

Weekly

September 16, 2005 / Vol. 54 / No. 36

Direct and Indirect Effects of Routine Vaccination of Children with 7-Valent Pneumococcal Conjugate Vaccine on Incidence of Invasive Pneumococcal Disease — United States, 1998–2003

Streptococcus pneumoniae (pneumococcus) is a leading cause of pneumonia and meningitis in the United States and disproportionately affects young children and the elderly. In 2000, a 7-valent pneumococcal conjugate vaccine (PCV7) was licensed in the United States for routine use in children aged <5 years (1). Surveillance data from 2001 and 2002 indicated substantial declines in invasive pneumococcal disease (IPD) in children and adults compared with prevaccine years (2,3). This report updates assessment of the impact of PCV7 on IPD through 2003 by using population-based data from the Active Bacterial Core surveillance (ABCs) of the Emerging Infections Program Network, a cooperative surveillance program conducted by several state health departments and CDC.* The results of this analysis indicated that 1) routine vaccination of young children with PCV7 continued to result in statistically significant declines in incidence of IPD through 2003 in the age group targeted for vaccination and among older children and adults, 2) the vaccine prevented more than twice as many IPD cases in 2003 through indirect effects on pneumococcal transmission (i.e., herd immunity) than through its direct effect of protecting vaccinated children, and 3) increases in disease caused by pneumococcal serotypes not included in the vaccine (i.e., replacement disease) occurred in certain populations but were small compared with overall declines in vaccine-serotype disease. Ongoing surveillance is needed to assess whether reductions in vaccine-serotype IPD are sustained and whether replacement disease will erode the substantial benefits of routine vaccination.

ABCs conducted active surveillance for IPD cases through regular contact with all clinical microbiology laboratories in defined surveillance areas; periodic audits of laboratory records ensured complete case finding. Pneumococcal isolates were sent to reference laboratories for serotyping by the quellung reaction and were categorized as vaccine-type (VT) (serotypes included in PCV7) or nonvaccine-type (NT) (all other serotypes). A case of IPD was defined as isolation of pneumococcus from a normally sterile body site (e.g., blood or cerebrospinal fluid) in an ABCs area resident. Participating areas during 1998-2003 included in this analysis were the state of Connecticut and selected counties in California, Georgia, Maryland, Minnesota, New York, and Oregon, representing a total surveillance population of approximately 16 million persons in 2000. Annual incidence rates were calculated for 1998–1999 by using U.S. Census Bureau population estimates for those years; incidence rates for 2001-2003 were based on National Center for Health Statistics (NCHS) bridged-race postcensal population estimates for those years (4). For national projections of annual numbers of IPD cases, age- and race-specific rates of disease were applied from the aggregate ABCs surveillance area to the age and racial distribution of the U.S. population.

The impact of PCV7 introduction on IPD was assessed in three ways. First, to assess the change in incidence of IPD after PCV7 introduction, IPD rates for 2001–2003 were

INSIDE

- 897 Improper Disposal of Hazardous Substances and Resulting Injuries — Selected States, January 2001– March 2005
- 899 Update: Influenza Activity United States and Worldwide, May 22–September 3, 2005, and 2005–06 Season Vaccination Recommendations
- 902 Update: West Nile Virus Activity United States, 2005
- 904 QuickStats

^{*}Available at http://www.cdc.gov/ncidod/dbmd/abcs.

The *MMWR* series of publications is published by the Coordinating Center for Health Information and Service, 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 2005;54:[inclusive page numbers].

Centers for Disease Control and Prevention

Julie L. Gerberding, MD, MPH Director

Dixie E. Snider, MD, MPH Chief Science Officer

Tanja Popovic, MD, PhD Associate Director for Science

Coordinating Center for Health Information and Service

Blake Caldwell, MD, MPH, and Edward J. Sondik, PhD (Acting) Directors

National Center for Health Marketing*

Jay M. Bernhardt, PhD, MPH Director

Division of Scientific Communications*

Maria S. Parker (*Acting*) *Director*

Mary Lou Lindegren, MD Editor, MMWR Series

Suzanne M. Hewitt, MPA Managing Editor, MMWR Series

Douglas W. Weatherwax (Acting) Lead Technical Writer-Editor

> Stephanie M. Neitzel Jude C. Rutledge *Writers-Editors*

Lynda G. Cupell Malbea A. LaPete *Visual Information Specialists*

Quang M. Doan, MBA Erica R. Shaver Information Technology Specialists

Notifiable Disease Morbidity and 122 Cities Mortality Data Patsy A. Hall Rosaline Dhara

> Deborah A. Adams Felicia J. Connor

Rosaline Dhara Tambra McGee Pearl C. Sharp compared with the average rate for 1998–1999 (baseline). Second, the projected number of VT IPD cases directly prevented by PCV7 in 2003 was calculated as the product of 1) the nationally projected number of VT IPD cases at baseline among children aged <5 years, 2) the 3-dose coverage of PCV7 in 2003 among all U.S. children aged 19–35 months identified from National Immunization Survey (NIS) data (68.1%) (5), and 3) vaccine efficacy against VT IPD from a large clinical trial (93.9%) (6). Third, the projected number of VT IPD cases indirectly prevented by PCV7 in 2003 was estimated across all ages aggregately by calculating the difference between the average annual projected number of VT cases in 1998–1999 and the projected number of VT cases directly prevented by the vaccine.

From 1998–1999 to 2003, the incidence of VT IPD among children aged <5 years decreased from 80.0 cases per 100,000 population to 4.6, a decline of 94% (95% confidence interval [CI] = 92%-96% (Figure 1). The total incidence of IPD (VT and NT) in this age group declined 75% (CI = 72%-78%), from 96.7 at baseline to 23.9 in 2003. Incidence rates of VT IPD also declined substantially among persons outside of the PCV7 target population (Figure 1). For persons aged \geq 5 years, VT disease decreased 62% (CI = 59%–66%) from 1998–1999 to 2003, with the largest absolute rate reduction occurring among those aged ≥ 65 years (rate difference: 21.7) cases per 100,000 [rate 33.6 during 1998-1999 and 11.9 during 2003]). Total IPD incidence declined 29% (CI = 25%-33%), again with the majority of the absolute rate reduction occurring among those aged ≥ 65 years (rate difference: 18.4) cases per 100,000 [rate 60.1 during 1998-1999 and 41.7





* Per 100,000 population.

For each age group, the decrease in VT IPD rate for 2003 compared with the 1998–1999 baseline is statistically significant (p<0.05).

* Proposed.

during 2003]). The incidence of IPD caused by the 16 serotypes included in the 23-valent polysaccharide pneumococcal vaccine (PPV23) and not in PCV7 among persons aged \geq 5 years increased 11% (CI = 3%–21%) from 1998–1999 to 2003.

Analysis of the projected 29,599 VT IPD cases prevented nationally by PCV7 in 2003 compared with 1998–1999 (Table) revealed that the majority (69%) of cases were prevented through indirect effects of the vaccine. An estimated 9,140 cases of VT IPD were directly prevented by vaccinating children aged <5 years with PCV7; an additional 20,459 cases of VT IPD were prevented through indirect effects of the vaccine across all ages (Figure 2). Incidence of IPD caused by pneumococcal serotypes not included in PCV7 increased among children aged <5 years and adults aged \geq 40 years, with a total of 4,721 projected additional cases of NT IPD in 2003 compared with the 1998–1999 baseline (Table). After

TABLE. Changes in projected numbers of invasive pneumococcal disease (IPD) cases, by age group and serotype category — Active Bacterial Core surveillance (ABCs), United States, 1998–1999 and 2003

Age		1998–1999 average projected	2003 projected	Change in annual projected
group (yrs)	Serotype category*	no. of cases [†]	no. of cases†	no. of cases
<5				
	Vaccine	14,293	876	-13,417
	Nonvaccine	2,947	3,578	631
	Total	17,240	4,454	-12,786
5–17				
	Vaccine	1,195	569	-626
	Nonvaccine	880	824	-56
	Total	2,075	1,393	-682
18–39				
	Vaccine	5,023	1,610	-3,413
	Nonvaccine	3,419	3,407	-12
	Total	8,442	5,017	-3,425
40–64				
	Vaccine	8,945	4,167	-4,778
	Nonvaccine	7,545	10,237	2,692
	Total	16,490	14,404	-2,086
>65				
_	Vaccine	11,595	4,230	-7,365
	Nonvaccine	9,169	10,635	1,466
	Total	20,764	14,865	-5,899
All ages				
0	Vaccine	41,051	11,452	-29,599
	Nonvaccine	23,960	28,681	4,721
	Total	65,011	40,133	-24,878

* Serotypes included in the 7-valent pneumococcal conjugate vaccine are defined as vaccine serotypes (4, 6B, 9V, 14, 18C, 19F, and 23F). All other serotypes are considered nonvaccine serotypes.

Annual national projections of IPD cases were calculated by applying age- and race-specific disease rates for the aggregate ABCs surveillance area to the age and racial distribution of the U.S. population on the basis of 2000 U.S. Census data.

FIGURE 2. Estimated number of cases of vaccine-type (VT) invasive pneumococcal disease (IPD) prevented by direct* and indirect[†] effects of pneumococcal conjugate vaccine (PCV7) — Active Bacterial Core surveillance, United States, 2003



* Direct VT IPD cases prevented in 2003 = 1998–1999 average number of VT IPD cases in children aged <5 years x 2003 PCV7 coverage with _3 doses (68.1%) x PCV7 effectiveness for VT IPD (93.9%).

¹ Indirect VT IPD cases prevented in 2003 = (1998–1999 average number of VT IPD cases across all age groups – 2003 number of VT IPD cases across all age groups) – 2003 direct VT IPD cases prevented. Calculation of indirect cases prevented does not account for replacement disease.

accounting for this increase, 24,878 net cases of IPD were prevented in 2003; net prevented cases were evenly distributed between the age group targeted for vaccination with PCV7 (12,786 prevented cases [51%]) and older children and adults outside the target population (12,092 prevented cases [49%]) (Table).

Reported by: A Reingold, MD, California Emerging Infections Program, Oakland, California. J Hadler, MD, Emerging Infections Program, Connecticut Dept of Public Health. MM Farley, MD, Georgia Emerging Infections Program, Veterans Affairs Medical Center and Emory Univ School of Medicine, Atlanta, Georgia. L Harrison, MD, Maryland Emerging Infections Program, Johns Hopkins Bloomberg School of Public Health, Baltimore, Maryland. R Lynfield, MD, J Besser, MS, Minnesota Dept of Health. N Bennett, MD, Monroe County Dept of Public Health, Rochester, New York. A Thomas, MD, Oregon Dept of Human Svcs. W Schaffner, MD, Tennessee Emerging Infections Program, Vanderbilt Univ Medical Center, Nashville, Tennessee. B Beall, PhD, Streptococcus Laboratory; T Pilishvili, MPH, Office of Surveillance, Active Bacterial Core surveillance/Emerging Infections Program Network; CG Whitney, MD, M Moore, MD, Div of Bacterial and Mycotic Diseases, National Center for Infectious Diseases; DC Burton, MD, EIS Officer, CDC.

Editorial Note: Routine use of PCV7 in young children has reduced the incidence of VT and overall IPD in children and adults, and these reductions have increased since 2001 (2).

The most substantial decline in the rate of VT disease has been in the target population of children aged <5 years. Data from 2003 also demonstrate statistically significant reductions in the rates of both VT IPD and total IPD for children aged 5–17 years, whereas no statistically significant change in disease rate was observed among persons aged 5–19 years in 2001 (2). As of 2003, the total incidence of IPD in persons aged \geq 65 years declined to 41.7 cases per 100,000 population in ABCs surveillance areas, meeting the *Healthy People 2010* objective of no more than 42 cases per 100,000 for this age group (7).

Indirect benefits of PCV7 (i.e., cases prevented in unvaccinated persons) exceeded direct protective benefits among immunized children, with more than twice as many cases of VT IPD prevented indirectly as directly in 2003. The indirect effects of PCV7 are believed to be caused by decreased nasopharyngeal carriage of VT strains among immunized children, which results in decreased transmission to nonimmunized children and adults (i.e., herd immunity) (2,8). On the basis of this mechanism, indirect benefits from PCV7 might be expected to increase as its vaccination coverage increases. In certain populations (e.g., children aged <5 years and adults aged \geq 40 years), the reduction in VT IPD attributable to PCV7 was partially offset by an increase in disease caused by non-VT strains. However, during 2003, the overall magnitude of this replacement disease was small compared with the reduction in VT disease.

The findings in this report are subject to at least two limitations. First, secular trends cannot be excluded as a factor in the changing pattern of IPD in the United States. However, these trends would be expected to affect disease caused by all serotypes; the reductions in IPD after introduction of PCV7 have been specific to vaccine serotypes, suggesting a vaccine effect. The decline in adult IPD likely is not attributable to PPV23, given that no decline occurred in the incidence of IPD caused by serotypes included in PPV23 but not in PCV7, and given that the slight increase in vaccine coverage of PPV23 since 1998 (9) would not be expected to cause a measurable change in IPD rate. Second, the calculations of direct and indirect effects of the conjugate vaccine were based on data estimates from several sources, each with an associated margin of error; the calculations in this report provide only crude estimates of the relative magnitudes of direct and indirect vaccine effects. In addition, the number of doses of vaccine needed to provide direct protection is unknown, and partial protection might be provided by fewer than 3 doses.

The robustness of the direct and indirect effects of PCV7 has important implications for cost-benefit analyses of similar vaccines in the United States and internationally. Initial estimates of cost-effectiveness for the United States (10) did not account for indirect effects and therefore underestimated the cost-effectiveness of PCV7. In addition, ongoing surveillance will be required to monitor the balance of disease reduction versus replacement in the conjugate vaccine era, particularly in vulnerable populations (e.g., the elderly and immunocompromised persons), who might be more susceptible to less virulent non-VT strains of pneumococci. Such information will be critical for determining whether the composition of conjugate vaccines should be revised or expanded over time.

Acknowledgments

This report is based, in part, on contributions by P Daily, MPH, G Rothrock, MPH, California Emerging Infections Program, Oakland, California. N Barrett, MPH, Emerging Infections Program, Connecticut Dept of Public Health. W Baughman, MSPH, J Howgate, MPH, C Payne, MPH, L Rainer, MPH, P Martell-Cleary, MSW, Georgia Emerging Infections Program, Veterans Affairs Medical Center and Emory Univ School of Medicine, Atlanta, Georgia. L Sanza, Maryland Emerging Infections Program, Johns Hopkins Bloomberg School of Public Health, Baltimore, Maryland. C Lexau, PhD, R Danila, PhD, B Juni, MS, G Kupferschmidt, Minnesota Dept of Health. C Long, Univ of Rochester, Rochester, New York; B Anderson, D Hoefer, New York State Dept of Health. K Stefonek, MPH, Oregon Dept of Human Svcs. B Barnes, Vanderbilt Univ Medical Center, Nashville; AS Craig, MD, Tennessee Dept of Health. TH Skoff, MS, ER Zell, MStat, C Wright, Div of Bacterial and Mycotic Diseases, National Center for Infectious Diseases, CDC.

References

- 1. CDC. Preventing pneumococcal disease among infants and young children: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 2000;49(No. RR-9).
- 2. Whitney CG, Farley MM, Hadler J, et al. Decline in invasive pneumococcal disease after introduction of protein-polysaccharide conjugate vaccine. N Engl J Med 2003;348:1737–46.
- 3. Flannery B, Schrag S, Bennett NM, et al. Impact of childhood vaccination on racial disparities in invasive *Streptococcus pneumoniae* infections. JAMA 2004;291:2197–203.
- 4. National Center for Health Statistics. Bridged-race vintage 2003 postcensal population estimates for July 1, 2000–July 1, 2003, by year, county, single-year of age, bridged-race, Hispanic origin, and sex. Available at http://www.cdc.gov/nchs/about/major/dvs/popbridge/ datadoc.htm.
- National Immunization Program. Estimated vaccination coverage with individual vaccines and selected vaccination series among children 19–35 months of age by state and immunization action plan area, US, National Immunization Survey (NIS), 2003. Available at http:// www.cdc.gov/nip/coverage/NIS/03/toc-03.htm.
- Black S, Shinefield H, Fireman B, et al. Efficacy, safety and immunogenicity of heptavalent pneumococcal conjugate vaccine in children. Pediatr Infect Dis J 2000;19:187–95.
- 7. US Department of Health and Human Services. Healthy people 2010, 2nd ed. Understanding and improving health and objectives for improving health (2 vols.). Washington, DC: US Department of Health and Human Services; 2000.

- O'Brien KL, Dagan R. The potential indirect effect of conjugate pneumococcal vaccines. Vaccine 2003;21:1815–25.
- 9. CDC. Behavioral Risk Factor Surveillance System (BRFSS). Available at http://www.cdc.gov/brfss/index.htm.
- Lieu TA, Ray GT, Black SB, et al. Projected cost-effectiveness of pneumococcal conjugate vaccination of healthy infants and young children. JAMA 2000;283:1460–8.

Improper Disposal of Hazardous Substances and Resulting Injuries — Selected States, January 2001–March 2005

Many consumer and industrial products, including fuels, solvents, fertilizers, pesticides, paints, and household cleaning disinfectants, contain hazardous substances. Improper disposal of these materials can lead to unexpected releases of toxins that are hazardous to humans and harmful to the environment. This report summarizes all known events involving improper disposal of hazardous substances reported to the Agency for Toxic Substances and Disease Registry (ATSDR) during January 2001–March 2005, describes four illustrative case reports, and provides recommendations for preventing injury resulting from improper disposal.

ATSDR maintains the Hazardous Substances Emergency Events Surveillance (HSEES) system to collect and analyze data about the public health consequences (i.e., morbidity, mortality, and evacuation) of hazardous-substance-release events.* The information in this report is based on events reported to HSEES from 18 participating state health departments[†] during January 2001–March 2005.[§] Improper disposal events are defined as events in which a hazardous substance is placed in municipal waste and subsequently causes a release or potential release that requires (or would have required) removal, clean-up, or neutralization according to federal, state, or local law.

Summary of HSEES Data

A total of 36,784 events involving release of hazardous substances were reported to HSEES during January 2001–March 2005. Of these, 107 (0.3%) were associated with improper disposal. All 18 states reported this type of event, with New York (47 [44%] events) and Washington (13 [12%]) reporting the most events. Sixteen (15%) events involved fires or explosions. Of the 159[¶] known improper disposal locations, releases occurred most frequently in residential (59 [37%]) and commercial settings (53 [33%]). Of the 284** total substances involved in improper disposal events, the most common substances were hydrochloric acid (24 [8%]), acid not otherwise specified (15 [5%]), and iodine-131 (six [2%]).

Of the 107 events, 35 (33%) resulted in injuries to 69 persons, 64 (93%) of whom were categorized as employees. HSEES does not collect specific information on type of employee injured (e.g., sanitation worker). However, evaluation of the comment field on incidence reports indicated that more than half (39 [57%]) of the 64 injured employees were sanitation workers.

The 69 injured persons had a total of 101 reported injuries, most frequently respiratory irritation (46 [46%]), dizziness or other central nervous system symptoms (12 [12%]), eye irritation (11 [11%]), and burns (nine [9%]). Forty-two (61%) injured persons were treated at hospitals but not admitted, 11 (16%) were treated at the scene, four (6%) were examined by private physicians, three (4%) were treated at hospitals and admitted, and three (4%) were sent to hospitals for observation. The remaining six (9%) persons experienced adverse health effects within 24 hours of exposure; these injuries were reported through official channels (e.g., fire or police departments, emergency medical services, or poison control centers). No deaths occurred.

Evacuation was ordered for 13 (12%) of the 107 events. The number of evacuees was known for nine of the events, for which 74 persons were known to have evacuated; the number of persons per event ranged from two to 25 (median: six persons per event). The median length of evacuation was 3 hours (range: 1–82 hours).

Of the 97 (91%) events for which decontamination status was known, decontamination of potentially exposed persons was necessary in 31 (32%) events. Ninety-two persons were decontaminated; of these, 61 (66%) were emergency responders, 29 (32%) were employees (i.e., sanitation workers or

^{*}An HSEES event is the acute release or threatened release of a hazardous substance into the environment in an amount that requires (or would have required) removal, cleanup, or neutralization according to federal, state, or local law (1). A hazardous substance is one that can reasonably be expected to cause an adverse health effect upon exposure.

[†] Alabama, Colorado, Florida, Iowa, Louisiana, Michigan, Minnesota, Mississippi, Missouri, New Jersey, New York, North Carolina, Oregon, Rhode Island, Texas, Utah, Washington, and Wisconsin.

[§] Data through March 31, 2005, were the most recent available when the analysis was conducted; data for 2004 and 2005 are provisional.

⁵ Exceeds the number of events because some events may occur in mixed-use locations (e.g., in a residential and commercial area).

^{**} Exceeds the number of events because certain events involved multiple substances.

employees of the industry involved in the release), and two (2%) were members of the general public.

Case Reports

The following case reports illustrate the danger involved in improper disposal of hazardous substances.

New York. In June 2004, a sanitation truck compacted an improperly disposed of container of hydrochloric acid, releasing approximately 10 gallons of the hazardous substance into a commercial/residential area. Two male sanitation workers sustained chemical burns and were decontaminated on the scene, treated at a hospital, and released. A hazardous materials (HazMat) team, law enforcement officials, fire department officials, and emergency medical services personnel responded to the event.

Colorado. In March 2003, a hospital employee improperly disposed of an unknown quantity of radioactive waste in a dumpster. The dumpster contents were picked up by a garbage truck. Later, as the garbage truck approached the land-fill, the contents activated radiation detectors at the landfill. No injuries were reported; however, four first responders were decontaminated at the site. Access to the landfill was restricted until the radioactive waste was removed. A company emergency response team, fire department officials, and hospital personnel responded to the event.

Washington. In June 2002, hydrochloric acid used in an illicit methamphetamine laboratory was disposed of in an apartment building dumpster. Later, a male sanitation worker sustained respiratory irritation when the acid was dumped into the back of his truck. After the exposure occurred, his supervisor took the worker to a physician for observation. Law enforcement officials, fire department officials, emergency medical services personnel, and an environmental agency responded to the event.

Wisconsin. In August 2001, a sanitation truck compacted an improperly disposed of container of hydrochloric acid, releasing approximately 1 gallon of the hazardous substance into a residential area. The sanitation truck driver sustained chemical burns after coming into contact with the acid. He was transported to a hospital, treated for his injury, and released. A HazMat response team responded to the event.

Reported by: *DK Horton, MSPH, S Rossiter, MPH, MF Orr, MS, Div of Health Studies, Agency for Toxic Substances and Disease Registry.*

Editorial Note: This report illustrates the dangers associated with improper disposal of hazardous substances. Although improper disposal events accounted for a limited number of hazardous-substance–release events overall, HSEES has been recording approximately 25 such events per year, and the potential for additional events appears substantial. Persons in

the United States generate approximately 1.6 million tons of household hazardous waste each year (2). An average household can accumulate as much as 100 pounds of hazardous waste in basements, garages, and storage closets (2). In addition, industries and businesses in the United States generate more than 40 million tons of hazardous waste annually (3).

Because many hazardous substances are toxic, flammable, corrosive, explosive, or even radioactive, they can be dangerous when disposed of improperly. Of particular concern is the hazard to sanitation workers because sanitation trucks, especially those with compactors, can easily breach hazardous substance containers, resulting in releases and mixing of substances. During this reporting period, more than half the injured persons were sanitation workers.

At least five of the events were caused by improper disposal of hazardous substances used in illicit methamphetamine laboratories (e.g., hydrochloric acid, ether, and acetone). Substances used in methamphetamine production, many of which are volatile, are often disposed of in municipal waste containers. When these substances are discarded and compacted, the potential for a hazardous release, fire, and explosion is increased.

The majority of the 101 reported injuries examined in this analysis were not life threatening, and no deaths occurred during the reporting period. However, a previous HSEES analysis described the death of a sanitation worker exposed to an improperly disposed of container of hydrofluoric acid (4).

The findings in this report are subject to at least two limitations. Reporting of events to HSEES is not mandatory; therefore, participating state health departments might not be informed about every event. Second, only 18 state health departments provided data to HSEES during the reporting period; therefore, these data underrepresent the total hazardous-substance-release events in the United States.

The findings suggest the need for development and implementation of effective public health strategies to prevent improper disposal practices or injuries resulting from those practices (2,5-7). Such strategies include educating the public regarding proper methods for disposing of hazardous substances, promoting the use of alternative products that do not contain hazardous substances, and organizing community collection days for disposal of hazardous substances (Box).

Acknowledgments

The findings in this report are based, in part, on contributions by T Arant, Alabama Dept of Public Health. C Kelley, Colorado Dept of Health. A Becker, PhD, Florida Dept of Health. D Cooper, Iowa Dept of Public Health. K Lanier, Louisiana Dept of Health and Hospitals. M Stanbury, MPH, Michigan Dept of Community

BOX. Preventing improper disposal of hazardous substances and resulting injuries

General public

- Learn the proper methods for disposing of hazardous substances.
- Understand the dangers associated with improper disposal of hazardous substances.
- Read container labels for proper use and disposal recommendations.
- Be certain a toxic product is needed before using it.
- Use alternative products that do not contain hazardous substances.
- Purchase the smallest possible quantity of a product.
- Use leftover chemicals for other projects, or share them with other persons (e.g., neighbors).
- Community leaders
- Place waste containers (e.g., dumpsters) in well-lit, secured areas.
- Train sanitation workers to recognize discarded methamphetamine laboratory chemicals and equipment.
- Establish collection days for hazardous substances.

Health. N Rice, MPH, Minnesota Dept of Health. R Mozingo, Mississippi State Dept of Health. C Henry, Missouri Dept of Health and Senior Svcs. J Savrin, New Jersey Dept of Health and Senior Svcs. R Wilburn, MPH, New York State Dept of Health. S Giles, MPH, North Carolina Dept of Health and Human Svcs. T Tsongas, PhD, Oregon Public Health Svcs. L Phillips, Rhode Island Dept of Health. R Harris, Texas Dept of Health. W Ball, PhD, Utah Dept of Health. W Clifford, Washington Dept of Health. J Drew, Wisconsin Dept of Health and Family Svcs.

References

- Agency for Toxic Substances and Disease Registry. Hazardous Substances Emergency Events Surveillance System (HSEES), 2001 annual report. Atlanta, GA: US Department of Health and Human Services, Agency for Toxic Substances and Disease Registry; 2001. Available at http:// www.atsdr.cdc.gov/HS/HSEES/annual2001.html.
- US Environmental Protection Agency. Household hazardous waste. Washington, DC: US Environmental Protection Agency; 2005. Available at http://www.epa.gov/epaoswer/non-hw/muncpl/hhw.htm.
- US Environmental Protection Agency. Basic facts about waste. Washington, DC: US Environmental Protection Agency; 2004. Available at http://www.epa.gov/epaoswer/osw/facts.htm.
- Horton DK, Berkowitz Z, Kaye WE. Secondary contamination of emergency department personnel from hazardous materials events, 1995– 2001. Am J Emerg Med 2003;21:28–33.
- Minnesota Pollution Control Agency. Household hazardous waste disposal. St. Paul, MN: Minnesota Pollution Control Agency; 1993. Available at http://www.moea.state.mn.us/hhw/w-hhw2-02.pdf.
- CDC. Acute public health consequences from illicit methamphetamine laboratories—selected states, January 2000–June 2004. MMWR 2005;54:356–9.

Update: Influenza Activity — United States and Worldwide, May 22–September 3, 2005, and 2005–06 Season Vaccination Recommendations

Influenza A (H3N2) viruses circulated worldwide, and Influenza A (H1)* and B viruses were reported less frequently during May 22–September 3, 2005. In North America, isolates of influenza A (H3N2), A (H1), and influenza B were identified sporadically. This report summarizes influenza activity in the United States and worldwide since the last *MMWR* update.[†]

United States

In the United States, CDC uses seven systems for national influenza surveillance, including the following four that operate year-round: 1) collaborating laboratories of the World Health Organization (WHO) and the National Respiratory and Enteric Virus Surveillance System (NREVSS) report the number, types, and subtypes of influenza viruses detected; 2) approximately 2,250 sentinel health-care providers report patient visits for influenza-like illness (ILI), and approximately 500 of these providers continue regular reporting throughout the summer; 3) 122 U.S. cities report mortality attributed to influenza and pneumonia on a weekly basis; and 4) a national surveillance system records pediatric deaths associated with laboratory-confirmed influenza (1).

During May 22–September 3,[§] WHO and NREVSS collaborating laboratories tested 14,016 respiratory specimens; 120 (0.9%) were positive for influenza. Of the positive results, 66 (55%) were influenza B viruses, 33 (28%) were influenza A (H3N2) viruses, one (0.8%) was an influenza A (H1) virus, and 20 (17%) were influenza A viruses that were not subtyped. The majority (78%) of these isolates were tested from mid-May through late June, during which time 1.3% of specimens tested were positive for influenza. Since July, 0.4% of specimens tested were positive for influenza.

^{7.} Ruckart PZ, Orr MF, Kaye WE. Hazardous-chemical releases in the home. J Environ Health 2004;67:14–9.

^{*} Includes both the A (H1N1) and A (H1N2) influenza virus types. Although H1N2 viruses have not been identified since February 2004, not all isolated H1 viruses have been tested for the subtype of their neuraminidase. Thus, H1N2 viruses might continue to circulate in some parts of the world. Influenza A (H1N2) viruses appear to have resulted from reassortment of the genes of the circulating influenza A (H1N1) and A (H3N2) subtypes. Because the hemagglutinin proteins of the A (H1N2) viruses are similar to those of the circulating A (H1N1) viruses, and the neuraminidase proteins are similar to the circulating A (H3N2) viruses, the 2005–06 influenza vaccine should provide protection against A (H1N2) viruses.

[†]CDC. Update: influenza activity—United States and worldwide, 2004–05 season. MMWR 2005;54:631–4.

[§]As of September 9, 2005; reporting is incomplete.

During May 22–September 3, the weekly percentage of patient visits to sentinel providers for ILI remained below the national baseline of 2.5%[¶] and ranged from 0.7% to 1.3%. The percentage of deaths attributable to pneumonia and influenza (P&I) as reported by the 122 Cities Mortality Reporting System remained below the epidemic threshold,** and no influenza-related pediatric deaths were reported as occurring during this period.

Worldwide

During May 22–September 3, influenza A (H3N2) viruses predominated in Asia (China, Hong Kong, Japan, Korea, and Thailand). Influenza A (H3N2) viruses were also identified in Oman and Singapore. Influenza A (H1) viruses were reported in China, Hong Kong, India, Indonesia, Japan, Korea, and Malaysia. Influenza B viruses were reported in China, Hong Kong, Indonesia, Korea, Nepal, Philippines, and Thailand.

In Oceania, during the same period, influenza A (H3N2 and non-subtyped) viruses predominated in Australia; influenza B viruses were responsible for outbreaks in New Zealand. Influenza B viruses were also reported in Australia and New Caledonia. In Africa, both influenza A virus subtypes (H3N2 and H1) and influenza B viruses were reported in South Africa, and influenza A (H3N2) and influenza B viruses were reported in Madagascar. Influenza B viruses also were reported in Kenya.

In South America, influenza A (H3N2 and non-subtyped) viruses were associated with regional outbreaks in Argentina and Chile during May 22–September 3 and were reported in Brazil, Colombia, Peru, and Uruguay. Influenza B viruses were associated with an outbreak in Colombia in July and also were reported in Argentina, Brazil, Chile, and Uruguay. Influenza A (H1) viruses were reported in Peru. In North America, influenza A viruses (H3N2 and non-subtyped) and influenza B viruses were reported in Canada, Mexico, and the United States. The United States reported one influenza A (H1) virus. Influenza A (H3N2) viruses also were reported in El Salvador and Panama (2–4).

Characterization of Influenza Virus Isolates

The WHO Collaborating Center for Surveillance, Epidemiology, and Control of Influenza, located at CDC, analyzes influenza-virus isolates received from laboratories worldwide. During May 22–September 3, a total of 77 influenza A (H3N2) viruses (47 from Latin America, 21 from Asia, eight from the United States, and one from Oceania) were collected and characterized antigenically. All 77 influenza A (H3N2) viruses were antigenically related to the A/California/07/2004 reference virus. However, four South American viruses and nine Asian viruses had reduced titers to A/California/07/2004. An A/California/07/2004-like virus was recommended as the H3 component for the 2005–06 Northern Hemisphere vaccine. No influenza A (H1) viruses collected during this period were received and characterized by CDC.

Influenza B viruses circulating worldwide can be divided into two antigenically distinct lineages: B/Yamagata/16/88 and B/Victoria/2/87. Before 1991, B/Victoria lineage viruses circulated worldwide; from late 1991 to early 2001, no viruses of the B/Victoria lineage were identified outside Asia. However, since March 2001, B/Victoria-lineage viruses have been identified in many countries outside Asia, including the United States. Viruses of the B/Yamagata lineage began circulating worldwide in 1990 and continue to be identified. The type-B component of the 2005-06 influenza vaccine (B/Shanghai/ 361/2002-like) belongs to the B/Yamagata lineage. Of the 46 influenza B isolates collected during May 22-September 3 and characterized antigenically at CDC, three belonged to the B/Yamagata lineage, and 43 belonged to the B/Victoria lineage. All three of the B/Yamagata-lineage viruses had reduced titers to B/Shanghai/361/2002. Two of the B/ Yamagata-lineage viruses were from Asia, and one was from the United States. Of the 43 B/Victoria-lineage viruses, 18 came from Asia, 18 from South America, and seven from the United States.

Avian Influenza A (H5N1)

Since December 2003, a total of 11 countries (Cambodia, China, Indonesia, Japan, Kazakhstan, Laos, Malaysia, Russia, South Korea, Thailand, and Vietnam) have reported outbreaks of highly pathogenic avian influenza A (H5N1) virus affecting poultry. Russia and Kazakhstan reported outbreaks of H5N1 virus among poultry for the first time in late July 2005 (5). Mongolia reported detection of H5N1 virus in migratory birds in August (6). In Southeast Asia, where H5N1 continues to be detected among poultry, approximately 150 million birds have died or been culled since 2003 (5).

⁵ The national baseline was calculated as the mean percentage of patient visits for ILI during noninfluenza weeks plus two standard deviations. Wide variability in regional data precludes calculating region-specific baselines and makes applying the national baseline to regional data inappropriate. National and regional percentages of patient visits for ILI are weighted on the basis of state population.

^{**} The expected seasonal baseline proportion of P&I deaths reported by the 122 Cities Mortality Reporting System is projected by using a robust regression procedure in which a periodic regression model is applied to the observed percentage of deaths from P&I during the previous 5 years. The epidemic threshold is 1.654 standard deviations above the seasonal baseline.

Since December 2003, a total of 112 H5N1 cases in humans have been reported to WHO in four countries (Cambodia, Indonesia, Thailand, and Vietnam); 57 (51%) persons died. In August 2005, three cases (including two deaths) were reported in Vietnam. In July, one fatal case was reported in Indonesia (5).

Influenza Vaccine Supply and Recommendations

Vaccination is the primary method for preventing influenza (1). For the 2005–06 influenza vaccine, four manufacturers expect to provide influenza vaccine to the U.S. population. Sanofi Pasteur, Inc., projects production of up to 60 million doses of trivalent inactivated influenza vaccine (TIV). Chiron Corporation projects production of 18–26 million doses of TIV. GlaxoSmithKline, Inc. projects production of 8 million doses of TIV. MedImmune Vaccines, Inc., producer of the nasal-spray, live attenuated influenza vaccine (LAIV), projects production of approximately 3 million doses (7).

Because of the uncertainties regarding production of influenza vaccine, the exact number of available doses and timing of vaccine distribution for the 2005–06 influenza season remain unknown. As a result, CDC recommends that only the following priority groups receive TIV before October 24, 2005:

- persons aged \geq 65 years with comorbid conditions
- residents of long-term-care facilities
- persons aged 2-64 years with comorbid conditions
- persons aged \geq 65 years without comorbid conditions
- children aged 6-23 months
- pregnant women
- health-care personnel who provide direct patient care
- household contacts and out-of-home caregivers of children aged <6 months

These groups correspond to tiers 1A-1C in the previously published table of TIV priority groups in the event of vaccination supply disruption (8). Beginning October 24, 2005, influenza vaccine should be made available to all persons. Healthy persons aged 5–49 years who are not pregnant, including health-care workers who are not caring for severely immunocompromised patients in special-care units, can receive LAIV at any time (1).

Vaccination Recommendations for Persons Displaced by Hurricane Katrina

On September 6, 2005, CDC issued interim vaccination recommendations for persons displaced by Hurricane Katrina (9). Any displaced persons aged ≥ 6 months living in crowded group settings should be administered influenza vaccine;

children aged ≤ 8 years should be administered 2 doses, at least 1 month apart.

Reported by: WHO Collaborating Center for Surveillance, Epidemiology, and Control of Influenza; L Brammer, MPH, A Postema, MPH, R Dhara, MPH, A Balish, T Wallis, H Hall, A Klimov, PhD, T Uyeki, MD, N Cox, PhD, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases; M Katz, MD, EIS Officer, CDC.

Editorial Note: During May 22–September 3, 2005, influenza A (H3N2) viruses were the most frequently reported virus worldwide; however, influenza A (H1) and influenza B viruses also circulated. In North America, sporadic cases of influenza were identified each month. The identification of influenza isolates and even sporadic outbreaks in the summer in North America is not unusual. Neither the influenza virus that will predominate in the United States nor the severity and timing of the 2005–06 season can be predicted.

The ongoing widespread epizootic of highly pathogenic avian influenza A (H5N1) viruses in Asia remains a major public health concern. Since December 2003, a total of 12 countries have reported H5N1 outbreaks in poultry or migratory birds, with human cases reported from four of these countries. Since July 2005, H5N1 infections in poultry and migratory birds have spread beyond their initial focus in Southeast Asia to Kazakhstan, Mongolia, and Russia; a human case was reported in Indonesia for the first time. No evidence of sustained person-to-person transmission has been identified to date, although probable limited person-to-person transmission has been reported (10). To date, no evidence has indicated genetic reassortment among avian influenza A (H5N1) and human influenza A viruses. CDC recommends enhanced surveillance for suspected H5N1 cases among travelers with unexplained severe respiratory illness returning from H5N1affected countries. Additional information about avian influenza is available at http://www.cdc.gov/flu/avian.

Influenza surveillance reports for the United States are posted online weekly during October–May and are available at http://www.cdc.gov/flu/weekly/fluactivity.htm. Additional information about influenza viruses, influenza surveillance, and the influenza vaccine is available at http://www.cdc.gov/ flu.

Acknowledgments

This report is based, in part, on data contributed by WHO collaborating laboratories; National Respiratory and Enteric Virus Surveillance System laboratories; Sentinel Providers Influenza Surveillance System; 122 Cities Mortality Reporting System; WHO National Influenza Centers, Communicable Diseases, Surveillance and Response, WHO, Geneva, Switzerland. A Hay, PhD, WHO Collaborating Centre for Reference and Research on Influenza, National Institute for Medical Research, London, England. I Gust, MD, A Hampson, WHO Collaborating Center for Reference and Research on Influenza, Parkville, Australia. M Tashiro, MD, WHO Collaborating Center for Reference and Research on Influenza, National Institute of Infectious Diseases, Tokyo, Japan.

References

- 1. CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 2005;54(No. RR-8).
- 2. World Health Organization. Influenza. Wkly Epidemiol Rec 2005;80:239–40.
- 3. World Health Organization. Global influenza surveillance. Geneva, Switzerland: World Health Organization; 2005. Available at http:// www.who.int/csr/disease/influenza/influenzanetwork/en.
- 4. Public Health Agency of Canada. FluWatch. Ottawa, Canada: Public Health Agency of Canada; 2005. Available at http://www.phac-aspc.gc.ca/fluwatch/index.html.
- World Health Organization. Communicable disease surveillance and response: avian influenza. Geographical spread of H5N1 avian influenza in birds—Update 28. Geneva, Switzerland: World Health Organization; 2005. Available at http://www.who.int/csr/don/2005_08_18/ en.
- 6. World Organisation for Animal Health (OIE). Highly pathogenic avian influenza in Mongolia in migratory birds. Paris, France: World Organisation for Animal Health (OIE); 2005. Available at http:// www.oie.int/eng/info/hebdo/ais_55.htm.
- 7. CDC. Update: influenza vaccine supply and recommendations for prioritization during the 2005–06 influenza season. MMWR 2005;54:850.
- 8. CDC. Tiered use of inactivated influenza vaccine in the event of a vaccine shortage. MMWR 2005;54:749–50.
- 9. CDC. Interim immunization recommendations for individuals displaced by Hurricane Katrina. Atlanta, GA: US Department of Health and Human Services, CDC; 2005. Available at http://www.bt.cdc.gov/ disasters/hurricanes/katrina/vaccrecdisplaced.asp.
- Ungchusak K, Auewarakul P, Dowell SF, et al. Probable person-toperson transmission of avian influenza A (H5N1). N Engl J Med 2005;352:333–40.

Update: West Nile Virus Activity — United States, 2005

This report summarizes West Nile virus (WNV) surveillance data reported to CDC through ArboNET as of 3 a.m. Mountain Daylight Time, September 13, 2005.

Thirty-seven states have reported 1,299 cases of human WNV illness in 2005 (Figure and Table 1). By comparison, in 2004, a total of 1,386 WNV cases had been reported as of September 14, 2004 (Table 2). A total of 671 (56%) of the 1,193 cases for which such data were available occurred in males; the median age of patients was 50 years (range: 3 months–98 years). Date of illness onset ranged from January 2 to September 8; a total of 29 cases were fatal.

During 2005, a total of 230 presumptive West Nile viremic blood donors (PVDs) have been reported to ArboNET. Of

FIGURE. Areas reporting West Nile virus (WNV) activity — United States, 2005*



* As of September 13, 2005.

these, 71 were reported from California; 37 from Nebraska; 32 from Texas; 20 from South Dakota; 15 from Louisiana; nine from Kansas; seven from Iowa; six each from Arizona and Illinois; five from New Mexico; four from Minnesota; three from Oregon; two each from Alabama, Colorado, Mississippi, and Utah; and one each from Idaho, Michigan, Montana, Nevada, North Carolina, North Dakota, and Penn-sylvania. Of the 230 PVDs, four persons aged 35, 53, 56, and 71 years subsequently had neuroinvasive illness; three persons aged 17, 41, and 51 years subsequently had other illnesses; and 60 persons (median age: 47 years [range: 17–78 years]) subsequently had West Nile fever.

In addition, 2,926 dead corvids and 627 other dead birds with WNV infection have been reported from 39 states. WNV infections have been reported in horses from 28 states, three dogs from Minnesota and Nebraska, four squirrels from Arizona, and two unidentified animal species in two states (Arizona and Illinois). WNV seroconversions have been reported in 675 sentinel chicken flocks from 12 states. One seropositive sentinel horse was reported from Minnesota. A total of 7,822 WNV-positive mosquito pools have been reported from 38 areas (Alabama, Arizona, Arkansas, California, Colorado, Connecticut, District of Columbia, Florida, Georgia, Idaho, Illinois, Indiana, Iowa, Kansas, Louisiana, Maryland, Massachusetts, Michigan, Minnesota, Missouri, Montana, Nebraska, Nevada, New Jersey, New Mexico, New York, Ohio, Oklahoma, Oregon, Pennsylvania, South Carolina, South Dakota, Tennessee, Texas, Utah, Virginia, Washington, and Wisconsin).

Additional information about national WNV activity is available from CDC at http://www.cdc.gov/ncidod/dvbid/ westnile/index.htm and at http://westnilemaps.usgs.gov.

TABLE 1. Number of human cases of West Nile virus (WNV) illness reported, by state - United States, 2005*

State	Neuroinvasive disease [†]	West Nile fever§	Other clinical/ unspecified ¹	Total**	Deaths
Alabama	3	2	0	5	0
Arizona	14	10	5	29	0
Arkansas	1	5	0	6	0
California	175	319	54	548	9
Colorado	5	36	0	41	0
Connecticut	2	0	0	2	0
Florida	4	7	1	12	0
Georgia	1	1	1	3	0
Idaho	2	5	2	9	0
Illinois	71	42	10	123	2
Indiana	1	0	0	1	0
Iowa	2	3	1	6	1
Kansas	2	2	0	4	0
Kentuckv	1	0	0	1	0
Louisiana	50	16	0	66	4
Maryland	1	0	0	1	0
Michigan	5	1	1	7	0
Minnesota	7	13	0	20	1
Mississippi	12	11	0	23	2
Missouri	3	4	2	9	1
Montana	5	6	0	11	0
Nebraska	18	39	0	57	1
Nevada	6	11	0	17	0
New Mexico	10	4	0	14	1
New York	2	1	0	3	0
North Carolin	na 1	1	0	2	0
North Dakota	2	14	0	16	0
Ohio	10	2	0	12	0
Oklahoma	1	0	0	1	0
Oregon	0	3	0	3	0
Pennsylvania	6	5	0	11	0
South Carolir	na 1	0	0	1	1
South Dakota	a 28	140	1	169	1
Tennessee	2	1	0	3	0
Texas	30	6	0	36	4
Utah	10	13	0	23	1
Wisconsin	3	1	0	4	0
Total	497	724	78	1.299	29

* As of September 13, 2005. † Cases with neurologic manifestations (i.e., West Nile meningitis, West Nile encephalitis, and West Nile myelitis).

§ Cases with no evidence of neuroinvasion.
 Illnesses for which sufficient clinical information was not provided.

** Total number of human cases of WNV illness reported to ArboNET by state and local health departments.

TABLE 2. Comparison of human cases and deaths from West Nile virus — United States, 2002–2005

Year	Human cases	Deaths
2002*	1,201	43
2003†	2,923	54
2004 [§]	1,386	35
2005¶	1,299	29

*Data through September 11, 2002. Data through September 10, 2003. Data through September 14, 2004. Data through September 13, 2005.



From 1995 to 1999, the percentage of nursing home residents aged \geq 65 years who received 23-valent pneumococcal polysaccharide vaccine (PPV23) increased by 58.5%. This increase might be attributable, in part, to a 36% increase in the number of residents living in nursing homes with pneumococcal immunization programs. The Advisory Committee on Immunization Practices continues to recommend PPV23 vaccination for all persons aged \geq 65 years and all residents of nursing homes and other long-term–care facilities (CDC. Recommended adult immunization schedule—United States, October 2004–September 2005. MMWR 2004;53:Q1–Q4.)

SOURCES: Bardenheier B, Shefer A, Tiggle RB, Marsteller J, Remsburg RE. Nursing home resident and facility characteristics associated with pneumococcal vaccination: National Nursing Home Survey, 1995–1999. J Am Geriatr Soc 2005;53:1543–51.

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals September 10, 2005, with historical data CASES CURRENT



* 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.

TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending September 10, 2005 (36th Week)*

Disease	Cum. 2005	Cum. 2004	Disease	Cum. 2005	Cum. 2004
Anthrax	_	_	Hemolytic uremic syndrome, postdiarrheal [†]	110	116
Botulism:			HIV infection, pediatric [†]	181	273
foodborne	9	6	Influenza-associated pediatric mortality**	43	_
infant	56	56	Measles	57††	25 ^{§§}
other (wound & unspecified)	19	10	Mumps	193	146
Brucellosis	73	63	Plague	3	1
Chancroid	17	19	Poliomyelitis, paralytic	—	_
Cholera	3	4	Psittacosis [†]	15	8
Cyclosporiasis [†]	673	189	Q fever [†]	83	47
Diphtheria	_		Rabies, human	1	4
Domestic arboviral diseases			Rubella	9	9
(neuroinvasive & non-neuroinvasive):	_	—	Rubella, congenital syndrome	1	_
California serogroup ^{†§}	14	84	SARS [†] **	—	_
eastern equine ^{†§}	11	3	Smallpox [†]	_	_
Powassan ^{†§}	_	1	Staphylococcus aureus:		
St. Louis ^{† §}	2	11	Vancomycin-intermediate (VISA) [†]	_	_
western equine ^{† §}	_		Vancomycin-resistant (VRSA) [†]	_	1
Ehrlichiosis:	_	_	Streptococcal toxic-shock syndrome [†]	91	102
human granulocytic (HGE)†	359	276	Tetanus	15	14
human monocytic (HME) [†]	246	206	Toxic-shock syndrome	71	63
human, other and unspecified [†]	51	50	Trichinellosis ^{¶¶}	13	1
Hansen disease [†]	54	70	Tularemia [†]	88	74
Hantavirus pulmonary syndrome [†]	17	18	Yellow fever	_	_

No reported cases.

Incidence data for reporting years 2004 and 2005 are provisional and cumulative (year-to-date).

Not notifiable in all states. §

Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Infectious Diseases (ArboNet Surveillance).

ſ Updated monthly from reports to the Division of HIV/AIDS Prevention, National Center for HIV, STD, and TB Prevention. Last update June 26, 2005.

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

^{TT} Of 57 cases reported, 46 were indigenous and 11 were imported from another country. ^{SS} Of 25 cases reported, eight were indigenous and 17 were imported from another country. 11

Formerly Trichinosis.

	A	IDS	Chla	mydia [†]	Coccidioidomycosis Cryptosporidiosis		oridiosis	
Reporting area	Cum. 2005§	Cum. 2004	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004
	2000	26.652	622 560	622 921	2 101	2 05/	2 457	2004
	20,400	20,000	020,000	000,021	5,101	0,004	159	107
	//8	208	22,054	20,625	N	N	158	127
NH	20	29	1 283	1 163			20	21
Vt. ¹	4	13	674	780	_	_	23	21
Mass.	368	283	9,864	9,038	—	—	58	51
R.I.	68	98	2,254	2,367			5	4
Conn.	307	422	6,470	5,877	N	N	38	14
MID. ATLANTIC	4,352	5,934	77,907	78,183	_	_	1,502	329
Upstate N.Y.	800	723	15,376	15,552	N	N	1,290	75
N.Y. City	2,327	3,242	24,501	24,283	N	N	53	85
Pa.	651	952	25.189	26.035	N	N	143	132
	1 000	0.000	20,100	110,100		10	600	740
	1,938	2,339	90,088 24 139	27 789	c N	IU N	326	743 169
Ind.	236	264	12.923	12.699	N	Ň	34	55
111.	983	1,106	29,604	32,779	_	_	52	128
Mich.	322	383	16,703	25,946	5	10	64	110
Wis.	85	121	12,719	12,907	N	N	163	281
W.N. CENTRAL	463	578	38,634	38,572	5	5	391	284
Minn.	123	141	7,120	8,110	3	N	80	91
Iowa	50 198	47	4,830	4,630	IN 1	IN 3	70 188	59 55
N. Dak.	5	15	830	1.236	N	N		9
S. Dak.	10	7	1,881	1,685	_	_	16	23
Nebr. ¹	18	35	3,837	3,560	1	2	4	24
Kans.	59	79	4,756	5,130	N	N	27	23
S. ATLANTIC	6,473	8,273	121,873	119,223	1	—	344	351
Del.	100	105	2,259	1,954	N	N	_	
	812	988	12,807	13,043	1	_	23	14
Va. ¹	307	472	14.310	15.266	_	_	22	38
W. Va.	36	55	1,808	1,949	Ν	Ν	9	4
N.C.	531	416	22,985	20,104	N	N	44	54
S.C. ¹	386	504	15,058	13,348	—	—	9	17
Ga. Fla	2 731	4 049	20,000	22,242	N	N	165	89
	1,000	1,010	45,001	20,000			70	07
E.S. CENTRAL	1,093	1,322	45,391	41,119	N	5 N	72	97
Tenn. ¹	434	533	16.560	15.521	N	N	22	29
Ala. ¹	295	305	8,258	9,394	_	_	15	16
Miss.	229	327	14,127	12,313	_	5	2	23
W.S. CENTRAL	2,206	3,151	75,337	78,744	1	2	59	72
Ark.	72	135	5,872	5,578		1	3	13
La.**	436	639	12,572	16,217	1	1	3	3
Okia. Tex 1	1 5 3 1	2 247	7,710 49.183	7,735	IN N	N	33	17
	700	2,247	40,100	40,214	0.100	0.470	20	100
MOUN IAIN Mont	789	933	36,237	38,605	2,136 N	2,470 N	90 14	129
Idaho ¹	9	16	1.655	1.946	N	Ň	6	18
Wyo.	2	13	765	747	3	2	2	3
Colo.	163	162	9,229	9,783	N	N	32	43
N. Mex.	72	138	3,272	6,140	2 090	18	3	11
Utah	33	51	2 976	2 559	2,009	2,392	15	3
Nev. ¹	177	193	4,523	4,470	31	45	8	2
PACIFIC	2 313	3 258	110 039	106 630	953	1 462	202	200
Wash.	229	288	12,787	12,084	N	N	30	23
Oreg. ¹	136	216	5,641	5,635	_	—	45	28
Calif.	1,874	2,658	86,219	82,469	953	1,462	125	147
Alaska Hawaii	14	29	2,683	2,638	_	_	1	2
Our	00	10	2,703	3,004	_	—	I	2
Guam	1 527	1	2 5 9 4	/89 2 5 2 2	N		N	N
V.I.	10	10	119	259				
Amer. Samoa	Ŭ	Ŭ	Ŭ	Ŭ	U	U	U	U
C.N.M.I.	2	U		U		U	_	U

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending September 10, 2005, and September 11, 2004 (36th Week)*

N: Not notifiable. U: Unavailable. —: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. *Incidence data for reporting years 2004 and 2005 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, National Center for HIV, STD, and TB Prevention. Last update June 26, 2005. [¶] Contains data reported through National Electronic Disease Surveillance System (NEDSS). ** Because of Hurricane Katrina, weekly reporting has been disrupted.

(Sour week)										
		Escher	<i>richia coli</i> , Ente	rohemorrhagio	: (EHEC)					
			Shiga tox	in positive,	Shiga toxi	n positive,			_	
	015	7:H7	serogrou	p non-O157	not sero	grouped	Giard	iasis	Gond	orrhea
Reporting area	2005	2004	2005	2004	2005	2004	2005	2004	2005	2004
UNITED STATES	1,336	1,639	186	183	170	119	11,081	12,712	212,703	222,870
NEW ENGLAND	96	114	35	38	17	10	1,018	1,186	4,085	4,810
Maine	12	9	6		_	—	136	99	93	154
Vt.	10	14	2		_	_	107	20 121	38	61
Mass.	36	51	6	13	17	10	425	524	1,788	2,136
R.I. Conn	3 24	6 23		1 19	_	_	70 242	68 348	308 1 744	604 1 768
MID ATLANTIC	171	190	19	28	26	28	2 051	2 692	22 149	25 259
Upstate N.Y.	73	81	11	12	8	14	737	875	4,390	5,085
N.Y. City	7	33	_	_	_	_	523	762	6,458	7,835
N.J. Pa.	27 64	36 40	2	5 11	5 13	6 8	240 551	348 707	3,864 7,437	4,736 7.603
E.N. CENTRAL	266	317	17	38	8	19	1.748	1.989	38.651	46.777
Ohio	80	66	4	7	3	10	504	541	11,476	14,330
Ind.	37	36					N 254	N 570	5,175	4,570
Mich	45 57	58		7	4	3	354 491	572 455	6 506	14,228
Wis.	47	86	12	17	_	_	399	421	3,584	3,294
W.N. CENTRAL	222	347	22	25	29	20	1,270	1,393	12,405	11,650
Minn.	54	79		10	17	4	562 173	504	2,010	2,023
Mo.	57	56	9	12	5	6	290	381	6.397	6.070
N. Dak.	3	11	_	_	_	6	7	18	50	80
S. Dak.	16	27	3			—	63	42	252	186
Kans.	28	53 26			4 3	4	58 117	151	1,726	1,726
S. ATLANTIC	120	114	48	20	67	25	1,567	1,972	52,625	53,984
Del.	3	2	N	N	N	N	31	34	563	616
Md.	22	20	19	3	6	3	123	81 51	4,795	5,598
Va.	19	23	16	9	12	_	323	334	5,101	6,149
W.Va.	1	2	_	—	1		30	27	491	622
N.C.			—	—	38	16	N 67	N 70	10,811	10,643
Ga.	17	15	9	6	_	_	318	615	9.515	9.721
Fla.	54	42	4	2	10	6	640	751	13,304	12,255
E.S. CENTRAL	90	75	1	3	16	13	268	261	17,480	17,905
Ky. Tenn	28	18	1	1	13	6	N 136	N 142	2,139	1,700
Ala.	22	14	_	_	_	_	132	119	4,851	5,699
Miss.	5	10	—	2		—	—	—	4,533	4,791
W.S. CENTRAL	34	64	4	3	3	4	193	215	30,592	30,259
Ark.	3	3	3	1	2	_	27	84 37	6,950	2,896
Okla.	16	14	_	_	_	_	108	94	3,125	3,289
Tex.	9	36	1	2	1	4	N	N	17,445	16,558
MOUNTAIN	119	160	34	27	4	_	888	1,036	7,879	8,100
Idaho	10	37	8	7	2	_	53	123	68	57
Wyo.	4	6	2	1	—	_	17	16	49	40
Colo.	25	41	1	1	1	—	344	368	2,095	2,091
Ariz.	26	10	N	N	N	N	43 97	130	2.750	2.610
Utah	27	26	18	12	_	_	246	217	457	401
Nev.	9	14	—	1	1	—	41	83	1,757	2,029
PACIFIC	218	258	6	1	—		2,078	1,968	26,837	24,126
Oreg	50	88 50	6	1	_	_	234 244	309	2,490	1,829
Calif.	91	114	_	_	_	_	1,490	1,319	22,479	20,197
Alaska	12	1	—	—	—	—	67	57	375	423
	9	5	_	_	_		43	59	494	895
Guam PR	N 1	N 1	_	_	_	_	97	2 181	245	122 184
V.I.		_	_	_	_	_			35	76
Amer. Samoa	U	U	U	U	U	U	U	U	U	U
U.IN.IVI.I.		U		U		U		U		L1

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending September 10, 2005, and September 11, 2004 (36th Week)*

N: Not notifiable. U: Unavailable. —: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Incidence data for reporting years 2004 and 2005 are provisional and cumulative (year-to-date).

<u>.</u>		Haemophilus influenzae, invasive										
	All	ages			Age <	5 years						
	All sei	rotypes	Serc	otype b	Non-se	rotype b	Unknown	serotype				
Reporting area	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004				
UNITED STATES	1,484	1,412	3	9	81	80	145	135				
NEW ENGLAND	120	126	_	1	10	8	5	1				
Maine	6	10	—	—	—		1	—				
Vt.	6	5	_	_	_		2	1				
Mass.	56	62	—	1	3	3	1	—				
K.I. Conn.	39	332	_	_	2 5	3	1	_				
MID. ATLANTIC	293	287	_	1	_	4	37	32				
Upstate N.Y.	85	98	_	1	_	4	8	5				
N.Y. City	53 55	65 53	_	_	_	_	10	12				
Pa.	100	71	—	_	—	—	10	13				
E.N. CENTRAL	214	268	1	_	3	8	15	41				
Ohio	91 52	78	_	_		2	9	14				
III.	35	94	_	_	_		3	20				
Mich.	15	17	1	—	—	2	2	4				
WIS.	21	41	—				1	2				
Minn.	36	34	_	2 1	3	3	9	<u> </u>				
Iowa	1	1	_	1	_	_	_	_				
Mo. N Dak	28 1	29	_	_	_	_	5	6				
S. Dak.	_	_	_	_	_	_	_	_				
Nebr.	7	4	_	_	_	_	1	1				
S ATLANTIC	356	323	1	_	22	21	20	22				
Del.			_	_				<u> </u>				
Md.	52	50	—	—	5	5	—	-				
Va.	34	30	_	_	_	_	1	3				
W.Va.	22	15	_	_	1	4	4	_				
N.C. S.C.	63 20	44 10	1	_		5	1	1				
Ga.	71	89	—	_		—	10	16				
Fla.	94	83	—	—	9	7	4					
E.S. CENTRAL	85	57	_	1	1	_	14	7				
Tenn.	59	38	_	_	_	_	8	5				
Ala.	18	12	—	1	—	—	4	2				
		2	_				_	_				
Ark.	84 4	55			1	6	6					
La.	28	10	1	_	2	_	6	1				
Okla. Tex	51	43	_	1	4	6	_	_				
ΜΟΙΙΝΤΔΙΝ	167	148	_	3	13	20	29	17				
Mont.		_	_	_				_				
Idaho	3	5	_	—	—	—	1	2				
Colo.	34	36	_	_	_	_	9	4				
N. Mex.	16	31	—	—	4	6	2	6				
Ariz. Utah	84 13	53 12	_	2		9	8	2				
Nev.	13	11	—	1	2	3	2	1				
PACIFIC	84	71	—	_	22	10	10	6				
Wash.	3	1 20	—	—	—	—	2	1				
Calif.	39	25	_	_	22	10	2	1				
Alaska	5	5	—	—	—	—	1	1				
Guam	ŏ	ð	_	—	_	_	_	I				
P.R.	3	2	_	_	_	_	1	2				
V.I.												
C.N.M.I.	<u> </u>	U	<u> </u>	U	<u> </u>	U	<u> </u>	U				

 TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 10, 2005, and September 11, 2004

 (36th Week)*

N: Not notifiable. U: Unavailable. —: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Incidence data for reporting years 2004 and 2005 are provisional and cumulative (year-to-date).

908

Vo	l. 54 /	/ No.	36

(Sour week)		Hepatitis (viral, acute), by type									
		Α		B		С					
Reporting area	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004					
UNITED STATES	2,588	4,062	3,712	4,027	556	525					
NEW ENGLAND Maine N.H.	353 1 69	688 11 15	194 12 16	251 1 26	9	12 — —					
Mass. R.I. Conn.	5 231 10 37	8 579 17 58	135 1 28	5 133 3 83	- - U	4 7 					
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	439 69 200 93 77	514 57 216 120 121	737 57 74 453 153	530 51 108 152 219	74 13 — 61	85 5 80					
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	242 36 36 55 98 17	328 37 39 107 107 38	324 98 31 79 116 —	392 84 31 63 183 31	91 3 19 69 	74 4 7 13 50 —					
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. S. Dak. Nebr.	62 3 16 28 — 4	118 28 34 25 1 3 10	190 20 18 111 <u>-</u> 3 19	243 37 14 147 4 1 27	30 5 23 1 - 1	18 15 3 					
Kans. S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	11 447 4 45 2 53 4 57 23 75 184	17 746 5 84 5 82 3 70 37 258 202	19 951 38 108 99 27 112 95 115 347	13 1,245 30 114 15 166 28 129 98 326 339							
E.S. CENTRAL Ky. Tenn. Ala. Miss.	187 25 124 22 16	122 29 75 6 12	242 49 90 57 46	348 43 169 56 80	72 13 14 10 35	70 23 23 4 20					
W.S. CENTRAL Ark. La. Okla. Tex.	140 6 44 4 86	496 59 37 18 382	286 28 31 25 202	240 85 42 49 64	49 — 9 3 37	70 2 3 3 62					
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	231 7 15 30 18 137 17 7	317 5 14 4 38 18 193 31 14	375 3 1 34 7 268 33 22	315 1 7 44 14 158 27 54	32 1 16 7 7	33 2 1 2 8 U 5 3 12					
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	487 30 33 404 3 17	733 40 52 617 4 20	413 50 69 283 7 4	463 39 79 327 10 8	29 U 13 16 —	37 U 14 22 — 1					
Guam P.R. V.I. Amer. Samoa C.N.M.I.	52 — U	1 30 	30 U	12 59 — U U	 	9 U U					

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending September 10, 2005, and September 11, 2004 (36th Week)*

N: Not notifiable. U: Unavailable. —: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Incidence data for reporting years 2004 and 2005 are provisional and cumulative (year-to-date).

(36th Week)*	Logion		Lioto	riagio	Lumo	diagona	Mal	orio
	Cum	Cum	Cum	Cum	Cum	Cum	Cum	
Reporting area	2005	2004	2005	2004	2005	2004	2005	2004
UNITED STATES	1,179	1,340	464	475	13,355	12,863	791	999
NEW ENGLAND	71	57	36	28	1,423	2,238	49	72
Naine N.H.	3	1 4	1 5	5	99 138	29 155	5	6 4
Vt.	3	3	1	1	21	39	1	3
Mass. R I	25 12	26 8	10 5	9	/14 25	1,248 152	24	44
Conn.	22	15	14	10	426	615	12	13
MID. ATLANTIC	408	352	123	114	9,343	7,985	211	264
Upstate N.Y. N Y City	109 50	66 49	37 21	31 19	2,508	2,497	32 102	30 137
N.J.	79	54	27	24	3,176	2,129	51	59
Pa.	170	183	38	40	3,659	3,077	26	38
E.N. CENTRAL Ohio	204 102	337 159	47 21	88 31	585 53	1,087	63 16	95 24
Ind.	13	31	2	16	18	19		10
III. Mich.	12 64	35 95	1 17	18 21	31	16	23 18	32 17
Wis.	13	17	6	2	483	935	6	12
W.N. CENTRAL	55	41	23	9	500	313	34	48
Minn. Iowa	16 3	6 4	6 7	2	422	242 37	11 5	18
Mo.	21	19	4	3	15	23	14	15
N. Dak. S. Dak.	2 10	2	2	_	_	1	_	3
Nebr.	1	2	1	3	_	7	1	2
	2	C	3		1 007	1 002	3	0
Del.	250 12	209	N	/4 N	406	174	192	6
Md.	75	59	14	10	699	651	73	46
Va.	30	31	7	13	113	99	9 17	32
W.Va.	11	7	3	2	7	16	1	
S.C.	21	25 8	4	5	40	16	5	14
Ga.	18	35	17	12	4	12	27	47
ES CENTRAI	50	85 70	20	20	40	35	18	27
Ky.	15	26	3	4	4	13	4	4
Tenn. Ala	22 10	29 12	8 7	10 4	25	18 4	10 4	7
Miss.	3	3	3	2	_	—	—	5
W.S. CENTRAL	24	104	23	31	46	37	49	109
Ark. La	4	7	7	3	4	8	4	7
Okla.	3	3	3				3	7
	13	94	13	26	38	27	40	90
MountAin Mont.	5	63 1	8	17	14	15	35	38
Idaho	3	7	—	1	1	5	_	1
Colo.	3 17	5 15	3	8	3	3	18	14
N. Mex.	2	4	3	—	1	_	2	2
Utah	11	16	_	1	2	1	6	6
Nev.	7	4	2	7	1	—	2	5
PACIFIC Wash	46	47 8	92 7	94 8	78	60 8	140	115
Oreg.	N	N	6	5	15	21	7	14
Calif. Alaska	45	39	79	78	54 3	29	106	87
Hawaii	1	_	_	3	Ň	Ň	13	3
Guam	—	_	—	—				_
P.R. VI	_	_	_	_	<u>N</u>	N	2	_
Amer. Samoa	U	U	U	U	U	U	U	U
C.N.M.I.	—	U	—	U	—	U	—	U

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 10, 2005, and September 11, 2004

N: Not notifiable. U: Unavailable. —: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Incidence data for reporting years 2004 and 2005 are provisional and cumulative (year-to-date).

(John Week)	Meningococcal disease										
	All sero	aroups	Sero	group and W-135	Serog	roup B	Other se	erogroup	Serogroup unknown		
Reporting area	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004	Cum. 2005	Cum. 2004	
UNITED STATES	864	875	63	70	44	35	_	1	757	769	
NEW ENGLAND	60	52	1	5	_	6	_	1	59	40	
Maine	2	9	_	_	_	1	—	—	2	8	
N.H.	10	4	_	_	—	_	_	_	10	4	
VI. Mass	6 28	2	_	5	_		_	_	6 28	2	
R.I.	20	1	_		_		_	_	20	1	
Conn.	12	6	1	—		_	—	1	11	5	
MID. ATLANTIC	116	123	31	35	5	5	_	_	80	83	
Upstate N.Y.	30	33	4	5	3	3	—		23	25	
N.Y. City	17	22	—	—			—	—	17	22	
N.J. Pa	30	25	27	30			_	_	30	25 11	
	03	40	27	50	2	2		_	10	11	
E.N. CENTRAL	87	97	18	22	9	6	_	_	60 26	69 40	
Ind	16	15	_	1	4	1	_	_	12	13	
III.	12	1	_	_	_	_	_	_	12	1	
Mich.	18	18	18	18	_		—	_			
Wis.	10	15	—	—			—		10	15	
W.N. CENTRAL	56	61	2	_	1	4	—	_	53	57	
Minn.	11	18	1	—			—	_	10	18	
Iowa Mo	13	13	1	_	_	2	_	_	12	16	
N. Dak.		2	_	_	_	_	_	_		2	
S. Dak.	2	2	—	—		1	_	_	2	1	
Nebr.	4	4	_	—			_	_	4	4	
Kans.	8	5	_	_	—	_	_	_	8	5	
S. ATLANTIC	165	163	4	2	9	2	—		152	159	
Del.	3	3		—			—		3	3	
	10	5		2		_	_	_		3	
Va.	21	12	_	_	_	_	_	_	21	12	
W. Va.	6	5	1	_	_		—	_	5	5	
N.C.	27	26	1	—	7	2	—	_	19	24	
5.0. Ga	14	13	_	_	_	_	_	_	14	13	
Fla.	61	80	_	_	_	_	_	_	61	80	
E S CENTRAL	42	44	1	1	3	1	_		38	42	
Ky.	14	8	_	1	3	1	_	_	11	6	
Tenn.	18	14	_	_	_	_	_	_	18	14	
Ala.	6	11	1	—	—	—	—	—	5	11	
IVIISS.	4	11	_						4	11	
W.S. CENTRAL	72	50	1	1	5	1	_	_	66	48	
Ark.	11	13	_	1		_	_	_	11	13	
Okla.	12	7	1	_	3	1	_	_	8	6	
Tex.	24	3	—	—			—	_	24	3	
MOUNTAIN	69	51	4	1	5	5	_	_	60	45	
Mont.	_	3	_	_	_		_	_	_	3	
Idaho	2	6	—	—			—	—	2	6	
Wyo.	16	3		—			_		12	3	
N. Mex.	2	6		1	_	3	_	_	2	2	
Ariz.	35	10	_	_	2	1	_	_	33	9	
Utah	9	4	1	—	2		—	_	6	4	
Nev.	5	7	—	—	1	1	—		4	6	
PACIFIC	197	234	1	3	7	5	—	—	189	226	
vvash.	38	21	1	3	4	5	—	_	33	13	
Calif	∠o 119	40 158	_	_	_	_	_	_	∠o 119	40 158	
Alaska	1	4	_	_	_	_	_	_	1	4	
Hawaii	11	5	_	_	3	_	_	_	8	5	
Guam	_	_	_	_	_	_		_	_	_	
P.R.	6	13	—	—	_	—	—	—	6	13	
V.I. Amor Sames	_	-	—	—	—	—	_	—	_	-	
C.N.M.I.			_	_	_	_	_	_		_	

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending September 10, 2005, and September 11, 2004 (36th Week)*

N: Not notifiable. U: Unavailable. —: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Incidence data for reporting years 2004 and 2005 are provisional and cumulative (year-to-date).

<u>,</u> ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Per	nesis	Babies	animal	Rocky M	lountain d fever	Salmo	nellosis	Shige	llosis
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
Reporting area	2005	2004	2005	2004	2005	2004	2005	2004	2005	2004
UNITED STATES NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn	13,039 755 17 41 73 571 21	11,776 1,226 5 42 60 1,058 16	3,817 516 40 11 40 272 15	4,589 440 39 19 17 183 31	1,064 3 N 1 1 1	1,002 12 N — 10 1	25,540 1,520 105 130 79 803 74 200	28,001 1,488 77 103 39 875 75 210	8,339 207 8 6 14 129 12 29	8,987 210 5 6 2 138 13
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	914 355 57 154 348	43 1,958 1,385 135 134 304	669 374 20 N 275	680 369 11 N 300		58 1 20 10 27	3,191 832 690 532 1,137	4,176 847 952 800 1,577	822 196 258 214 154	881 347 283 173 78
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	2,483 810 208 494 165 806	3,908 387 75 768 142 2,536	152 56 29 17 31 19	132 53 7 36 31 5	33 26 2 1 4	29 8 5 12 2 2	3,451 939 383 963 613 553	3,698 892 351 1,187 592 676	582 76 105 128 162 111	785 120 134 313 82 136
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	2,021 868 372 305 81 1 152 242	1,235 157 104 264 626 22 11 51	335 55 94 59 21 43 — 63	470 59 79 42 49 80 81 80	134 2 3 111 5 4 9	99 1 82 4 12 —	1,660 389 262 529 24 106 99 251	1,685 411 341 452 30 75 113 263	995 62 56 663 2 25 43 144	303 45 56 117 3 9 19 54
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	890 5 119 7 237 36 64 253 27 142	465 — 87 107 17 62 83 17 85	1,147 214 378 39 356 5 151 4	1,622 9 225 348 50 439 111 235 205	528 2 62 35 5 329 32 48 13	474 4 47 17 4 269 49 69 15	6,936 560 39 615 104 1,005 731 1,029 2,797	7,208 78 604 43 789 172 912 729 1,315 2,566	1,335 8 60 8 75 — 133 61 312 678	2,127 6 103 30 106 5 225 457 455 740
E.S. CENTRAL Ky. Tenn. Ala. Miss.	373 106 167 65 35	221 51 134 23 13	103 7 36 58 2	102 18 34 41 9	192 2 144 42 4	148 2 81 39 26	1,763 309 496 482 476	1,782 240 487 459 596	915 215 449 190 61	579 53 293 189 44
W.S. CENTRAL Ark. La. Okla. Tex.	863 203 30 	514 51 13 17 433	614 26 — 61 527	845 41 1 87 716	67 44 5 7 11	159 83 5 70 1	2,058 492 458 274 834	2,669 362 615 284 1,408	1,759 47 83 488 1,141	2,373 51 221 330 1,771
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	2,801 500 94 33 917 107 752 370 28	944 32 25 16 466 124 155 113 13	168 12 14 14 6 105 12 5	151 20 3 4 38 4 76 3 3 3	27 1 2 5 1 13 4	19 3 4 4 2 2 1	1,565 63 70 63 437 159 458 239 76	1,624 130 121 42 407 195 457 157 115	483 5 2 81 56 277 34 26	552 4 9 4 117 94 268 27 29
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	1,939 569 517 694 59 100	1,305 447 324 506 11 17	113 U 4 108 1 —	147 U 6 130 11	8 1 7 	4 2 2 —	3,396 359 264 2,531 39 203	3,671 352 335 2,685 41 258	1,241 72 90 1,047 7 25	1,177 76 56 998 6 41
Guam P.R. V.I. Amer. Samoa C.N.M.I.	5 U	3 	52 — U	40 — U U	N U	N U U	323 	48 292 — U U	2 U	41 22 — U U

 TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 10, 2005, and September 11, 2004

 (36th Week)*

N: Not notifiable. U: Unavailable. —: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Incidence data for reporting years 2004 and 2005 are provisional and cumulative (year-to-date).

			Strepto	coccus pneum	oniae, invasiv	e disease					
	Streptococ	cal disease,	Drug res	sistant,			Brimany &	Syp	nilis Congonital		
	Cum.	Cum.	Cum.	ges Cum.	Age <5 Cum.	years Cum.	Cum.	Cum.	Cum.	Cum.	
Reporting area	3 153	<u>1 2004 </u> 3 346	2005 1.615	1 58/	618	<u>2004</u> 5/3	5 338	5 309	172	2004	
NEW ENGLAND	120	223	82	1,564	47	75	145	141	1	274	
Maine	9	9	N	N	— —	4	1	2		_	
N.H.	13	15		_	4	N	12	3		3	
Vt. Mass.	9 81	8 101	59	26	4 39	41	92	87	_	_	
R.I.	8	17	13	14		6	8	19	_	1	
Conn.	—	73	U	55	U	23	31	30	1	_	
MID. ATLANTIC	699	573	155	114	108	80	695	690	20	27	
N.Y. City	126	90	60 U	48 U	48 19	55 U	427	418	5	12	
N.J.	153	122	N	N	19	7	96	111	10	13	
Pa.	213	173	95	66	22	18	112	95		1	
E.N. CENTRAL	625	775	434	356	159	128	533	622	25	36	
Ind.	81	78	148	107	42	26	43	44	1	2	
III.	116	208	10		48	1	262	258	9	9	
Mich. Wis	243	237 71	N	N	7	N 41	56 23	133	11	23	
WN CENTRAL	205	238	33	17	66	73	170	120	- 1	З	
Minn.	79	119		<u> </u>	40	49	45	17	_	1	
Iowa	N	N	N	N	_	N	2	5			
Mo. N Dak	52	52 10	27	12	2	10	102	<u>72</u>	1	1	
S. Dak.	19	12	3	5			1	_	_	_	
Nebr.	14	15	2		6	6	4	6	_	-	
CATLANTIC	52	30	N 640	010	11	0	1041	20		1	
Del.	1	3	042	4	63	37 N	1,341	1,322	30	40	
Md.	144	102		_	41	25	229	252	10	7	
D.C. Va	7	7	15 N	8 N	2	4 N	72	41		1	
W. Va.	21	20	95	90	20	8	3	3			
N.C.	96	95	N	N	U	U	189	133	8	8	
S.C. Ga	24 126	50 161	112	79 201	_	N	42 220	233	3	10	
Fla.	172	166	419	436	—	N	490	497	5	14	
E.S. CENTRAL	128	172	125	111	7	12	294	284	16	19	
Ky. Tann	27	51	24	22	N	N	31	30		1	
Ala.		121	101		_	N	92	126	3	9	
Miss.	_	_	—	2	7	12	28	40	1	2	
W.S. CENTRAL	195	261	94	46	122	108	864	815	50	54	
Ark. La	14	16	12 82	6 40	13	23	33 176	37 194	6	3	
Okla.	87	49	N	Ň	19	32	29	19	1	2	
Tex.	88	194	N	N	68	46	626	565	43	46	
MOUNTAIN	459	364	50	20	38	30	270	276	15	35	
Mont. Idaho	1	8	N	N	_	N	5 20	15	1	2	
Wyo.	3	7	21	8	_	_		1	_	_	
Colo.	173	75	N	N	37	30	29	48			
Ariz.	183	164	N	N	_	N	104	119	12	30	
Utah	61	30	28	10	1	_	5	7	—	1	
Nev.	1	2	1	2	_	_	73	21		_	
PACIFIC Wash	71 N	77 N	N	1 N	8 N	N	1,026	1,039 83	14	50	
Oreg.	N	N	N	N	6	N	19	22	_	_	
Calif.	—	—	N	N	N	N	901	929	14	50	
Hawaii	71	77	_	1	2	IN	o 4	5	_	_	
Guam	_	_	_	_	_	_	_	1	_	_	
P.R.	Ν	N	Ν	Ν	—	Ν	141	95	8	3	
V.I. Amer Samoa								4			
C.N.M.I.	<u> </u>	ŭ	_	ŭ	_	ŭ	_	Ŭ	_	Ŭ	

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending September 10, 2005, and September 11, 2004 (36th Week)*

N: Not notifiable. U: Unavailable. —: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Incidence data for reporting years 2004 and 2005 are provisional and cumulative (year-to-date).

(Jour Week)					1						
					Vari	cella	, ,	West Nile viru	s disease [†]		
	Tube	uberculosis Typhoid fever ((chick	(chickenpox)		nvasive	Non-neuroinvasive [§]			
Reporting area	2005	2004	2005	2004	2005	2004	2005	2004	Cum. 2005		
UNITED STATES	7,336	8,876	157	222	16,390	19,598	497	964	725		
NEW ENGLAND	227	299	18	17	990	2.065	2	_	_		
Maine	10	13	1	_	213	181	_	_	—		
N.H.	4	10	—	—	203	412	—	—	_		
Mass.	141	172	10	14	538	177	_	_	_		
R.I.	18	40	1	1			_	—	_		
	50	62	6	2	U	1,294	2	_	_		
MID. AI LAN I IC Unstate N Y	1,338	1,386	32	55 7	3,161	73	8	9	6		
N.Y. City	651	697	10	20	_	_	2	2	1		
N.J.	322	299	9	16	2 161			1	 F		
	194	199	0	12	3,101	73	0	5	5		
Ohio	895 171	808 141	12	28	4,501	8,454 1.048	90 10	53	46		
Ind.	88	86			482	N	1	6			
III. Mich	435	360	3	12	64	4,328	71	23	42		
Wis.	58	62	4	2	2,034	502	3	4	1		
W.N. CENTRAL	300	309	2	7	301	136	62	69	216		
Minn.	128	114	2	3			7	10	13		
Iowa Mo	32 68	26 82	_	2	N 210	N 5	2	23	3		
N. Dak.	2	3	_		13	75	2	20	14		
S. Dak.	9	8	—		78	56	28	5	140		
Kans.	39	23 53	_	2	_	_	18	4 16	39		
S. ATLANTIC	1.634	1.835	26	31	1.397	1.734	8	53	9		
Del.	7	17			21	4		_	_		
Md.	184 38	185	9	11	24	20	1	7	_		
Va.	214	148	5	5	284	411	_	3	_		
W.Va.	17	14			716	976			N		
S.C.	147	214 131		3	352	323	1	- 3			
Ga.	254	409	2	4	_	_	1	11	1		
Fla.	588	651	8	8	—	—	4	28	7		
E.S. CENTRAL	362	434	5	6		34	18	53	14		
Tenn.	161	146		4			2	9	1		
Ala.	129	134	1	_	_	34	3	15	2		
MISS.	_	82	2		_	_	12	28	11		
W.S. CENTRAL	776 70	1,366	9	20	4,245	5,464	82 1	175	27		
La.			_	_	107	48	50	57	16		
Okla.	92 614	112		1	4 129	 5 /16	1	10	6		
	014	255	9	19	4,136	1 629	50	201	95		
Mont.	250	4	<u> </u>	<u> </u>	1,795	1,030	5	1	6		
Idaho	—	3	—	_			2	1	5		
vvyo. Colo	46	2 86	3	1	45 1 268	26 1.300	5	2 39			
N. Mex.	14	21	_	_	123	U	10	28	4		
Ariz.	149	146	3	2	250	212	14	203	10		
Nev.	18	65	1	2			6	22	11		
PACIFIC	1.548	2.084	45	52	_	_	175	251	322		
Wash.	172	151	5	4	Ν	Ν	_		_		
Oreg. Calif	54 1 227	69 1 751	2 31	1 41	_	_	175	251	3 319		
Alaska	18	27	_	—	_	_			_		
Hawaii	77	86	7	6	_	—	—	_	—		
Guam	_	41	—	—		108	_	_	_		
Р.К. V.I.	_	/4	_	_	499	300	_	_	_		
Amer. Samoa	U	U	U	U	U	U	U	U	—		
C.N.M.I.		U		U		U	_	U	_		

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending September 10, 2005, and September 11, 2004 (36th Week)*

N: Not notifiable. U: Unavailable. —: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Incidence data for reporting years 2004 and 2005 are provisional and cumulative (year-to-date). † Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Infectious Diseases (ArboNet Surveillance). * Not previously notifiable.

TABLE III. Deaths in 122 U.S. cities,* week ending September 10, 2005 (36th Week)

	All causes, by age (years)							All causes, by age (years)							
Reporting Area	All Ages	<u>≥</u> 65	45-64	25–44	1–24	<1	P&l⁺ Total	Reporting Area	All Ages	<u>≥</u> 65	45-64	25–44	1–24	<1	P&I [†] Total
NEW ENGLAND	430	296	88	30	6	10	43	S. ATLANTIC	1,099	669	270	106	27	27	60
Boston, Mass.	115	68	23	15	2	7	16	Atlanta, Ga.	103	65	26	8	3	1	6
Bridgeport, Conn.	22	20	2				2	Baltimore, Md.	159	83	51	18	5	2	11
Cambridge, Mass.	11	/	2	1	1	_	_	Charlotte, N.C.	101	68	1/	11	3	2	9
Hartford Conn	51	21	5 15	3	1	1	6	Miami Ela	138	92 81	27	12	4	3	0
Lowell Mass	13	11	1	1	_	_	3	Norfolk Va	36	18	10	2	1	5	1
Lvnn. Mass.	10	5	3	2	_	_	2	Richmond, Va.	59	31	12	3	4	9	4
New Bedford, Mass.	15	12	2	1	_	_	_	Savannah, Ga.	37	26	5	5	1	_	5
New Haven, Conn.	U	U	U	U	U	U	U	St. Petersburg, Fla.	40	31	8	1	—	_	1
Providence, R.I.	53	44	5	4	_	_	3	Tampa, Fla.	164	110	37	15	1	1	6
Somerville, Mass.	4	2	2	_	_	_	_	Washington, D.C.	100	55	31	10	_	4	1
Springfield, Mass.	28	17	9	1	1		2	vviimington, Del.	18	9	5	4	_	_	1
Worcester Mass	58	46	9 10	_	_	2	6	E.S. CENTRAL	669	423	154	57	15	20	36
			10					Birmingham, Ala.	124	73	29	8	2	12	7
MID. AI LANTIC	1,891	1,305	389	136	30	31	84	Chattanooga, Ienn.	57	34	20	1	_	2	4
Albany, N.Y.	56	42	9	3	_	2	3	Knoxville, lenn.	89	64	15	8	2	-	1
Ruffalo N Y	110	9 79	21	7	1	2	8	Memphis Tenn	135	45 79	33	18	3	1	6
Camden, N.J.	18	10	6	1	_	1	2	Mobile, Ala.	44	26	12	2	2	2	2
Elizabeth, N.J.	17	14	1	2	_	_	3	Montgomery, Ala.	40	33	5	2	_	_	4
Erie, Pa.	48	37	10	—	_	1	4	Nashville, Tenn.	115	69	29	13	2	2	6
Jersey City, N.J.	35	19	10	5	1	—	_	W.S. CENTRAL	1 015	660	213	83	30	29	57
New York City, N.Y.	910	623	186	70	13	18	34	Austin, Tex.	46	28	10	5	2	1	2
Newark, N.J.	4/	19	17	5	2	4	_	Baton Rouge, La.	15	12	2	1	_	_	_
Paterson, N.J. Philadelphia Pa	250	14	4 57	26	6	_	6	Corpus Christi, Tex.	31	22	4	1	1	3	1
Pittsburgh Pa §	233	15	7	20	_	_	1	Dallas, Tex.	137	88	24	16	5	4	3
Reading, Pa.	24	21	3	_	_	_	_	El Paso, Tex.	46	33	13		—		5
Rochester, N.Y.	138	104	25	3	3	3	13	Ft. Worth, Tex.	79	47	22	5	10	5	4
Schenectady, N.Y.	26	18	6	1	1	_	_	Little Bock Ark	209	101	13	21	12	3	22
Scranton, Pa.	20	16	3	1		_	2	New Orleans La ¹	U	U 37	U	Ű	ú	Ŭ	U
Syracuse, N.Y.	67	55	8	3	1		7	San Antonio, Tex.	189	129	38	14	4	4	10
Irenton, N.J.	26	16	6	3	1		_	Shreveport, La.	55	40	10	5	_	_	5
Yonkers NY	22	10	3	3	1	_	_	Tulsa, Ökla.	91	63	15	7	4	2	5
	1 005	1 1 0 0		100		0.4	104	MOUNTAIN	704	455	145	64	21	19	46
Akron Ohio	1,085	30	363	129	40 4	24 3	104	Albuquerque, N.M.	105	63	23	15	3	1	7
Canton, Ohio	24	17	6	1	_	_	4	Boise, Idaho	45	36	6	3	_		6
Chicago, III.	277	166	69	35	5	2	18	Colo. Springs, Colo.	50	41	7	1	_	1	1
Cincinnati, Ohio	46	31	6	6	3	_	3	Denver, Colo.	80	45	19	22	3	6	5 16
Cleveland, Ohio	182	132	37	10	2	1	12	Orden Utah	31	24	40	23	_	-	3
Columbus, Ohio	192	114	58	13	5	2	13	Phoenix, Ariz.	56	33	11	5	5	2	2
Dayton, Onio	114	82	22	4	3	3	6	Pueblo, Colo.	17	14	3	_	_	_	2
Evansville Ind	46	70 39	5	9	1		5	Salt Lake City, Utah	93	53	22	9	4	5	4
Fort Wayne, Ind.	40	29	10	1		_	3	Tucson, Ariz.	U	U	U	U	U	U	U
Gary, Ind.	23	12	6	2	2	1	2	PACIFIC	1,299	928	250	75	27	19	102
Grand Rapids, Mich.	39	26	5	4	1	3	6	Berkeley, Calif.	14	9	4	1	—	—	2
Indianapolis, Ind.	134	84	37	9	2	2	2	Fresno, Calif.	78	58	14	5	1	_	5
Lansing, Mich.	43	37	5	1	_	_	_	Glendale, Calif.	10	6	3	1	_	_	1
Milwaukee, Wis.	85	54	14	12	2	3	8	Honolulu, Hawali	70	52	12	3	1	2	6
Rockford III	63	25	13	4 5	1	_	1	Long Beach, Calif.	40 108	1/5	12	11	5		33
South Bend Ind	45	34	8	1	1	1	_	Pasadena Calif	22	16	4	1		1	3
Toledo, Ohio	74	56	11	4	2	1	6	Portland, Oreg.	103	74	22	5	1	1	4
Youngstown, Ohio	35	27	5	3	_	—	2	Sacramento, Calif.	167	112	42	10	3	_	6
WN CENTRAL	449	307	93	21	15	13	32	San Diego, Calif.	120	86	20	8	2	4	7
Des Moines. Iowa	73	50	19	1	_	3	4	San Francisco, Calif.	81	54	17	7	2	1	2
Duluth, Minn.	31	23	5	_	2	1	2	San Jose, Calif.	149	112	28	6	3	_	12
Kansas City, Kans.	19	14	2	3	_	_	1	Santa Uruz, Calif.	102	10	10			~	10
Kansas City, Mo.	72	42	17	3	6	4	5	Spokane Wash	41	34	01 A	0	5	ა 1	12
Lincoln, Nebr.	45	37	6	2	_	<u> </u>	7	Tacoma, Wash.	82	58	14	6	4	_	1
Minneapolis, Minn.	43	28	12	2	_	1	6	TOTAL	0.044++	0.150	1 005	707		100	
Omana, Nebr.	62	36	17	6	2	1	5	TOTAL	9,241**	6,152	1,985	701	211	192	564
St. LOUIS, IVIO.	16	1 27	6	1	1	1	_								
Wichita, Kans.	57	39	9	3	4	2	2								

U: Unavailable. —: No reported cases.

*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. [¶]Because of Hurricane Katrina, weekly reporting of deaths has been temporarily disrupted.

** Total includes unknown ages.

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 K-95, 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.

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.

☆U.S. Government Printing Office: 2005-733-116/00112 Region IV ISSN: 0149-2195