelson A, Nichol K. Validation of self-report of influenza and pneumococcal vaccination status in elderly outpatients. Am J Prev Med 1999;16:173–7.
- Mieczkowski A, Wilson S. Adult pneumococcal vaccination: a review of physician and patient barriers. Vaccine 2002;20:1383–92.
- Task Force on Community Preventive Services. Recommendations regarding interventions to improve vaccination coverage in children, adolescents, and adults. Am J Prev Med 2000;18(suppl 1):S92–S96.

## Notice to Readers

# Use of Anthrax Vaccine in Response to Terrorism: Supplemental Recommendations of the Advisory Committee on Immunization Practices

In December 2000, the Advisory Committee on Immunization Practices (ACIP) released its recommendations for using anthrax vaccine in the United States (1). Because of recent terrorist attacks involving the intentional exposure of U.S. civilians to *Bacillus anthracis* spores and concerns that the current anthrax vaccine supply is limited, ACIP developed supplemental recommendations on using anthrax vaccine in response to terrorism. These recommendations supplement the previous ACIP statement in three areas: use of anthrax vaccine for pre-exposure vaccination in the U.S. civilian population, the prevention of anthrax by postexposure prophylaxis (PEP), and recommendations for additional research related to using antimicrobial agents and anthrax vaccine for preventing anthrax.

## Use of Anthrax Vaccine for Pre-Exposure Vaccination

In December 2001, the U.S. Department of Health and Human Services obtained a limited supply of anthrax vaccine (BioThrax [formerly Anthrax Vaccine Adsorbed (AVA)], BioPort, Lansing, Michigan), allowing ACIP to reconsider using anthrax vaccine in the U.S. civilian population. ACIP reaffirms that pre-exposure use of anthrax vaccine should be based on a quantifiable risk for exposure (1). ACIP recommends that groups at risk for repeated exposures to B. anthracis spores should be given priority for pre-exposure vaccination. Groups at risk for repeated exposure include laboratory personnel handling environmental specimens (especially powders) and performing confirmatory testing for B. anthracis in the U.S. Laboratory Response Network (LRN) for Bioterrorism Level B laboratories or above, workers who will be making repeated entries into known B. anthracis-sporecontaminated areas after a terrorist attack (2), and workers in other settings in which repeated exposure to aerosolized B. anthracis spores might occur. Laboratory workers using standard Biosafety Level 2 practices in the routine processing of clinical samples or environmental swabs (Level A laboratories [3]) are not considered by ACIP to be at increased risk for exposure to *B. anthracis* spores.

For persons not at risk for repeated exposures to aerosolized *B. anthracis* spores through their occupation, pre-exposure vaccination with anthrax vaccine is not recommended. For the general population, prevention of morbidity and mortality

associated with anthrax will depend on public vigilance, early detection and diagnosis, appropriate treatment, and PEP.

## **Prevention of Anthrax by PEP**

Because of a potential preventive benefit of combined antimicrobial PEP and vaccine and the availability of a limited supply of anthrax vaccine for civilian use, ACIP endorses CDC making anthrax vaccine available in a 3-dose regimen (0, 2, 4 weeks) in combination with antimicrobial PEP under an Investigational New Drug (IND) application with the Food and Drug Administration for unvaccinated persons at risk for inhalational anthrax. However, anthrax vaccine is not licensed for postexposure use in preventing anthrax.

Use of anthrax vaccine for PEP could have additional benefits, including reducing the need for long-term antimicrobial therapy with its associated problems of nonadherence and possible adverse events. After the anthrax-related terrorist attacks in 2001, approximately 10,000 persons were recommended to receive a 60-day regimen of antimicrobial prophylaxis for suspected or confirmed exposure to *B. anthracis* spores, but adherence to the recommended 60-day antibiotic regimens was as low as 42% (4). In addition, because studies of the 2001 terrorist attacks suggest that some persons might be exposed to *B. anthracis* spores in excess of those studied in animal models, the effectiveness of antimicrobial prophylaxis in such persons is unclear (4). However, no cases of anthrax have been detected among persons recommended to take antimicrobial prophylaxis after the terrorist attacks of 2001.

The provision of anthrax vaccine for PEP under an IND application should provide an opportunity to reduce the risk to the greatest extent possible with current medical knowledge and might provide data to support developing additional recommendations for preventing anthrax. To better document the immunogenicity of anthrax vaccine in the postexposure setting, ACIP encouraged CDC to obtain serologic testing on a subset of vaccinees.

ACIP recommended previously that if antimicrobial therapy is used alone for postexposure prevention of anthrax, at least a 30-day course of treatment should be provided. Previous recommendations noted that longer courses (42–60 days) might be indicated. On the basis of limited data from both unintentional human exposures and animal studies (5–7), ACIP now recommends that the duration of postexposure antimicrobial prophylaxis should be 60 days if used alone for PEP of unvaccinated exposed persons.

Data are insufficient to clarify the duration of antimicrobial use in combination with vaccine for PEP against anthrax. Antibody titers among vaccinated persons peak at 14 days after the third dose (8). If antimicrobial prophylaxis is

administered in combination with postexposure vaccination, it might be prudent to continue antibiotics until 7–14 days after the third vaccine dose.

Few data exist about the effectiveness of postexposure antimicrobial prophylaxis among exposed persons who have been partially or fully vaccinated. In the only human clinical trial of anthrax vaccine, cases occurred among participants who had received <4 doses (9). Recognizing these limited data, but considering a potential undefined benefit, ACIP recommends that persons who have been partially or fully vaccinated receive at least a 30-day course of antimicrobial PEP and continue with the licensed vaccination regimen. Antimicrobial PEP is not needed for vaccinated persons working in Biosafety Level 3 laboratories under recommended conditions (10) nor for vaccinated persons (six vaccinations according to the current label) wearing appropriate personal protective equipment (PPE) while working in contaminated environments in which inhalational exposure to *B. anthracis* spores is a risk, unless their respiratory protection is disrupted.

## **Additional Considerations**

For most occupational settings, recommendations about anthrax vaccine and antimicrobial PEP might be implemented in combination with use of appropriate PPE (2). In addition to receiving PEP for preventing anthrax, potentially exposed persons should be observed for signs of febrile illness. CDC has published guidelines on clinical evaluation of persons with possible anthrax, including antimicrobial treatment (1,2). Because the current vaccine supply is limited, ACIP recommends expanded and intensive efforts to improve anthrax vaccine production.

## **Recommendations for Additional Research**

Because of the absence of data to guide public health recommendations in these critical areas, ACIP recommends studies on the safety and immunogenicity of anthrax vaccine for use in children, additional studies on the safety of anthrax vaccine during human pregnancy, and reproductive toxicology studies on anthrax vaccine in laboratory animals. To strengthen public health recommendations for PEP, ACIP recommends expanded animal studies to evaluate further the effectiveness of antimicrobial prophylaxis with and without anthrax vaccine, define the optimal duration of antimicrobial PEP for the prevention of inhalational anthrax, and evaluate alternative antimicrobial PEP regimens. Additional research also should be directed toward developing an improved vaccine for preventing anthrax and new therapeutic strategies, including use of antitoxin (e.g., hyperimmune globulin) for treating anthrax.

#### References

- CDC. Use of anthrax vaccine in the United States: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 2000;49(No. RR-15).
- CDC. Occupational health guidelines for remediation workers at Bacillus anthracis—contaminated sites—United States, 2001–2002. MMWR 2002;51;786–9.
- 3. CDC. Biological and chemical terrorism: strategic plan for preparedness and response: recommendations of the CDC Strategic Planning Workgroup. MMWR 2000;49(No. RR-4).
- 4. Shepard CW, Soriano-Gabarro M, Zell ER, et al. Antimicrobial postexposure prophylaxis for anthrax: adverse events and adherence. Emerg Infect Dis 2002;8:1124–32.
- Meselson M, Guillemin J, Hugh-Jones M, et al. 1994. The Sverdlosk anthrax outbreak of 1979. Science 1994;226:1202–7.
- Friedlander AM, Welkos SL, Pitt ML, et al. Postexposure prophylaxis against experimental inhalation anthrax. J Infect Dis 1993;167:1239– 42.
- 7. Henderson DW, Peacock S, Belton FC. Observations on the prophylaxis of experimental pulmonary anthrax in the monkey. J Hyg 1956;54:28–36.
- Pittman PR, Kim-Ahn G, Pifat DY, et al. Anthrax vaccine: immunogenicity and safety of a dose-reduction, route-change comparison study in humans. Vaccine 2002;20:1412–20.
- Brachman PS, Gold H, Plotkin SA, Fekety FR, Werrin M, Ingraham NR. Evaluation of human anthrax vaccine. Am J Public Health 1962;52:632–45.
- CDC. Biosafety in microbial and biomedical laboratories, 5th ed. In: Richmond JY, McKinney RW, eds. Washington, DC: U.S. Department of Health and Human Services, CDC, 2001.

# West Nile Virus Activity — United States, November 7–13, 2002

This report summarizes West Nile virus (WNV) surveillance data reported to CDC through ArboNET and by states and other jurisdictions as of 8 a.m. Mountain Standard Time, November 13, 2002.

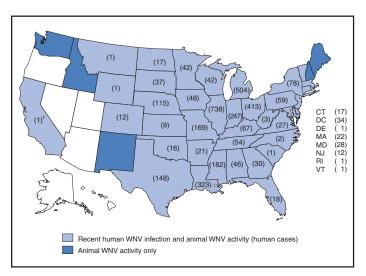
During November 7–13, a total of 80 laboratory-positive human cases of WNV-associated illness were reported from Michigan (n=21), Illinois (n=19), the District of Columbia (n=seven), Alabama (n=five), Missouri (n=four), New York (n=four), Kansas (n=three), Maryland (n=three), Virginia (n=three), Wisconsin (n=three), Colorado (n=two), Louisiana (n=two), Tennessee (n=two), Montana (n=one), and New Jersey (n=one). During this period, Montana reported its first-ever human case of WNV infection. Also, during the same period, WNV infections were reported in 210 dead crows and 294 other dead birds. A total of 169 veterinary cases and 79 WNV-positive mosquito pools were reported.

During 2002, a total of 3,587 human cases with laboratory evidence of recent WNV infection have been reported from

Illinois (n=738), Michigan (n=504), Ohio (n=413), Louisiana (n=323), Indiana (n=247), Mississippi (n=182), Missouri (n=169), Texas (n=148), Nebraska (n=115), New York (n=78), Kentucky (n=67), Pennsylvania (n=59), Tennessee (n=54), Iowa (n=48), Alabama (n=46), Minnesota (n=42), Wisconsin (n=42), South Dakota (n=37), the District of Columbia (n=34), Georgia (n=30), Maryland (n=28), Virginia (n=27), Massachusetts (n=22), Arkansas (n=21), Florida (n=18), Connecticut (n=17), North Dakota (n=17), Oklahoma (n=16), Colorado (n=12), New Jersey (n=12), Kansas (n=nine), West Virginia (n=three), North Carolina (n=two), California (n=one), Delaware (n=one), Montana (n=one), Rhode Island (n=one), South Carolina (n=one), Vermont (n=one), and Wyoming (n=one) (Figure). Among the 3,226 patients for whom data were available, the median age was 56 years (range: 1.5 months-99 years); 1,719 (54%) were male, and the dates of illness onset ranged from June 10 to October 21. A total of 196 human deaths have been reported. The median age of decedents was 78 years (range: 24–99 years); 119 (61%) deaths were among men. In addition, 7,522 dead crows and 5,730 other dead birds with WNV infection were reported from 42 states and the District of Columbia; 8,312 WNV infections in mammals (8,299 equines, three canines, and 10 other species) have been reported from 37 states (Alabama, Arkansas, Colorado, Connecticut, Delaware, Florida, Georgia, Idaho, Illinois, Indiana, Iowa, Kansas, Kentucky, Louisiana, Maryland, Massachusetts, Minnesota, Mississippi, Missouri, Montana, Nebraska, New Jersey, New Mexico, New York, North Carolina, North Dakota, Ohio, Oklahoma, Pennsylvania, South Carolina, South Dakota, Tennessee, Texas, Vermont, Virginia, Wisconsin, and Wyoming). During 2002, WNV seroconversions have been reported in 366 sentinel chicken flocks from Florida, Iowa, Nebraska, North Carolina, Pennsylvania, Texas, and New York City; 4,906 WNV-positive mosquito pools have been reported from 27 states (Alabama, Arkansas, Connecticut, Delaware, Georgia, Illinois, Indiana, Iowa, Kansas, Kentucky, Maryland, Massachusetts, Mississippi, Missouri, Nebraska, New Hampshire, New Jersey, New York, North Carolina, Ohio, Pennsylvania, Rhode Island, South Carolina, South Dakota, Texas, Vermont, and Virginia), New York City, and the District of Columbia.

Additional information about WNV activity is available at http://www.cdc.gov/ncidod/dvbid/westnile/index.htm and http://www.cindi.usgs.gov/hazard/event/west\_nile/west\_nile.html.

FIGURE. Areas reporting West Nile virus (WNV) activity — United States, 2002\*



 $<sup>^{\</sup>star}_{}$  As of 8 a.m. Mountain Standard Time, November 13, 2002.  $^{\dagger}_{}$  California has reported human WNV activity only.

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending November 9, 2002, with historical data

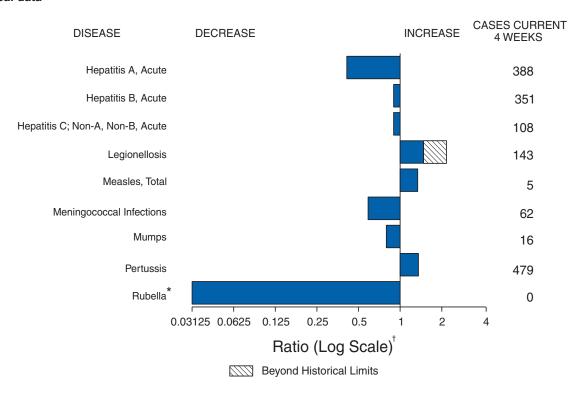


TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending November 9, 2002 (45th Week)\*

		Cum. 2002	Cum. 2001		Cum. 2002	Cum. 2001
Anthrax		2	21	Encephalitis: West Nile <sup>†</sup>	1,311	52
Botulism:	foodborne	12	33	Hansen disease (leprosy)†	58	60
	infant	47	85	Hantavirus pulmonary syndrome†	13	7
	other (wound & unspecified)	25	15	Hemolytic uremic syndrome, postdiarrheal <sup>†</sup>	170	158
Brucellosis†	, , , ,	68	112	HIV infection, pediatric <sup>†§</sup>	116	172
Chancroid		61	31	Plague	-	2
Cholera		5	4	Poliomyelitis, paralytic	-	-
Cyclosporiasi	S <sup>†</sup>	156	140	Psittacosis†	18	17
Diphtheria		1	2	Q fever <sup>†</sup>	40	22
Ehrlichiosis:	human granulocytic (HGE)†	300	194	Rabies, human	2	1
	human monocytic (HME)†	155	101	Streptococcal toxic-shock syndrome†	71	68
	other and unspecified	9	5	Tetanus	21	27
Encephalitis:	California serogroup viral†	114	107	Toxic-shock syndrome	99	104
·	eastern equine <sup>†</sup>	2	8	Trichinosis	12	21
	Powassan <sup>†</sup>	-	-	Tularemia <sup>†</sup>	54	120
	St. Louis <sup>†</sup>	8	76	Yellow fever	1	-
	western equine <sup>†</sup>	1	-			

<sup>-:</sup> No reported cases.

<sup>\*</sup> No rubella cases were reported for the current 4-week period yielding a ratio for week 45 of zero (0).

† 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.

<sup>\*</sup>Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TNot notifiable in all states.

Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP). Last update October 31, 2002.

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending November 9, 2002, and November 10, 2001 (45th Week)\*

(45th Week)*							Esch	erichia coli, E	nterohemorrha	gic
	ΔΙ	DS	Chlar	nydia <sup>†</sup>	Cryptos	poridiosis	015	57:H7		in Positive, p non-O157
Reporting Area	Cum. 2002§	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001
UNITED STATES	24,713	34,080	661,928	667,886	2,512	3,402	3,137	2,832	145	138
NEW ENGLAND	1,011	1,268	23,198	21,070	165	135	246	226	32	38
Maine	23	40	1,428	1,173	10	18	36	25	5	1
N.H. Vt.	20 8	31 13	1,360 815	1,198 527	29 31	15 31	31 12	31 13	1	3 1
vı. Mass.	519	654	9,511	8,947	60	51	110	109	9	10
R.I.	71	84	2,338	2,555	19	4	13	13	-	1
Conn.	370	446	7,746	6,670	16	16	44	35	17	22
MID. ATLANTIC	5,619	8,977	73,490	72,580	298	307	212	212	-	-
Upstate N.Y. N.Y. City	404 3,210	1,168 4,773	14,708 23,170	12,251 25,724	120 115	91 111	157 12	137 15	-	_
N.J.	925	1,509	10,290	11,934	10	18	43	60	-	-
Pa.	1,080	1,527	25,322	22,671	53	87	N	N	-	-
E.N. CENTRAL	2,494	2,499	111,777	124,101	803	1,490	766	729	17	11
Ohio Ind.	453 347	476 306	24,833 14,349	32,972 13,507	117 50	160 74	141 67	192 77	13 1	9
III.	1,170	1,110	31,418	37,391	85	475	164	162	-	-
Mich.	398	457	27,375	25,873	109	173	132	87	3	2
Wis.	126	150	13,802	14,358	442	608	262	211	-	-
W.N. CENTRAL	421	718	36,887	34,028	383	491	477	458	35	37
Minn. Iowa	90 54	118 80	8,235 4,761	7,109 4,296	198 42	166 80	153 115	185 75	30	28
Mo.	189	337	13,279	12,195	32	48	68	58	N	N
N. Dak.	1	2	740	878	20	13	15	19	-	2
S. Dak. Nebr.	3 43	23 72	1,884 2,456	1,557 2,832	28 47	6 175	39 54	41 59	2	6 1
Kans.	41	86	5,532	5,161	16	3	33	21	-	-
S. ATLANTIC	7,537	10,268	128,393	128,857	312	338	316	221	35	31
Del.	131	217	2,309	2,434	3	6	7	4	-	1
Md.	1,066	1,517	14,339	13,344	21 4	36	25	28	-	-
D.C. Va.	371 538	733 843	2,974 14,286	2,818 15,803	21	11 24	56	48	9	5
W. Va.	58	71	2,081	2,066	2	2	9	10	-	-
N.C.	555	778	21,423	18,468	32	27	102	46	-	-
S.C. Ga.	547 1,160	612 1,232	10,486 25,731	13,332 28,156	6 133	7 148	5 53	15 41	10	9
Fla.	3,111	4,265	34,764	32,436	90	77	59	29	16	16
E.S. CENTRAL	1,128	1,532	40,925	43,046	109	44	98	126	-	-
Ky.	173	299	7,681	7,820	8	5	30	63	-	-
Tenn. Ala.	483 197	488 378	13,911 11,034	12,598 12,206	52 42	12 13	43 18	36 16	-	-
Miss.	275	367	8,299	10,422	7	14	7	11	-	-
W.S. CENTRAL	2,696	3,435	92,369	93,127	36	120	63	180	-	_
Ark.	163	176	6,094	6,505	8	8	10	15	-	-
La. Okla.	693 133	699 204	16,704 9,496	16,053 9,202	5 17	7 14	2 21	7 30	-	-
Tex.	1,707	2,356	60,075	61,367	6	91	30	128	-	-
MOUNTAIN	790	1,175	40,694	39,922	146	221	332	264	18	15
Mont.	8	15	1,928	1,570	5	36	28	20	-	-
Idaho	18	19 3	2,171	1,710 712	29 9	21 7	47 14	64 9	8 2	3 2
Wyo. Colo.	6 157	262	802 11,790	11,476	51	39	86	84	4	6
N. Mex.	53	133	5,739	5,308	18	27	11	14	3	4
Ariz.	327 43	446	12,947 2,236	12,680	16	7 70	34 84	27 31	1	-
Utah Nev.	178	98 199	2,236 3,081	2,169 4,297	14 4	78 6	28	15	-	-
PACIFIC	3,017	4,208	114,195	111,155	260	256	627	416	8	6
Wash.	302	427	12,796	11,738	43	U	133	115	-	-
Oreg.	216	177	5,894	6,416	38	51	219	64	8	6
Calif. Alaska	2,416 17	3,525 19	88,649 3,121	87,222 2,273	176 1	201 1	230 7	216 4	-	-
Hawaii	66	60	3,735	3,506	2	3	38	17	-	-
Guam	2	11	-	354	-	-	N	N	-	-
P.R.	668	1,017	1,997	2,342	-	-	-	2	-	-
V.I. Amer. Samoa	66 U	2 U	125 U	130 U	U	U	- U	U	- U	- U
		0	_	Ŭ	0	Ü	0	0		0

N: Not notifiable. U: Unavailable. -: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands.

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

† Chlamydia refers to genital infections caused by *C. trachomatis*.

§ Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention. Last update October 31, 2002.

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending November 9, 2002, and November 10, 2001 (45th Week)\*

Reporting Area UNITED STATES NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr.	Enterohe Shiga Tox	ichia coli morrhagic in Positive, ogrouped  Cum. 2001  17  1  -  1  -  3  -  3  6  6	Giardiasis  Cum. 2002  14,638  1,468 185 40 124 738 138 243 3,114 1,080 1,108 306	Gono Cum. 2002 279,054 6,496 115 114 81 2,871 776 2,539 33,652 7,519	Cum. 2001 309,633 5,989 118 160 57 2,766 729 2,159	All Second 1,271 90 1 8 7 49 10	Ages, erotypes  Cum. 2001  1,261  93 2 4 3 40 5	Age <5 Serot B Cum. 2002 21	уре
UNITED STATES  NEW ENGLAND  Maine N.H.  Vt. Mass. R.I. Conn.  MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis.  W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	Cum. 2002  34  1	Cum. 2001  17  1  3 3 3 6	Cum. 2002 14,638 1,468 185 40 124 738 138 243 3,114 1,080 1,108 306	Cum. 2002 279,054 6,496 115 114 81 2,871 776 2,539 33,652 7,519	Cum. 2001 309,633 5,989 118 160 57 2,766 729 2,159	Cum. 2002 1,271 90 1 8 7 49 10	Cum. 2001 1,261 93 2 4 3 40	Cum. 2002	<b>Cum.</b> <b>2001</b> 21
UNITED STATES  NEW ENGLAND  Maine N.H.  Vt. Mass. R.I. Conn.  MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis.  W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	34 1 - - 1 - - - - - - - 1 1 1 1 - - - -	17 1 - - 1 - - - 3 - - 3	14,638 1,468 185 40 124 738 138 243 3,114 1,080 1,108 306	2002 279,054 6,496 115 114 81 2,871 776 2,539 33,652 7,519	309,633 5,989 118 160 57 2,766 729 2,159	1,271 90 1 8 7 49	1,261 93 2 4 3 40		21
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	1        11 10	1 - - 1 - - - 3 - - 3 - 3	1,468 185 40 124 738 138 243 3,114 1,080 1,108 306	6,496 115 114 81 2,871 776 2,539 33,652 7,519	5,989 118 160 57 2,766 729 2,159	90 1 8 7 49 10	93 2 4 3 40	21 - - - -	
Maine N.H. Vt. Mass. R.I. Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	- - 1 - - - - - - - 1 1 10	- - 1 - - - 3 - - - 3	185 40 124 738 138 243 3,114 1,080 1,108 306	115 114 81 2,871 776 2,539 33,652 7,519	118 160 57 2,766 729 2,159	1 8 7 49 10	2 4 3 40	- - - -	1 - -
N.H. Vt. Mass. R.I. Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	- - - - - - - 11 10	3 - - 3 - - 3 6	40 124 738 138 243 3,114 1,080 1,108 306	114 81 2,871 776 2,539 33,652 7,519	160 57 2,766 729 2,159	8 7 49 10	4 3 40	- - -	-
Mass. R.I. Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	- - - - - - - 11 10	3 - - 3 - - 3 6	738 138 243 3,114 1,080 1,108 306	2,871 776 2,539 33,652 7,519	2,766 729 2,159	49 10	40	-	
R.I. Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	10 - -	- - - 3 6	138 243 3,114 1,080 1,108 306	776 2,539 33,652 7,519	729 2,159	10			1
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	10 - -	- - - 3 6	3,114 1,080 1,108 306	33,652 7,519				-	-
Upstate N.Y. N.Y. City N.J. Pa. E.N. CENTRAL Ohio Ind. III. Wich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	10 - -	- - - 3 6	1,080 1,108 306	7,519	26 607	15	39	-	3
N.J. Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	10 - -	6	306		36,697 7,401	230 103	194 67	4 2	-
Pa. E.N. CENTRAL Ohio Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	10 - -	6		9,677 5,724	10,834 7,079	54 48	50 42	-	-
Ohio Ind. Ill. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	10 - -		620	10,732	11,383	25	35	2	3
Ind. III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	-	6	2,849	54,350	65,165	186	235	3	2
III. Mich. Wis. W.N. CENTRAL Minn. Iowa Mo. Mo. Dak. S. Dak.	- 1 -	_	835	13,948 6,189	18,408 6,006	71 37	62 43	- 1	1
Wis. W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	1 -	-	672	16,731	20,674	57	86	-	-
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.		-	816 526	12,480 5,002	14,802 5,275	14 7	13 31	2	- 1
Minn. Iowa Mo. N. Dak. S. Dak.	-	3	1,774	14,423	14,568	57	64	1	1
Mo. N. Dak. S. Dak.	-	-	697	2,514	2,268	42	36	i	-
N. Dak. S. Dak.	N	N	279 421	1,117 7,566	1,146 7,551	1 10	- 16	-	-
	-	3	27	42	41	-	7	-	-
14001.	-	-	66 133	232 713	244 1,035	1	3	-	- 1
Kans.	-	-	151	2,239	2,283	3	2	-	-
S. ATLANTIC	1	-	2,528	72,503	80,266	325	312	4	1
Del. Md.	-	-	47 104	1,376 7,658	1,491 7,961	- 78	- 74	2	-
D.C.	-	-	41	2,387	2,510	-	-	-	-
Va. W. Va.	- 1	-	271 50	8,053 812	9,437 613	29 15	27 14	-	1
N.C.	-	-	-	13,823	14,762	30	44	-	-
S.C. Ga.	-	-	118 774	6,304 14,223	9,487 15,533	12 82	5 85	-	-
Fla.	-	-	1,123	17,867	18,472	79	63	2	-
E.S. CENTRAL	8	3	324	23,545	27,862	59	67	1	-
Ky. Tenn.	8	3	153	3,287 8,191	3,116 8,483	4 30	2 37	-	-
Ala.	-	-	171	7,118	9,447	16	26	1	-
Miss.	<del>-</del>	-	-	4,949	6,816	9	2	-	-
W.S. CENTRAL Ark.	1 -	-	208 143	41,424 3,861	45,623 3,965	57 2	50 1	2	2
La.	-	-	3	10,295	10,962	8	9	-	-
Okla. Tex.	1	-	62	4,088 23,180	4,167 26,529	42 5	38 2	2	2
MOUNTAIN	12	1	1,458	8,759	8,999	151	128	3	7
Mont.	-	-	78	95	88	-	-	-	-
ldaho Wyo.	-	-	115 29	81 55	69 72	2 1	2 1	-	-
Colo.	12	1	483	2,959	2,752	31	35	-	-
N. Mex. Ariz.	-	-	135 189	1,204 3,220	877 3,417	25 63	21 52	1	1 4
Utah	-	-	292	227	163	17	6	1	-
Nev.	-	-	137	918	1,561	12	11	·	2
PACIFIC Wash.	-	-	915 353	23,902 2,504	24,464 2,606	116 3	118 5	3 2	4 -
Oreg.	-	-	386	755	995	57	33	- 1	- 1
Calif. Alaska	-	-	96	19,526 516	19,953 370	22 1	52 6	-	4 -
Hawaii	-	-	80	601	540	33	22	-	-
Guam	-	-	-	-	44 521	- 1	1	-	-
P.R. V.I.	-	-	38	292 31	521 23	1	ı	-	-
Amer. Samoa C.N.M.I.	-	U	-	0.	23	-	-	-	-

N: Not notifiable. U: Unavailable. -: No reported cases.

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending November 9, 2002, and November 10, 2001 (45th Week)\*

(45th Week)*										
	Ha	aemophilus in	fluenzae, Inva	sive						
		Age <	5 Years			Н	epatitis (Viral,	Acute), By Ty	<b>у</b> ре	
		rotype B	Unknown		+	A		В	C; Non-A	<del></del>
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001
UNITED STATES	203	212	15	26	7,309	8,905	5,688	6,275	2,895	3,454
NEW ENGLAND	11	15	-	-	263	613	205	123	22	33
Maine N.H.	-	- 1	-	-	8 11	10 15	9 20	5 13	-	-
Vt.	-	-	-	-	1	14	4	5	13	7
Mass. R.I.	8	7	-	-	126 30	299 59	112 24	30 25	9	26
Conn.	3	7	-	-	87	216	36	45	-	-
MID. ATLANTIC	27	31	-	3	883	1,120	1,256	1,201	1,416	1,175
Upstate N.Y. N.Y. City	11 8	9 11	-	1 -	165 412	226 389	120 624	108 559	63	26
N.J.	5	4	-	-	117	259	316	261	1,322	1,086
Pa.	3	7	-	2	189	246	196	273	31	63
E.N. CENTRAL Ohio	30 8	38 12	1 1	2	975 297	1,066 209	549 94	838 88	92 8	150 8
Ind.	7	6	-	1	44	90	42	46	-	1
III. Mich.	11 3	14	-	1	252 215	397 298	126 287	132 534	13 71	11 130
Wis.	1	6	-	-	167	72	-	38	-	-
W.N. CENTRAL	6	5	3	6	279	347	198	188	717	1,010
Minn. Iowa	5	3	1 -	2	38 74	39 32	27 17	21 21	1	9 -
Mo. N. Dak.	-	- 1	2	4	78 1	77	106	106	698	988
S. Dak.	-	-	-	-	3	3 3	4 2	1 1	1	-
Nebr. Kans.	1	1	-	-	17 68	31 162	22 20	26 12	13 4	6 7
S. ATLANTIC	44	42	2	6	2,124	2,156	1,446	1,296	167	94
Del.	-	-	-	-	12	16	7	25	5	10
Md. D.C.	4	7	-	1 -	273 70	226 47	107 22	127 11	6	8
Va.	4	5	-	-	129	115	176	157	16	-
W. Va. N.C.	1 3	1 2	1 -	1 4	17 195	18 202	18 207	20 173	3 25	9 19
S.C.	2	1	-	-	56	66	112	28	4	6
Ga. Fla.	17 13	17 9	1	-	402 970	845 621	338 459	381 374	29 79	42
E.S. CENTRAL	13	12	1	3	240	362	339	414	180	180
Ky. Tenn.	1 7	- 6	-	1 1	41 109	122 139	48 120	49 207	3 24	9 61
Ala.	3	5	1	1	35	70	95	78	10	4
Miss.	2	1	-	-	55	31	76	80	143	106
W.S. CENTRAL Ark.	14 1	9 1	-	-	549 42	766 64	468 75	748 86	143 7	643 10
La.	2	2	-	-	62	85	84	110	51	142
Okla. Tex.	9 2	6	-	-	47 398	106 511	44 265	85 467	5 80	4 487
MOUNTAIN	35	21	7	1	510	632	536	405	60	50
Mont.	-	-	-	-	13	11	9	3	1	1
ldaho Wyo.	1 -	-	-	-	26 3	52 7	6 17	11 3	1 5	2 7
Colo.	3	2	-	-	72	79	69	87	18	8
N. Mex. Ariz.	6 16	9 8	1 5	1 -	27 263	39 322	129 199	116 120	1 4	11 9
Utah	5 4	2	- 1	-	59	62	53	22	4	3
Nev. PACIFIC	23	39	1	5	47 1,486	60 1,843	54 691	43 1,062	26 98	9 119
Wash.	1	3	-	2	140	133	58	128	23	20
Oreg. Calif.	5 13	6 28	- 1	- 1	61 1,273	92 1,588	113 508	147 761	16 59	14 85
Alaska	1	1	-	-	10	14	4	9	-	-
Hawaii	3	1	-	2	2	16	8	17	-	-
Guam P.R.	-	1	-	-	96	1 201	- 84	237	-	- 1
V.I.	-	-	-	-	-	-	-	-	-	-
Amer. Samoa C.N.M.I.	U -	U U	U -	U U	U -	U U	U 37	U U	U -	U

N: Not notifiable. U: Unavailable. -: No reported cases.

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending November 9, 2002, and November 10, 2001 (45th Week)\*

(45th Week)*	· ·		1		1				Mea	slos
	Legior	ellosis	Liste	1	Lyme	Disease	Ma	laria	To	
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001
UNITED STATES	979	964	510	527	14,597	13,592	1,094	1,301	27 <sup>†</sup>	113§
NEW ENGLAND	89	65	54	52	4,266	3,949	56	86	-	5
Maine N.H.	2 5	7 10	5 4	2 4	111 230	96	5 7	4 2	-	-
Vt.	36	5	3	3	32	16	4	1	-	1
Mass. R.I.	30 2	19 10	28 1	27 1	1,150 314	1,098 449	21 5	47 9	-	3 -
Conn.	14	14	13	15	2,429	2,290	14	23	-	1
MID. ATLANTIC Upstate N.Y.	263 91	228 61	145 52	95 25	8,520 4,584	7,441 3,116	264 43	391 56	7 1	19 4
N.Y. City	46	43	30	23	142	61	163	233	6	6
N.J. Pa.	22 104	21 103	30 33	17 30	1,448 2,346	1,970 2,294	28 30	59 43	- -	1 8
E.N. CENTRAL	231	274	68	81	84	700	124	157	3	10
Ohio	105	116	24	13	66	38 22	22	23	1	3
Ind. III.	18 -	20 24	9 11	8 23	18	31	12 30	16 64	2	4 3
Mich. Wis.	74 34	72 42	18 6	24 13	- U	17 592	46 14	36 18	-	-
W.N. CENTRAL	50	45	17	16	333	364	56	33	3	5
Minn.	11	9	3	-	241	292	17	6	1	3
Iowa Mo.	11 14	8 19	2 8	2 9	36 39	34 32	4 15	6 13	2	2
N. Dak.	-	1	1	-	1	-	1	-	-	-
S. Dak. Nebr.	4 10	3 4	1 1	1	2 6	4	1 5	2	-	-
Kans.	-	1	1	4	8	2	13	6	-	-
S. ATLANTIC Del.	186 7	167 12	72	71 2	1,173 155	887 152	320 4	265 2	2	5
Md.	41	32	17	13	623	539	101	108	-	3
D.C. Va.	6 24	8 20	7	12	20 144	14 115	19 31	13 45	-	1
W. Va.	N	N 9	-	5 5	17	11	3	1	-	-
N.C. S.C.	11 8	13	6 8	5	122 20	38 5	21 7	17 6	-	-
Ga. Fla.	19 70	11 62	11 23	14 15	2 70	- 13	73 61	41 32	2	1
E.S. CENTRAL	40	54	17	21	46	62	20	35	5	2
Ky.	18	12	3	7	21	22 25	8	14	-	2
Tenn. Ala.	14 8	26 12	10 4	8 6	22 3	8	3 4	11 6	5	-
Miss.	-	4	-	-	-	7	5	4	-	-
W.S. CENTRAL Ark.	10	23	18	31 1	16 3	81	15 2	83 3	2	1
La.	1	6	-	-	3	8	4	6	-	-
Okla. Tex.	3 6	3 14	9 9	2 28	10	73	8 1	3 71	2	1
MOUNTAIN	43	49	27	34	20	11	42	53	1	2
Mont. Idaho	3	3	2	- 1	- 4	5	2	3 3	-	- 1
Wyo.	i	2	-	į	1	1	-	-	-	-
Colo. N. Mex.	7 2	14 3	6 3	9 7	3 1	-	22 3	22 3	- -	-
Ariz.	10	16	12	7	3	1	7	10	-	1
Utah Nev.	14 5	7 4	3 1	2 7	7 1	1 3	5 3	4 8	1	-
PACIFIC	67	59	92	126	139	97	197	198	4	64
Wash. Oreg.	7 N	9 N	8 9	10 12	10 15	7 11	21 9	9 15	-	15 3
Calif.	59	44	67	98	111	77	158	162	3	39
Alaska Hawaii	1	1 5	8	6	3 N	2 N	2 7	1 11	1	7
Guam	-	-	-	-	-	-	-	1	-	-
P.R. V.I.	-	2	1 -	-	N -	N -	-	5	-	1
Amer. Samoa	U	U	Ü	U	U	U	U	Ü	Ū	Ü
C.N.M.I.	-	U	-	U	-	U	-	U	-	U

N: Not notifiable.

N: Not notifiable. U: Unavailable. -: No reported cases.

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

† Of 27 cases reported, 14 were indigenous and 13 were imported from another country.

§ Of 113 cases reported, 59 were indigenous and 54 were imported from another country.

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending November 9, 2002, and November 10, 2001 (45th Week)\*

	Meningo Disea		Mun	nps	Pert	ussis	Rabies	, Animal
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001
INITED STATES	1,425	2,008	230	210	6,531	4,770	5,423	6,301
NEW ENGLAND	79 7	93	7	1	534	460	828	655
laine I.H.	7 11	4 12	4	-	17 17	22 17	54 46	63 19
t.	4	5	-	-	122	33	86	58
lass. .l.	38 5	50 4	2	1	340 13	365 5	269 69	243 64
onn.	14	18	1	-	25	18	304	208
IID. ATLANTIC	133	228	25	25	403	316	999	1,167
pstate N.Y. I.Y. City	40 21	62 41	7 2	3 12	292 13	128 51	622 10	711 34
I.J.	25	39	-	3	3	18	157	173
a.	47	86	16 32	7	95	119	210	249 147
i.N. CENTRAL Dhio	188 72	310 79	32 13	25 1	785 378	751 270	146 38	42
nd. I.	29 36	35 78	2 8	3 16	119 141	78 89	31 31	15 24
ı. 1ich.	39	78 70	8	3	47	133	46	46
Vis.	12	48	1	2	100	181	-	20
V.N. CENTRAL ⁄linn.	132 32	141 20	16 4	8 3	670 340	320 146	402 36	338 43
owa	21	29	1	-	130	46	72	76
1o. I. Dak.	43	50 6	5 1	1	127	91 4	49 26	40 35
. Dak.	2	5	-	-	6	4	65	53
lebr. ans.	26 8	17 14	- 5	1 3	8 59	5 24	154	4 87
. ATLANTIC	257	312	25	36	375	223	2,248	2,193
el.	7	5	-	-	3	-	24	30
1d. ).C.	8	38	5	5	57 2	37 1	321	455
a.	37	37	4	8	132	40	454	423
V. Va. I.C.	4 30	12 61	2	- 5	31 40	3 68	161 644	131 510
S.C.	28	31	3	5	41	31	133	102
ia. Ia.	33 110	45 83	4 7	8 5	21 48	20 23	347 164	365 177
S. CENTRAL	85	122	13	9	231	164	153	197
ý. enn.	13 37	21 55	3 2	3 1	89 101	68 57	26 97	26 106
vla.	21	30	3	-	32	35	26	61
liss.	14	16	5	5	9	4	4	4
V.S. CENTRAL ırk.	174 23	297 21	17 -	12 -	1,476 459	559 130	109 3	1,006
a.	30	74	1	2	7	8	-	8
Okla. ex.	19 102	28 174	16	10	66 944	27 394	105 1	57 941
IOUNTAIN	78	86	18	14	913	1,204	278	252
font.	2	4	-	1	5	34	18	37
laho √yo.	4 -	7 5	2	1	65 11	170 1	37 18	28 28
olo. I. Mex.	21	33 10	2 1	3 2	359 163	282 129	59 7	- 15
riz.	4 23	13	i	1	167	496	115	128
tah	4 20	8 6	7 5	1 4	96 47	74 18	13 11	15 1
lev. ACIFIC	299	419	5 77	80	1,144	773	260	346
Vash.	58	59	-	2	383	140	-	-
Oreg. Calif.	42 188	56 289	N 63	N 39	173 567	49 541	13 223	4 304
laska	4	2	-	1	4	9	24	38
lawaii	7	13	14	38	17	34	-	-
luam R.	- 5	- 5	-	- 1	3		49	- 85
II.	-	-		-	-	. <del>.</del>	-	-
mer. Samoa C.N.M.I.	U	U U	U	U U	U 1	U U	U	U

N: Not notifiable. U: Unavailable. -: No reported cases.

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending November 9, 2002, and November 10, 2001 (45th Week)\*

		Mountain	Rub	velle		enital pella	Salmor	allasia
Reporting Area	Cum. 2002	d Fever Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001
UNITED STATES	916	536	13	21	2	-	35,747	34,871
NEW ENGLAND	7	3	-	-	-	-	1,952	2,138
Maine	-	-	-	-	-	-	134	159
N.H. Vt.	-	1	-	-	-	-	125 70	155 74
Mass.	4	2	-	-	-	-	1,075	1,230
R.I.	3	-	-	-	-	-	149	120
Conn.	-		-	-	-	-	399	400
MID. ATLANTIC Upstate N.Y.	39 7	31 2	1 1	8 1	-	-	4,345 1,370	4,581 1,062
N.Y. City	8	2	-	6	-	-	1,161	1,156
N.J.	10	9	-	1	-	-	621	1,064
Pa.	14	18	-	-	-	-	1,193	1,299
E.N. CENTRAL Ohio	17 12	16 2	1	2	-	-	4,685 1,268	4,469 1,219
Ind.	2	1	-	-	-	-	425	472
III.	-	12	-	2	-	-	1,445	1,251
Mich. Wis.	3	1	1	-	-	-	791 756	775 752
			-	-	-	-		
W.N. CENTRAL Minn.	97	67	-	3	-	-	2,334 509	2,041 553
lowa	3	2	-	1	-	-	461	317
Mo. N. Dak.	89	61	-	1	-	-	768	555
S. Dak.	1	1 2	-	-	-	-	42 102	58 141
Nebr.	4	1	-	-	-	-	150	142
Kans.	-	-	-	1	-	-	302	275
S. ATLANTIC	482	264	5	5	-	-	9,918	8,145
Del. Md.	4 56	10 38	-	- 1	-	-	81 840	89 706
D.C.	2	-	-	-	-	-	69	700
Va.	38	25	-	-	-	-	1,086	1,193
W. Va. N.C.	2 270	149	-	-	-	-	128 1,334	118 1,185
S.C.	68	29	-	2	-	-	720	785
Ga.	27	9	-	-	-	-	1,827	1,514
Fla.	15	4	5	2	-	-	3,833	2,483
E.S. CENTRAL	97	104	-	-	1	-	2,811	2,436
Ky. Tenn.	5 73	2 72	-	-	1	-	339 715	336 576
Ala.	16	15	-	-	-	-	781	680
Miss.	3	15	-	-	-	-	976	844
W.S. CENTRAL	158	39	2	1	-	-	3,068	4,492
Ark. La.	97	8 2	-	-	-	-	914 659	828 782
Okla.	61	29	-	-	-	-	446	430
Tex.	-	-	2	1	-	-	1,049	2,452
MOUNTAIN	13	11	1	-	-	-	1,961	1,926
Mont.	1	1	-	-	-	-	80	68
Idaho Wyo.	4	1 2	-	-	-	-	131 91	126 58
Colo.	2	2	-	-	-	-	492	534
N. Mex.	1	1	-	-	-	-	283	254
Ariz. Utah	-	3	1	-	-	-	524 183	534 197
Nev.	5	1	-	-	-	-	177	155
PACIFIC	6	1	3	2	1	-	4,673	4,643
Wash.	-	<del>-</del>	-	-	-	-	457	462
Oreg. Calif.	2 4	1	3	- 1	-	-	322 3,577	248 3,579
Alaska	-	-	-	-	-	-	3,577 72	39
Hawaii	-	-	-	1	1	-	245	315
Guam	-	-	-	-	-	-	-	22
P.R.	-	-	-	3	-	-	201	830
V.I. Amer. Samoa	U	U	U	- U	- U	- U	U	Ū
Amer. Samoa								

N: Not notifiable. U: Unavailable. -: No reported cases.

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending November 9, 2002, and November 10, 2001 (45th Week)\*

(45th Week)*	Shig	ellosis	Streptococo Invasive,			s pneumoniae, ant, Invasive	Streptococcus pneumoniae, Invasive (<5 Years)		
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	
UNITED STATES	15,952	16,856	3,549	3,204	2,020	2,250	226	371	
NEW ENGLAND	287	276	167	197	17	112	3	40	
Maine N.H.	10 11	6 6	20 35	10 N	-	-	- N	N	
Vt. Mass.	1 171	7 195	9 88	14 59	5 N	7 N	2 N	1 N	
R.I.	16	17	15	12	12	4	1	3	
Conn.	78	45	-	102	-	101	-	36	
MID. ATLANTIC Upstate N.Y.	1,169 270	1,356 435	568 261	593 236	97 81	143 136	59 58	96 96	
N.Y. City	355	374	133	157	U	U	U	U	
N.J. Pa.	332 212	252 295	118 56	124 76	N 16	N 7	N 1	N -	
E.N. CENTRAL	1,577	3,918	646	714	199	161	99	115	
Ohio Ind.	570 89	2,613 198	190 45	181 56	53 141	1 160	19 55	- 54	
III.	611	545	145	231	2	-	-	61	
Mich. Wis.	162 145	279 283	266	195 51	3 N	- N	N 25	N -	
W.N. CENTRAL	909	1,728	214	337	414	137	49	54	
Minn. Iowa	201 115	387 340	108	156	292 N	63 N	49 N	45 N	
Mo.	166	286	42	69	5	9	-	-	
N. Dak. S. Dak.	16 150	21 543	- 13	17 11	1 1	6 4	-	9	
Nebr. Kans.	179 82	84 67	18 33	36 48	29 86	19 36	N N	N N	
S. ATLANTIC	5,892	2,396	716	529	1,067	1,187	7	5	
Del.	275	14	2	4	3	6	N	N	
Md. D.C.	1,024 53	137 52	123 7	N 21	N 48	N 5	N 1	N 3	
Va. W. Va.	870 9	336 8	68 19	70 19	N 39	N 37	N 6	N 2	
N.C.	396	312	112	134	N	N	U	U	
S.C. Ga.	106 1,350	237 449	34 153	10 166	169 268	243 370	N N	N N	
Fla.	1,809	851	198	105	540	526	N	N	
E.S. CENTRAL Ky.	1,261 157	1,516 717	103 18	107 35	119 17	216 24	- N	- N	
Tenn.	93	91	85	72	102	191	N	N	
Ala. Miss.	699 312	189 519	-	-	<del>-</del> -	1 -	N -	N -	
W.S. CENTRAL	1,523	2,600	100	292	67	253	5	61	
Ark. La.	164 372	532 218	6	- 1	6 61	15 238	2	- 61	
Okla.	518	76	41	39	N	N	3	-	
Tex. MOUNTAIN	469	1,774	53	252	N 10	N oz	-	-	
Mont.	806 3	864 8	514 -	361 -	40	37 -	4 -	-	
Idaho Wyo.	15 9	39 7	9 7	7 11	N 9	N 7	N	N	
Colo.	163	223	132	139	-	-	-	-	
N. Mex. Ariz.	192 346	112 354	95 241	75 126	30	28	N	N	
Utah Nev.	33 45	52 69	30	3	- 1	2	4	-	
PACIFIC	2,528	2,202	- 521	74	-	4	-	-	
Wash.	145	186	65	-	-	-	N	N	
Oreg. Calif.	103 2,213	98 1,857	N 364	N -	N N	N N	N N	N N	
Alaska Hawaii	6 61	6 55	92	- 74	-	4	N	N	
Guam	-	45	<i>3</i> ∠ -	1	- -	-	-	-	
P.R.	8	16	N	Ń	-	-	N	N	
V.I. Amer. Samoa	U	Ū	Ū	Ū	-	-	U	U	
C.N.M.I.	17	Ü	-	Ü	-	-	-	Ü	

N: Not notifiable. U: Unavailable. -: No reported cases.

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending November 9, 2002, and November 10, 2001 (45th Week)\*

(45th Week)*		Cum	hilis		1		Turn	hoid	
	Primary & S			genital	Tubero	culosis	Typhoid Fever		
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	
UNITED STATES	5,432	5,222	299	431	9,977	11,972	228	319	
NEW ENGLAND	121	54	-	5	332	396	14	16	
Maine N.H.	2 7	1 1	-	-	10 13	15 16	-	1 2	
Vt. Mass.	1 81	3 30	-	- 3	- 190	4 202	8	10	
R.I.	6	9	-	- 2	34	55	-	3	
Conn. MID. ATLANTIC	24 557	10 450	- 55	69	85 1,815	104 1,976	6 47	104	
Upstate N.Y.	29	17	8	5	250	321	9	15	
N.Y. City N.J.	344 127	247 106	21 25	32 32	933 418	978 431	23 11	42 38	
Pa.	57	80	1	-	214	246	4	9	
E.N. CENTRAL Ohio	937 141	913 69	51 4	61 2	997 126	1,224 244	18 6	32 4	
Ind.	58	139	1	9	105	85	2	2	
III. Mich.	286 428	334 348	29 17	40 6	508 217	570 256	1 4	17 5	
Wis.	24	23	-	4	41	69	5	4	
W.N. CENTRAL Minn.	94 48	89 31	<del>-</del> -	9 2	465 202	465 199	8 3	15 6	
Iowa Mo.	2 24	4 23	-	- 5	24 115	34 115	- 1	9	
N. Dak.	-	-	-	-	2	3	-	-	
S. Dak. Nebr.	3	- 8	-	-	10 23	12 32	4	-	
Kans.	17	23	-	2	89	70	-	-	
S. ATLANTIC Del.	1,459 10	1,770 13	67	102	2,021 13	2,224 15	44	41 1	
Md.	172	235	14	4	249	200	7	10	
D.C. Va.	55 59	34 92	1 1	2 5	163	51 224	7	11	
W. Va. N.C.	2 255	4 403	- 18	- 12	28 306	26 291	2	2	
S.C.	113	214	8	21	146	150	-	-	
Ga. Fla.	306 487	342 433	10 15	22 36	347 769	421 846	9 19	9 8	
E.S. CENTRAL	417	580	18	30	631	730	4	1	
Ky. Tenn.	83 153	43 286	3 8	1 17	114 251	115 265	4	1	
Ala. Miss.	140 41	118 133	4 3	5 7	177 89	235 115	-	-	
W.S. CENTRAL	742	648	64	72	1,345	1,822	5	18	
Ark. La.	31 133	33 155	2	7	109	134 114	-	-	
Okla.	61	55	3	5	122	131	2		
Tex.	517	405	59	60	1,114	1,443	3	18	
MOUNTAIN Mont.	259	193	15 -	29 -	299 6	477 6	11 -	8 1	
ldaho Wyo.	8 -	1 1	-	-	9 3	7 3	-	-	
Colo.	44 30	20	1	1 2	48	115	5 2	1	
N. Mex. Ariz.	162	15 139	14	26	22 171	47 193	-	1	
Utah Nev.	8 7	10 7	- -	- -	26 14	33 73	2 2	1 4	
PACIFIC	846	525	29	54	2,072	2,658	77	84	
Wash. Oreg.	53 20	42 13	1 1	- -	198 97	208 95	4 2	5 7	
Calif.	765	458	26	54	1,608	2,186	66	68	
Alaska Hawaii	8	12	1	-	43 126	43 126	5	1 3	
Guam	-	9	-	1	- 75	54	-	3	
P.R. V.I.	227 1	242	15 -	13 -	75 -	95 -	-	-	
Amer. Samoa C.N.M.I.	U 15	U U	U -	U U	U 32	U U	U -	U U	

N: Not notifiable. U: Unavailable. -: No reported cases.

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE III. Deaths in 122 U.S. cities.\* week ending November 9, 2002 (45th Week)

TABLE III. Deaths in 122 U.S. cities,* week ending November 9, 2002 (45th Week)  All Causes, By Age (Years)  All Causes, By Age (Years)															
-		All C	Jauses, E	By Age (Y	ears)					All	Causes, I	By Age (1	rears)	1	
Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	P&I <sup>†</sup> Total	Reporting Area	All Ages	<u>≥</u> 65	45-64	25-44	1-24	<1	P&I <sup>†</sup> Total
NEW ENGLAND	320	239	54	21	3	3	35	S. ATLANTIC	1,153	745	247	113	28	20	77
Boston, Mass.	U 45	U 33	U 8	U 3	U	U 1	U 3	Atlanta, Ga. Baltimore, Md.	U 185	U 116	U 46	U 17	U 3	U 3	U 7
Bridgeport, Conn. Cambridge, Mass.	14	12	2	-	-	-	2	Charlotte, N.C.	89	52	24	8	3	2	9
Fall River, Mass.	26	20	5	1	-	-	5	Jacksonville, Fla.	143	95	25	14	7	2	10
Hartford, Conn.	U	U	U	U	U	U	U	Miami, Fla.	122	83	25	10	1	3	10
Lowell, Mass.	22	16	3	3	-	-	4	Norfolk, Va.	51	37	13	-	-	1	-
Lynn, Mass.	18 30	15 26	1 3	2 1	-	-	2 2	Richmond, Va. Savannah, Ga.	49 48	29 34	13 6	5 6	1 1	1 1	2 5
New Bedford, Mass. New Haven, Conn.	40	26 26	5 5	4	3	2	5	St. Petersburg, Fla.	48 49	30	6	8	3	2	5 4
Providence, R.I.	Ü	Ü	Ŭ	Ū	Ŭ	Ū	Ŭ	Tampa, Fla.	195	127	42	19	4	3	21
Somerville, Mass.	3	1	2	-	-	-	-	Washington, D.C.	204	125	46	26	5	2	5
Springfield, Mass.	36	24	10	2	-	-	2	Wilmington, Del.	18	17	1	-	-	-	4
Waterbury, Conn.	26	19	6	1	-	-	2	E.S. CENTRAL	801	514	182	68	21	16	61
Worcester, Mass.	60	47	9	4		-	8	Birmingham, Ala.	193	120	41	22	7	3	21
MID. ATLANTIC	2,113	1,455	447	137	28	32	98	Chattanooga, Tenn.	64	43	14	3	2	2	4
Albany, N.Y. Allentown, Pa.	48 27	33 23	11 3	1 1	2	1	2 1	Knoxville, Tenn. Lexington, Ky.	83 62	52 43	23 12	7 2	1 4	1	4 6
Buffalo, N.Y.	103	71	21	7	-	4	6	Memphis, Tenn.	142	87	37	17	-	1	7
Camden, N.J.	33	23	7	3	-		1	Mobile, Ala.	75	56	14	2	1	2	5
Elizabeth, N.J.	13	6	4	2	1	-	-	Montgomery, Ala.	26	19	4	2	1	-	3
Erie, Pa.	68	55	10	2	-	1	4	Nashville, Tenn.	156	94	37	13	5	7	11
Jersey City, N.J.	58	38	10	8	- 15	2	-	W.S. CENTRAL	1,313	819	284	113	57	40	75
New York City, N.Y. Newark, N.J.	1,188 51	823 18	264 21	72 8	15 1	14 1	59 6	Austin, Tex.	76	40	25	4	2	5	7
Paterson, N.J.	23	17	-	3	2	1	1	Baton Rouge, La.	26	21	3	2	-	-	-
Philadelphia, Pa.	227	161	42	18	5	1	7	Corpus Christi, Tex.	68 207	38	23	5	1	1	4
Pittsburgh, Pa.§	36	17	13	5	-	1	-	Dallas, Tex. El Paso, Tex.	63	130 46	45 9	20 5	4 1	8 2	14 2
Reading, Pa.	27	19	.7		-	1	1	Ft. Worth, Tex.	105	65	24	5	3	8	3
Rochester, N.Y.	118 22	79 17	17 2	4 3	2	4	10	Houston, Tex.	322	175	64	38	35	10	16
Schenectady, N.Y. Scranton, Pa.	27	17	7	-	-	1	-	Little Rock, Ark.	55	34	18	1	2	-	-
Syracuse, N.Y.	Ū	Ü	Ú	U	U	Ü	U	New Orleans, La.	38	19	10	5	4	-	-
Trenton, N.J.	20	15	5	-	-	-	-	San Antonio, Tex. Shreveport, La.	200 53	132 39	41 9	20 2	3 2	4 1	19 5
Utica, N.Y.	24	21	3	-	-	-	-	Tulsa, Okla.	100	80	13	6	-	i	5
Yonkers, N.Y.	U	U	U	U	U	U	U	MOUNTAIN	829	550	185	59	25	10	66
E.N. CENTRAL	1,668	1,172	332	92	30	38	114	Albuquerque, N.M.	104	67	22	12	1	2	8
Akron, Ohio Canton, Ohio	61 43	43 29	9 13	4 1	1 -	-	7 2	Boise, Idaho	33	24	6	1	2	-	3
Chicago, III.	U	U	Ü	ΰ	Ū	Ū	Ū	Colo. Springs, Colo.	64	47	9	4	3	1	3
Cincinnati, Ohio	84	61	15	6	-	2	13	Denver, Colo.	103	69	21	8	4 9	1	9
Cleveland, Ohio	118	82	23	8	2	3	5	Las Vegas, Nev. Ogden, Utah	232 26	139 20	65 3	18 3	9	-	21 7
Columbus, Ohio	217	152	41	14	5	5	12	Phoenix, Ariz.	U	U	Ü	Ü	U	U	Ú
Dayton, Ohio Detroit. Mich.	132 185	105 104	16 56	6	4 5	1 5	7	Pueblo, Colo.	27	22	5	-	-	-	3
Evansville, Ind.	44	33	9	15 1	5	ວ 1	16 5	Salt Lake City, Utah	98	56	27	6	5	4	3
Fort Wayne, Ind.	66	53	6	5	2	-	5	Tucson, Ariz.	142	106	27	7	1	1	9
Gary, Ind.	17	9	5	1	-	2	1	PACIFIC	1,674	1,164	336	112	28	34	105
Grand Rapids, Mich.	61	45	8	4	-	4	5	Berkeley, Calif.	16	11	2	3	-	-	3
Indianapolis, Ind.	179 54	131 34	37 13	6 4	1	4 3	15	Fresno, Calif. Glendale, Calif.	157	112 16	30 3	10	2	3	17
Lansing, Mich. Milwaukee, Wis.	123	97	15	5	2	4	3 7	Honolulu, Hawaii	19 60	47	10	3	-		3
Peoria, III.	55	35	13	6	1		3	Long Beach, Calif.	62	40	15	5	1	1	5
Rockford, III.	65	36	23	2	2	2	3	Los Angeles, Calif.	354	230	79	29	10	6	14
South Bend, Ind.	39	26	6	1	5	1	-	Pasadena, Calif.	21	18	1	. 1	-	1	6
Toledo, Ohio	75 50	57	16	2	-	-	3	Portland, Oreg.	159	111	30	10	2	6	11
Youngstown, Ohio	50	40	8	1	-	1	2	Sacramento, Calif. San Diego, Calif.	168 144	121 101	31 21	10 15	2 4	4 3	8 7
W.N. CENTRAL	507	365	86	31	17	8	42	San Francisco, Calif.	U	Ü	Ü	Ü	Ü	Ü	ύ
Des Moines, Iowa	U 27	U	U	U	U	U	U	San Jose, Calif.	187	135	36	10	2	4	18
Duluth, Minn. Kansas City, Kans.	27 31	22 17	2 7	2 5	1 -	2	4 2	Santa Cruz, Calif.	27	22	5	-	-	-	3
Kansas City, Mo.	117	90	16	8	3	-	9	Seattle, Wash.	131	85	34	6	3	3	5
Lincoln, Nebr.	32	27	3	1	-	1	4	Spokane, Wash.	59	40 75	11	4	1	3	1
Minneapolis, Minn.	70	42	13	7	5	3	3	Tacoma, Wash.	110	75	28	6	-	-	4
Omaha, Nebr.	92	71	14	6	-	1	11	TOTAL	10,378 <sup>¶</sup>	7,023	2,153	746	237	201	673
St. Louis, Mo. St. Paul, Minn.	U 56	U 40	U 13	U 1	U 2	U	U 6								
Wichita, Kans.	56 82	40 56	18	1	6	1	3								
	02		10					I							

U: Unavailable. -: No reported cases.

Or. Orlavaliable.
 1.No reported classes.
 Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of ≥100,000. 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 this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks.
 Total includes unknown ages.

All MMWR references are available on the Internet at http://www.cdc.gov/mmwr. Use the search function to find specific articles.

Use of trade names and commercial sources is for identification only and does not imply endorsement by the U.S. Department of Health and Human Services.

References to non-CDC sites on the Internet are provided as a service to MMWR readers and do not constitute or imply endorsement of these organizations or their programs by CDC or the U.S. Department of Health and Human Services. CDC is not responsible for the content of these sites. URL addresses listed in MMWR were current as of the date of publication.

The Morbidity and Mortality Weekly Report (MMWR) Series is prepared by the Centers for Disease Control and Prevention (CDC) and is available free of charge in electronic format and on a paid subscription basis for paper copy. To receive an electronic copy each week, send an e-mail message to listserv@listserv.cdc.gov. The body content should read SUBscribe mmwr-toc. Electronic copy also is available from CDC's World-Wide Web server at http://www.cdc.gov/mmwr or from CDC's file transfer protocol server at ftp://ftp.cdc.gov/pub/publications/mmwr. To subscribe for paper copy, contact Superintendent of Documents, U.S. Government Printing Office, Washington, DC 20402; telephone 202-512-1800.

Data in the weekly *MMWR* are provisional, based on weekly reports to CDC by state health departments. The reporting week concludes at close of business on Friday; compiled data on a national basis are officially released to the public on the following Friday. Address inquiries about the *MMWR* Series, including material to be considered for publication, to Editor, *MMWR* Series, Mailstop C-08, CDC, 1600 Clifton Rd., N.E., Atlanta, GA 30333; telephone 888-232-3228.

All material in the MMWR Series is in the public domain and may be used and reprinted without permission; citation as to source, however, is appreciated.

☆U.S. Government Printing Office: 2003-533-155/69074 Region IV