

Weekly

July 19, 2002 / Vol. 51 / No. 28

### Tetanus — Puerto Rico, 2002

During February–May 2002, the Puerto Rico Department of Health (PRDOH) received reports of three tetanus cases, two of which were fatal. The last reported case of tetanus in Puerto Rico had occurred in 1999. This report summarizes the investigations of these three cases, which underscore that health-care providers should ensure that all patients have been vaccinated fully against tetanus (1,2).

### **Case Reports**

**Case 1.** On December 19, 2001, a man aged 86 years with a history of hypertension and coronary artery disease (CAD) sustained a splinter in his right hand while gardening. On December 22, the patient saw a physician for wound care. At that time, he was not treated with either a tetanus toxoid vaccine or prophylactic tetanus immune globulin (TIG). His tetanus vaccination history was not documented in the medical record; he had no history of military service.

On December 26, the patient received treatment for pharyngitis from a local physician. On December 29, he presented to an emergency department (ED) with difficulty talking, swallowing, and breathing and with chest pain and disorientation of 2 days' duration. He was admitted to a general medicine ward with a preliminary diagnosis of stroke.

On January 2, 2002, the patient had neck rigidity and respiratory failure requiring tracheotomy and mechanical ventilation and was transferred to the intensive care unit (ICU) with tetanus diagnosed. He was administered a dose of tetanus and diphtheria toxoids (Td); TIG was ordered but was unavailable. On January 11, the patient received nonspecific intravenous immune globulin (pooled plasma, 7.5 grams). His hospital course was complicated by two myocardial infarctions, congestive heart failure, a lacunar stroke, and pneumonia. He died on February 2. **Case 2.** On April 18, 2002, a man aged 68 years with a history of diabetes mellitus, CAD, and mitral valve replacement sustained a puncture wound in his right foot from stepping on a rusted nail. His spouse cleaned the wound with a surface antiseptic (benzalkonium chloride). The following day, the patient sought care from a primary-care physician who administered intravenous cefazolin and prescribed oral ciprofloxacin and oxycodone. The patient requested vaccination against tetanus but was told that the vaccine was unavailable. The patient did not know if he had been vaccinated previously against tetanus; he had not served in the military.

On April 22, the patient presented to an ED complaining of difficulty swallowing, mild shortness of breath, abdominal pain, throat pain, and mandibular rigidity. On physical examination, he had trismus, risus sardonicus, muscular rigidity, and difficulty speaking. He was admitted to the ICU with diagnoses of suspected tetanus and right foot cellulitis. He was treated with metronidazole, ciprofloxacin, and midazolam by continuous intravenous infusion. On April 23, the patient had seizures and respiratory failure requiring mechanical ventilation. He also was given intramuscular TIG (500 units) and Td (0.5 cc) at that time. Despite midazolam therapy and supplemental diazepam for seizures, the patient's muscle spasms persisted. He died on April 27.

### INSIDE

- 616 Pertussis Deaths United States, 2000
- 618 Hepatitis B Vaccination Among High-Risk Adolescents and Adults — San Diego, California, 1998–2001
- 621 Weekly Update: West Nile Virus Activity United States, July 10–16, 2002
- 622 Poliomyelitis Madagascar, 2002

CENTERS FOR DISEASE CONTROL AND PREVENTION SAFER • HEALTHIER • PEOPLE™ 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 **Case 3.** On April 10, 2002, a man aged 76 years with a history of hypertension sustained a splinter in his right hand. On April 18, the patient experienced weakness and dysphagia, and on the following day, trismus. At that time, he was treated for otitis media but refused Td vaccination. His previous tetanus vaccination status was unknown; he had not served in the military.

On April 20, the patient presented to an ED with difficulty walking, talking, and swallowing. He did not report any wound history to the attending physician. He was treated with an intramuscular corticosteroid injection and an antihistamine. On April 21, the patient sought care at another ED. He was admitted to the ICU with diagnosed tetanus and intubated preemptively. On April 22, he received 3,000 units of TIG and was started on metronidazole. His course was complicated by methicillin-sensitive *Staphylococcus aureus* pneumonia and pseudomembranous colitis. He was released from the hospital on June 17.

### **Case Summary**

During January 1990–April 2002, PRDOH received reports of 20 cases of tetanus (average annual incidence rate: 0.04 per 100,000 population). Of these, 18 (90%) were in men; the median age was 70 years (range: 55–86 years). Among the 11 (55%) for whom supplemental information was available, none had a definite history of previous vaccination with tetanus toxoid. Five (25%) patients had a history of diabetes mellitus. The overall case-fatality rate was 68%.

As a result of the Td shortage affecting the United States during 2000–2002, PRDOH instituted a protocol in March 2001 consistent with the modified guidelines for Td use during the shortage (3,4). Priority was given to persons requiring prophylaxis for wound management and to persons who had previously received fewer than 3 doses of tetanus-containing vaccine, and routine Td boosters in adolescents and adults were deferred. The shortage reduced Td use in Puerto Rico by 67% during 2000–2001 (Puerto Rico Immunization Program, unpublished data, 2002).

In response to the recent tetanus cases, PRDOH has 1) continued reminding health-care providers of the increased risk for tetanus among persons aged  $\geq 60$  years and those with no history of primary vaccination against tetanus; 2) promoted an increase in the availability of TIG for prophylactic and therapeutic use; and 3) notified physicians that the Td shortage has ended and that Td is available for routine indications (5). **Reported by:** JC Orengo, MD, Y García, MPH, A Rodríguez, MD, J Rullán, MD, Puerto Rico Dept of Health. MH Roper, MD, P Srivastava, MS, TV Murphy, MD, Epidemiology and Surveillance Div, National Immunization Program; F Alvarado-Ramy, MD, Div of Applied Public Health Training, Epidemiology Program Office, CDC.

**Editorial Note:** Tetanus is a rare disease in the United States; following the introduction of vaccination with tetanus toxoid in the 1940s, the overall incidence of tetanus declined from 0.4 per 100,000 population in 1947 to 0.02 during the latter half of the 1990s. The overall case-fatality ratio declined from 91% to 11% during the same period. The majority of tetanus cases reported during 1989–1997 occurred in persons who had not completed a 3-dose primary tetanus toxoid vaccination series or for whom vaccination histories were uncertain; no tetanus deaths occurred in persons who received primary tetanus vaccination (5–7; CDC, unpublished data, 2002).

Adults aged  $\geq 60$  years are at greatest risk for tetanus and tetanus-related mortality (5–7). During 1998–2000, the average annual incidence of tetanus in persons aged  $\geq 60$  years was 0.03 with a case-fatality ratio of 31%, both more than twice that of adults aged <60 years. The increased risk for tetanus with increasing age is thought to be related to the lower prevalence of protective immunity in older age groups. Protective levels of antibodies against tetanus toxoid decline with age; by age 70 years, only 30% of the population is protected (8). Older persons might never have received a primary vaccination series or might not have received subsequent Td boosters. Women are significantly less likely to be protected against tetanus than men (8) probably, in part, because women are less likely to have received a Td booster in conjunction with military service.

The Td shortage during 2000–2002 necessitated deferral of routine Td boosters in adolescents and adults. However, booster doses given as part of wound management and administration of primary series in unvaccinated persons remained priorities (3). Previous reports on tetanus cases occurring in the United States during the 1980s and 1990s indicated that even during periods in which Td was in ample supply, <60% of persons for whom Td was indicated received a dose during wound management (5–7).

Recommendations for the use of Td and TIG for wound care depend on the nature of the wound and the patient's vaccination history. Persons who have received a primary tetanus vaccination series but who have not had a Td booster during the 10 years preceding any injury should receive a booster dose. Persons who present with wounds contaminated with dirt, feces, or saliva, deep wounds, or wounds with necrotic tissue and who have not had a booster during the preceding 5 years also should receive a dose of Td. Persons who have never received tetanus vaccination or those with unknown or uncertain vaccination histories should receive the first dose of a primary series at the time of presentation. These patients also should receive TIG (250 units injected intramuscularly at a site distant from that used for Td administration) unless the wound is superficial and clean, because a single dose of Td in the absence of previous tetanus vaccination will not induce the production of protective levels of antibody. Therapeutic TIG (3,000–6,000 units as 1 dose) should be administered as soon as possible to any patient presenting with tetanus (9).

The majority of cases of tetanus and virtually all tetanusassociated deaths are preventable through adequate vaccination. Because all wounds, even minor and relatively clean wounds, confer a risk for tetanus, health-care providers should review the vaccination status of all patients and administer indicated tetanus toxoid vaccine to keep their patients fully protected (1,2).

#### References

- CDC. Diphtheria, tetanus, and pertussis: recommendations for vaccine use and other preventive measures—recommendations of the Immunization Practices Advisory Committee (ACIP). MMWR 1991;40(No. RR-10).
- CDC. Immunization of adolescents: recommendations of the Advisory Committee on Immunization Practices, the American Academy of Pediatrics, the American Academy of Family Physicians, and the American Medical Association. MMWR 1996;45(No. RR-13).
- 3. CDC. Shortage of tetanus and diphtheria toxoids. MMWR 2000;49:1029-30.
- CDC. Deferral of routine booster doses of tetanus and diphtheria toxoids for adolescents and adults. MMWR 2001;50:418, 427.
- CDC. Resumption of routine schedule for tetanus and diphtheria toxoids. MMWR 2002;51:529–30.
- Prevots R, Sutter RW, Strebel PM, Cochi SL, Hadler S. Tetanus surveillance—United States, 1989–1990. In: CDC Surveillance Summaries (December 11). MMWR 1992;41(No. SS-8).
- Izurieta HS, Sutter RW, Strebel PM, et al. Tetanus surveillance—United States, 1991–1994. In: CDC Surveillance Summaries (February 21). MMWR 1997;46(No. SS-2).
- Bardenheier B, Prevots R, Khetsuriani N, Wharton M. Tetanus surveillance—United States, 1995–1997. In: CDC Surveillance Summaries (July 3). MMWR 1998;47(No. SS-2).
- 9. McQuillan G, Kruszon-Moran D, Deforrest A, Chu SY, Wharton M. Serologic immunity to diphtheria and tetanus in the United States. Ann Int Med 2002;136:660–6.
- American Academy of Pediatrics. Tetanus. In: Pickering LK, ed. 2000 Red book: report of the Committee on Infectious Diseases. 25th ed. Elk Grove Village, Illinois: American Academy of Pediatrics, 2000:563–8.

# Pertussis Deaths — United States, 2000

Pertussis (i.e., whooping cough) is associated typically with an inspiratory "whoop," prolonged paroxysmal cough, and posttussive vomiting; however, persons infected with *Bordetella pertussis* sometimes experience atypical symptoms, making prompt recognition difficult (1) and probably increasing infection transmission. All infants aged <6 months and any infants who have not yet received 3 doses of diphtheria and tetanus toxoids and acellular pertussis (DTaP) vaccine are especially vulnerable to *B. pertussis* infection (2). This report summarizes the investigations of two pertussis deaths that occurred in 2000. Clinicians should consider pertussis as a cause of illness, especially among vulnerable infants who present with cough illness, respiratory distress, or apnea. Timely diagnosis of pertussis in caregivers and other contacts of infants could prevent infant pertussis fatalities.

### **Case Reports**

Colorado. On January 6, 2000, a full-term, white, non-Hispanic female infant aged 3 months was evaluated by her pediatrician for rhinorrhea and cough of 7 days' duration. A test for respiratory syncytial virus (RSV) was negative, and the infant received her first vaccinations, including DTaP vaccine. On January 17, the infant returned with persistent symptoms that had progressed during the preceding 2-3 days to include paroxysmal cough, breathing difficulty, and fever. Perioral cyanosis, intercostal retractions, tachypnea, and hypoxia were noted. A chest radiograph revealed marked hyperinflation and bilateral perihilar infiltrates. The infant's mother reported a cough illness with onset 3-4 weeks before the infant's cough onset; the infant's sibling aged 3 years (who had received 4 DTaP vaccinations) also had a mild cough illness. On hospital admission that day, the infant's leukocyte count was 129,000 (normal: 5,000-20,000). Specimens of nasopharyngeal (NP) secretions were collected for *B. pertussis* culture and repeat RSV testing. A blood sample was obtained for culture, and empiric treatment for pertussis was initiated with oral azithromycin, which was later replaced with oral erythromycin. On January 18, the infant became increasingly irritable, had a temperature of 104° F (40° C), and was transferred to a tertiary medical center. Pertussis complicated by bacterial pneumonia was diagnosed presumptively and the infant was treated with intravenous erythromycin, nafcillin, and cefotaxime. NP specimens were tested by polymerase chain reaction (PCR) assay for B. pertussis DNA; a positive assay result was reported on January 20. Recurrent apnea was followed on January 22 by acute respiratory decompensation,

requiring mechanical ventilation. Management of disseminated intravascular coagulation, hypotension, hyponatremia, and hypoalbuminemia was necessary. On January 24, the infant's antibiotic regimen was augmented empirically with ceftazidime and tobramycin, and a tracheal aspirate culture confirmed Pseudomonas aeruginosa infection later that day. An echocardiogram revealed severe pulmonary hypertension and right ventricular dilatation. The infant had multiple cardiac arrests, including one during initiation of extracorporeal membrane oxygenation (ECMO). On January 25, a cranial ultrasound revealed severe frontal hemorrhage; support was withdrawn, and the infant died. An autopsy confirmed that the infant died because of B. pertussis infection, superimposed P. aeruginosa sepsis, and severe necrotizing bronchopneumonia. Microscopic examination of the lung revealed necrosis, hemorrhage, and gram-negative bacilli. B. pertussis was isolated from nasopharyngeal secretions collected on January 17. A blood culture collected on January 23 and postmortem cultures from multiple sites yielded P. aeruginosa. No other pathogens were identified.

Texas. On November 10, 2000, a full-term, white, Hispanic female infant aged 3 weeks was evaluated by her pediatrician for a 3-day history of cough, posttussive emesis, and poor feeding; supportive care was recommended. That evening, the infant had worsening cough and posttussive emesis and was taken to the emergency department of hospital A. A chest radiograph revealed a right upper lobe infiltrate; the infant's leukocyte count was 8,800. A blood sample was obtained for culture. Intramuscular ceftriaxone was administered, and the patient was discharged. The next morning, because of respiratory distress and hypoxia, the infant was admitted to hospital B. A second chest radiograph revealed a right-sided infiltrate. Ampicillin, gentamicin, and vancomycin were administered empirically. The infant was intubated and transported to a tertiary care center. On her arrival at hospital C, a third chest radiograph revealed extensive bilateral infiltrates; the infant's leukocyte count was 112,000. Specimens of NP secretions were obtained to test by PCR assay for *B. pertussis* DNA. Ampicillin and cefotaxime were administered empirically. Following transfer, the maternal grandmother reported a 1-month history of severe cough; both parents reported 2 weeks of severe cough illness with posttussive emesis. The infant's cardiopulmonary status did not improve with either conventional or high-frequency oscillatory ventilation and was complicated by a right-sided pneumothorax and hypotension. An echocardiogram suggested pulmonary hypertension. Having failed to respond to inhaled nitric oxide therapy, the infant was placed on ECMO with transient stabilization on November 12. Because pathogens including *B. pertussis* and herpes simplex viruses were suspected, erythromycin, acyclovir, and clindamycin were administered empirically. Later that day, the infant had a cardiac arrest and died. An autopsy was not performed. After the infant's death, *B. pertussis* DNA was detected by PCR, and herpes simplex virus was detected by direct fluorescent antibody testing. Blood cultures from hospitals A and C, and viral cultures from hospital C, did not identify other pathogens.

### **United States**

A total of 17 deaths of persons having pertussis symptom onset in 2000 were reported to CDC by 12 states. All deaths occurred among infants born in the United States, with onset of pertussis symptoms at age <4 months. Nine (53%) deaths occurred among males. Of the 17 deceased infants, 14 (82%) were white, one (6%) was black, and one (6%) was American Indian/Alaska Native; race was not reported for one (6%). Data on ethnicity were reported for 15 (88%) infants; seven (41%) of the 17 deceased infants were Hispanic.

**Reported by:** K Plott, MPH, Association of Schools of Public Health, Atlanta, Georgia. FB Pascual, MPH, KM Bisgard, DVM, C Vitek, MD, TV Murphy, MD, Epidemiology and Surveillance Div, National Immunization Program; CR Curtis, MD, EIS Officer, CDC.

Editorial Note: Despite record high vaccination coverage levels with 3 doses of DTaP among U.S. children aged 19-35 months (3), pertussis continues to cause fatal illness among vulnerable infants. During 1980-1998, the average annual incidence of reported pertussis cases and deaths among U.S. infants increased 50% (4). The increased morbidity and mortality occurred primarily among infants aged <4 months, who were too young to have received the recommended three DTaP vaccinations at ages 2, 4, and 6 months (1,2,4). During 1990-1999, a disproportionately high number of pertussis deaths occurred among Hispanic infants; of 89 infants who died from pertussis for whom data on ethnicity were available, 31 (35%) were Hispanic (5; CDC, unpublished data, 2002). Academic investigators and public health agencies, including CDC, are initiating studies to identify the risk factors for severe and fatal pertussis.

Infants with severe pertussis often are suspected initially of having systemic infection and are treated with broadspectrum antibiotics. The two cases described in this report illustrate that pertussis can be fatal despite broad-spectrum antimicrobial therapy, specific therapy for pertussis, and supportive interventions. Severe respiratory insufficiency (caused by primary pertussis pneumonia, secondary bacterial pneumonia, or both) is the most commonly recognized immediate cause of death among infants with underlying pertussis infection (5-8). Co-infection with viral pathogens also has occurred (7).

Refractory pulmonary hypertension is associated with fatal outcomes among very young infants with pertussis (8, 9). During 2000, of the eight deceased infants for whom medical records were available, six (including the two cases in this report) received ECMO for management of pulmonary hypertension before their deaths (CDC, unpublished data, 2002). Risk factors and optimal treatment for pulmonary hypertension associated with pertussis are not defined clearly and require further investigation (9).

Adults and children with pertussis sometimes experience mild respiratory symptoms or typical symptoms (e.g., an inspiratory "whoop," prolonged paroxysmal cough, and posttussive vomiting) (6). Although some vulnerable infants exhibit these manifestations, infants with pertussis also can present with respiratory distress or apnea. Because the spectrum of symptoms among infected persons is broad, a timely diagnosis of pertussis can be difficult. Clinicians should consider pertussis as a possible cause of acute respiratory illness and apnea among vulnerable infants and as a possible cause of acute cough illness among noninfants, especially parents, siblings, and other contacts of infants. After collection of an NP specimen for *B. pertussis* culture, empiric macrolide antibiotic treatment should be initiated. Erythromycin is generally effective for *B. pertussis* treatment and chemoprophylaxis. Because published data describing the safety and efficacy of macrolides other than erythromycin are limited, erythromycin remains the preferred antibiotic for these indications (6).

Caregivers should minimize exposure of vulnerable infants to any persons with respiratory illness. As illustrated by these two cases, adult and adolescent caregivers and other family members have been linked epidemiologically as sources of pertussis infection for vulnerable infants (10). All suspected pertussis cases should be reported promptly to local public health officials, who will assist with control measures in households and communities.

Timely vaccination of infants and children according to current recommendations of the Advisory Committee on Immunization Practices remains the most effective way for infants' caregivers and health-care providers to prevent pertussis (2). Infants should receive the first DTaP vaccine at age 2 months, followed by doses at ages 4, 6, and 15–18 months and a booster dose at age 4–6 years. During a communitywide pertussis outbreak, an accelerated DTaP vaccination schedule may be used. Infants vaccinated with the accelerated DTaP vaccination schedule receive the first DTaP dose at age 6 weeks and the next 2 doses at 4-week intervals (6).

### Acknowledgments

This report is based on data contributed by state health departments to the National Notifiable Disease Surveillance System and by: S Rios, D Woods-Stout, R Hoffman, MD, State Epidemiologist, Colorado Dept of Public Health. D Bastis, MPH, L Tabony, MPH, J Pelosi, MPH, D Perrotta, PhD, State Epidemiologist, Texas Dept of Health; S Whitworth, MD, L Snow, Cook Children's Medical Center, Fort Worth, Texas.

#### References

- 1. CDC. Pertussis—United States, 1997–2000. MMWR 2002;51:73– 6.
- CDC. Pertussis vaccination: use of acellular pertussis vaccines among infants and young children. Recommendations of the Immunization Practices Advisory Committee (ACIP). MMWR 1997;46(No. RR-7).
- CDC. National, state, and urban area vaccination coverage levels among children aged 19–35 months—United States, 1999. MMWR 2000;49:585–9.
- Tanaka M, Vitek C, Pascual FB, Bisgard KM, Murphy T. Increasing incidence of pertussis among young infants in the United States, 1980– 98. In: Abstracts of the 38th Annual Meeting of the Infectious Diseases Society of America [Abstracts]. New Orleans, Louisiana: Infectious Diseases Society of America, 2000.
- Vitek C, Pascual B, Murphy T. Pertussis deaths in the United States in the 1990s. In: Abstracts of the 40th Interscience Conference on Antimicrobial Agents and Chemotherapy [Abstracts]. Washington, DC: American Society for Microbiology, 2000.
- 6. CDC. Guidelines for the control of pertussis outbreaks. Available at http://www.cdc.gov/nip/publications/pertussis/guide.htm.
- Smith C, Vyas H. Early infantile pertussis; increasingly prevalent and potentially fatal. Euro J Pediatr 2000;159:898–900.
- Goulin GD, Kaya KM, Bradley JS. Severe pulmonary hypertension associated with shock and death in infants infected with *Bordetella pertussis*. Crit Care Med 1993;21:1791–4.
- 9. Williams GD, Numa A, Sokol J, Tobias V, Duffy BJ. ECLS in pertussis: does it have a role? Intensive Care Med 1998;24:1089–92.
- Bisgard KM, Cianfrini CL, Pascual FB, et al. Infant pertussis—who is the source? Prospective investigation of cases from GA, IL, MN, MA, January 1999–October 2000. In: Abstracts of the annual meeting of the Pediatric Academic Societies [Abstracts]. Baltimore, Maryland: Pediatric Academic Societies, 2001;49:110a.

### Hepatitis B Vaccination Among High-Risk Adolescents and Adults — San Diego, California, 1998–2001

The national strategy to eliminate hepatitis B virus (HBV) transmission is based on 1) screening all pregnant women for hepatitis B surface antigen and post-exposure vaccination of infants of infected mothers; 2) vaccinating all infants as part of the childhood vaccination schedule; 3) vaccinating children and adolescents not vaccinated previously; and 4) vaccinating adolescents and adults in groups at increased risk for infection (1,2). These strategies have been implemented successfully in the United States except for the vaccination of

adults and older adolescents at high risk (2). This report describes the initial findings of a hepatitis B vaccination program for potentially high-risk adolescents and adults conducted in areas of San Diego County, California. The findings indicate that high rates of hepatitis B vaccination can be achieved in clinics and programs that serve persons at high risk for HBV infection through the integration of hepatitis B vaccination into routine preventive health-care services. Improved efforts to vaccinate adolescents and adults at increased risk for HBV infection are critical to reduce disease incidence and prevent chronic HBV infection.

The San Diego Viral Hepatitis Prevention Project (VHPP) began in February 1998 with the selection of a convenience sample of sites\* located primarily in the central and southeast areas of San Diego County, where the incidences of gonorrhea and chlamydia are higher than in other parts of the county. The population of San Diego County is approximately 2.9 million persons, and the population of the central and southeast areas is approximately 500,000 persons. Sites that serve both clients at high risk and those with a lower risk for HBV infection were selected. Hepatitis B vaccine was provided at no cost to participating sites, and project staff assisted site personnel in developing educational materials and administrative procedures and in monitoring vaccine coverage and completion. At sites that did not provide clinical services, the project provided a vaccination nurse on selected days.

At all participating sites, clinic managers/program administrators agreed to offer vaccine to all clients without collecting client-specific risk information. At most sites, clients starting vaccination were asked to complete a selfadministered sexually transmitted disease (STD)/hepatitis riskassessment form that included information about previous hepatitis B vaccination or infection. All STD clinic clients were asked to complete the risk-assessment form to determine the percentage of clients eligible to start vaccination (i.e., those with no self-reported history of previous hepatitis B vaccination or infection). Approximately 85% of STD clients were eligible to start hepatitis B vaccination; this percentage was used at other project sites to estimate the number of eligible clients. Risk criteria were not used to determine eligibility.

<sup>\*</sup>Sites serving primarily persons with a high risk for HBV infection included clinics providing treatment for sexually transmitted diseases, centers providing services for men having sex with men, the Job Corps program for disadvantaged youth, clinics providing methadone treatment for injection-drug users, drugoffender rehabilitation programs, and correctional institutions. Sites serving primarily persons with a lower risk for HBV infection included clinics providing family planning services, teen services, university/college health care, and community primary care.

### **STD Clinics**

Hepatitis B vaccination was offered to all clients of the county health department's STD clinics. During February 1998–January 2001, risk-assessment forms were completed by 18,221 clients, of whom 1,900 (10%) reported previous completion of the hepatitis B vaccination series. Among men who have sex with men (MSM) and injection-drug users (IDUs), 16% (286 of 1,755) and 6% (67 of 1,106), respectively, reported having completed the vaccination series previously; among those aged <25 years, 12% (31 of 265) of MSM and 8% (12 of 153) of IDUs reported completion of the series.

Of 18,221 clients completing risk-assessment forms, 15,502 (85%) were eligible to begin the vaccination series, of whom 11,405 (74%) received the first dose of vaccine. Of the 9,697 clients for whom  $\geq 6$  months had elapsed since they received the first dose, 5,123 (53%) received the second dose, and 2,910 (30%) completed the 3-dose series (Table).

To improve vaccination acceptance rates, during October 1999–December 2000, the main clinic offered all clients a 5-minute counseling session about hepatitis B vaccination. The acceptance rate for the first dose increased from 66% (4,390 of 6,615) during February 1998–September 1999 (before counseling was initiated) to 77% (3,094 of 4,040) during the 15-month counseling period (rate ratio [RR]=1.15; 95% confidence interval [CI]=1.13–1.18; p<0.001). Because of staff shortages and scheduling difficulties, counselors were

not available on all days; as a result, some clients were not counseled. Among the 1,861 clients counseled, the acceptance rate for the first dose was 80%, compared with 74% (1,610 of 2,189) for clients who were not counseled (RR=1.08; 95% CI=1.05–1.12; p<0.001). HIV counselors now provide hepatitis prevention and vaccination information as part of pretest HIV counseling offered to all clients.

### **Other Sites**

Other sites serving primarily clients at high-risk attained first-dose vaccination coverage rates of 4%–66%, with correctional institutions (i.e., county juvenile detention and adult jail) and a health-care clinic serving MSM having the lowest first-dose coverage rates (Table). At sites serving primarily clients at lower-risk, vaccine coverage was <30% at all sites except teen clinics, which had a first-dose coverage rate of 69%. Although community primary-care clinics vaccinated the most clients each month, their first-dose vaccination coverage rate was 11%. Clinic managers had agreed to implement a policy of offering vaccination to all new eligible clients; however, some clinics might have offered vaccine selectively based on clinical judgment of risk or were unable to integrate vaccination into their regular schedules.

Project support for hepatitis B vaccination continues at most high-risk sites. In addition, other viral hepatitis prevention services (e.g., selective hepatitis B and hepatitis C serologic screening, hepatitis A vaccination, and STD screening

TABLE. Number and percentage of adults and adolescents eligible for and receiving Hepatitis B vaccin	ation at sites serving high-
and lower-risk clients, by site, dose, and number of months vaccinating — San Diego, California, Febru	ary 1998–January 2001

Site	No. sites	Eligible no.* monthly/dose 1	Dose 1	Dose 2 <sup>†</sup>	Dose 3 <sup>†</sup>	No. months	Estimated
Uigh rick eliente	01100	montiny/dobe 1	(/0)	(/0)	(/0)	vaconiating	
righ-risk clients							
STD clinic	4	428	(74)§	(53)	(30)	36	20,772
Job Corps	1	64	(66)	(67)	(26)	32	2,592
Center for MSM <sup>¶</sup>	1	26	(50)	(62)	(38)	20	520
Methadone clinic	1	34	(44)	(53)	(40)	10	290
Drug rehabilitation	2	56	(36)	(40)	(35)	20	700
Clinic for MSM	2	24	(25)	(67)	(33)	24	288
Juvenile detention	1	340	(18)	(94)	(31)	18	2,502
Women's jail	1	221	(12)	(65)	(8)	23	1,035
Men's jail	3	1,020	(4)	(51)	(2)	24	1,656
Lower-risk clients							
Teen clinic	2	163	(69)	(80)	(61)	18	4,896
Family planning	1	102	(25)	(68)	(36)	17	867
College health	5	340	(19)	(68)	(40)	15	1,965
University health	1	1,530	(11)	(69)	(44)	19	6,821
Community clinic	4	2,040	(11)	(49)	(34)	28	11,312

\* Estimated as 85% of new client visits (except for jail sites, which used 85% of sick call visits); 85% was selected based on experience of clinics treating sexually transmitted diseases (STDs) that 15% of clients self-reported previous hepatitis B vaccination or disease and were therefore ineligible to start the \_vaccine.

<sup>+</sup>Dose 2–3 percentages determined from individual dose-completion forms of persons receiving first dose and having  $\geq$ 6 months of follow-up at STD clinics, Job Corps, methadone clinic, drug rehabilitation clinic, clinic for men having sex with men, and university health clinic; quarterly aggregate dose  $_{2}$  2–3 reports used at all other sites.

<sup>3</sup>Actual vaccine dose 1 acceptance rate among eligible clients determined from risk-assessment form given all clients at clinics for treatment of STDs.

<sup>11</sup>Men having sex with men.

services) have been or are being integrated into STD clinics, court-ordered drug-offender rehabilitation programs, and anonymous HIV counseling and testing sites. The San Diego VHPP developed a guide for establishing hepatitis B vaccination services in an STD clinic (http://www.cdc.gov/hepatitis/spotlights/integration.htm). The guide has been distributed to all state health department STD, hepatitis C prevention, and vaccination programs.

**Reported by:** P Murray, MPH, C Brennan, MPH, S O'Neill, MS, P Gonzales, R Gilchick, MD, Public Health Svcs, Health and Human Svcs Agency, San Diego County, California. Div of Viral Hepatitis, National Center for Infectious Diseases; R Gunn, MD, Div of Sexually Transmitted Diseases Prevention, National Center for HIV, STD and TB Prevention; D Callahan, MD, Div of Applied Epidemiology and Training, Epidemiology Program Office, CDC.

Editorial Note: Data from the San Diego VHPP indicate that high rates of hepatitis B vaccination can be achieved in some clinics and programs that serve persons at high risk for HBV infection through the integration of hepatitis B vaccination into routine clinic and program services. In the United States, the incidence of reported cases of acute hepatitis B has declined 76% since the late 1980s (3). The greatest decline has occurred among persons aged 10-29 years, and the median age of persons with acute hepatitis B has increased approximately 5 years during the 1990s (3). Universal vaccination of infants and adolescents prevents HBV infections within these age groups and eventually will prevent transmission among adults. However, because it will take several decades to achieve the secondary benefit of hepatitis B vaccination of infants and young adolescents, vaccination of older adolescents and of adults at increased risk for HBV infection is needed to reduce disease incidence and chronic HBV infection prevalence in the near future (3).

As with other vaccines recommended to prevent disease among older adolescents and adults, achieving high levels of hepatitis B vaccine coverage among these groups at increased risk for HBV infection has been difficult. Several obstacles account for low vaccine coverage including 1) inability of health-care providers to identify and deliver vaccine to at-risk populations; 2) lack of a public health infrastructure to support adult vaccination; 3) lack of familiarity by health-care providers with practices required to achieve high rates of adult vaccination; and 4) limited private- and public-sector reimbursement for adult vaccination.

Many persons at increased risk for HBV infection are clients of programs that provide other prevention and clinical services, at times in nonclinical settings. The San Diego VHPP tested the feasibility of vaccinating adults and older adolescents at increased risk for HBV infection at sites that provide services to such persons. For example, hepatitis B vaccination is recommended for all persons seeking care at STD clinics, a setting that provides services to the greatest number of adults at increased risk for HBV infection. Among persons with acute hepatitis B reported annually to a CDC hepatitis surveillance system, approximately 35% have been treated previously for STDs, which indicates the importance of this setting in the prevention of HBV infections (*3*). Earlier attempts at hepatitis B vaccination in STD clinics had limited success; first-dose acceptance rates varied (range: 44%–70%), and <30% of persons completed the 3-dose series (*4*; CDC, unpublished data, 1993, 1997). By providing counseling as part of an integrated service, the San Diego VHPP was able to achieve first-dose acceptance rates as high as 80%.

The goal of hepatitis B vaccination programs is to achieve the highest possible rate of 3-dose vaccination coverage. However, not being able to ensure high 3-dose completion rates should not preclude the initiation of hepatitis B vaccination in STD clinics. Among healthy young adults, protective levels of antibody develop in 30%–55% following a single dose of hepatitis B vaccine and in 75% after 2 doses (5–7). Although long-term (i.e., >10 years) protection cannot be ensured with incomplete vaccination, most persons responding to the first dose are expected to have protection for at least 5 years, which parallels their expected loss of antibody (8). Vaccination completion rates should be monitored, and efforts to increase series completion, especially among those at the highest risk (e.g., MSM and IDUs), should be strongly considered.

Reimbursement remains a major barrier to hepatitis B vaccination of persons at increased risk for infection. Sites (e.g., STD clinics) that serve adolescents aged <19 years can obtain and offer vaccination through reimbursement under the Vaccines for Children (VFC) program (http://www.cdc.gov/nip/ vfc). In the San Diego VHPP, the majority of sites were enrolled with the state vaccination program as VFC providers. However, vaccination of adults was supported only through funding provided by the project. Private- and public-sector health insurance plans rarely cover hepatitis B vaccination for adults. Although some states and local jurisdictions provide hepatitis B vaccine in STD clinics (9), drug-treatment clinics, and prison health programs, many adults with high-risk medical or behavioral conditions have limited access to recommended vaccinations. Providing additional funding to purchase vaccine for uninsured and underinsured adult populations (10) would overcome a major barrier to vaccinating persons at high risk.

The findings in this report are subject to at least three limitations. First, sites for integration of hepatitis B vaccination services were selected on the basis of convenience and might not be representative of all sites. Second, the eligibility criteria used in the STD clinic (i.e., no self-report of previous hepatitis B vaccination or disease) also was used to estimate the percent eligible in all other sites, including sites (e.g., community clinics) that might serve persons for whom hepatitis B vaccination is not specifically recommended. Clinicians at these sites might not have encouraged vaccination for adults without specific risk factors; however, because written risk assessments were not completed for most clients in these settings, the actual percentage of high-risk clients who were offered and received hepatitis B vaccination cannot be determined. Finally, completion rates might be underestimated because persons receiving a first dose of hepatitis B vaccine might not have been followed long enough to track subsequent doses.

The findings in this report suggest that a sustained vaccination program, when combined with a short counseling session, might achieve high levels of vaccine acceptance. Even when vaccination cost is not a barrier, achieving high rates of vaccination coverage requires that program managers set vaccination-coverage goals, train staff, review the vaccination status of all clients routinely, and use appropriate health-education materials and counseling services.

#### References

- CDC. Hepatitis B virus: a comprehensive strategy for eliminating transmission in the United States through universal childhood vaccination. Recommendations of the Immunization Practices Advisory Committee (ACIP). MMWR 1991;40(No. RR-13).
- CDC. Hepatitis B vaccination—United States, 1982–2002. MMWR 2002;51:549–552,563.
- 3. Goldstein ST, Alter MJ, Williams IT, et al. Incidence and risk factors for acute hepatitis B in the United States, 1982–1998: implications for vaccination programs. J Infect Dis 2002;185:713–9.
- Weinstock HS, Bolan G, Moran JS, Peterman TA, Polish L, Reingold AL. Routine hepatitis B vaccination in a clinic for sexually transmitted diseases. Am J Public Health 1995;85:846–9.
- Andre FE. Summary of safety and efficacy data on a yeast-derived hepatitis B vaccine. Am J Med 1989;87:145–20S.
- Davidson M, Krugman S. Recombinant yeast hepatitis B vaccine compared with plasma-derived vaccine: immunogenicity and effect of a booster dose. J Infect 1986;13:31–8.
- 7. Jilg W, Deinhardt F. Results of immunization with a recombinant yeastderived hepatitis B vaccine. J Infect 1986;13:47–51.
- Wainwright RB, McMahon BJ, Bulkow LR, et al. Duration of immunogenicity and efficacy of hepatitis B vaccine in a Yupik Eskimo population. JAMA 1989;261:2362–6.
- 9. Wilson BC, Moyer L, Schmid G, et al. Hepatitis B vaccination in sexually transmitted disease (STD) clinics: a survey of STD programs. Sex Transm Dis 2001;28:148–52.
- Institute of Medicine. Calling the Shots: Immunization Finance Policies and Practices. Washington, DC: National Academy Press, 2000.

### Weekly Update: West Nile Virus Activity — United States, July 10–16, 2002

This report summarizes West Nile virus (WNV) surveillance data reported to CDC through ArboNET and verified by states and other jurisdictions as of July 16, 2002.

During the reporting week of July 10–16, two human cases of WNV were reported, both in Louisiana. During the same period, WNV infections were reported in 55 dead crows, 115 other dead birds, nine horses, and 19 mosquito pools.

During 2002, three human cases of WNV encephalitis or meningitis have been reported, all from Louisiana. Among these cases, all were men, the median age was 62 years (range: 53–78 years), and the dates of illness onset ranged from June 10–28; no cases were fatal. In addition, 171 dead crows and 266 other dead birds with WNV infection were reported from 20 states and the District of Columbia (Figure); 23 WNV infections in horses have been reported from four states (Florida, Kentucky, Louisiana, and Texas). During 2002, WNV seroconversions have been reported in 10 sentinel chicken flocks from Florida; WNV seropositivity has been reported from two states (Indiana and Louisiana) in two wild birds that were caught and released; and 26 WNV-positive mosquito pools have been reported from six states (Alabama, Illinois, Indiana, Massachusetts, New Jersey, and Ohio).

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



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

\* As of July 16, 2002.

### Public Health Dispatch

### Poliomyelitis — Madagascar, 2002

Surveillance for acute flaccid paralysis (AFP) in Madagascar has detected a cluster of four cases of paralytic poliomyelitis from which type-2 vaccine-derived polioviruses have been isolated. Preliminary data indicate that these patients, residing in the Tolagnaro district of Toliara province in southeastern Madagascar, had onset of paralysis during March 20–April 12, 2002. None of the children affected was vaccinated fully. During March–April 2002, provincial authorities conducted a small-scale house-to-house vaccination response. Genetic sequencing studies of these vaccine-derived viruses indicate substantial genetic drift and recombination with nonpolio enteroviruses. These findings are compatible with an outbreak of paralytic polio associated with a circulating vaccine-derived poliovirus (cVDPV); however, further investigation is required.

The three outbreaks of cVDPV described previously occurred in areas where routine oral polio vaccine (OPV) coverage is low, AFP surveillance is suboptimal, and supplementary vaccination activities have not been conducted for years (*1,2*). Vaccination coverage data suggest that during 1999, 37% of children aged <1 year had received 3 doses of OPV. In 2001, the nonpolio AFP rate of 0.3 case per 100,000 population aged <15 years was below the target level of 1.0.

A joint mission by the Ministry of Health of Madagascar, the Pasteur Institute of Madagascar, the World Health Organization, and United Nations Children's Fund (UNICEF) is ongoing to 1) conduct a field investigation of the cases to verify early reports, 2) review health facility records for any missed cases, 3) enhance the quality of AFP surveillance nationwide, and 4) plan for a nationwide house-to-house polio vaccination response. The work of this mission is being complemented by laboratory work in Madagascar, South Africa, France, and the United States.

**Reported by:** Ministry of Health; Pasteur Institute, Madagascar. National Institute for Communicable Diseases, South Africa. Pasteur Institute, Paris, France. World Health Organization Regional Office for Africa, Harare, Zimbabwe. Vaccines and Biologicals Dept, World Health Organization, Geneva, Switzerland. Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases; Global Immunization Div, National Immunization Program, CDC.

### References

- CDC. Acute flaccid paralysis associated with circulating vaccine-derived poliovirus—Philippines, 2001. MMWR 2001;50:874.
- CDC. Outbreak of poliomyelitis—Dominican Republic and Haiti, 2000–2001. MMWR 2001;50:147.

### Erratum: Vol. 51, No. 27

In the Notice to Readers, "Resumption of Routine Schedule for Diphtheria and Tetanus Toxoids and Acellular Pertussis Vaccine and for Measles, Mumps, and Rubella Vaccine," on page 599 under the heading "DTaP Vaccine," an error occurred in the first sentence of the second paragraph. The sentence should read, "During the DTaP vaccine shortage beginning in 2000 (5), ACIP recommended that health-care providers vaccinate infants with the initial 3 DTaP doses, if they did not have *sufficient* supply of DTaP to vaccinate all children in their practice."

## FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending July 13, 2002, with historical data



\* 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 July 13, 2002 (28th Week)\*

		Cum. 2002	Cum. 2001		Cum. 2002	Cum. 2001
Anthrax		2	1	Encephalitis: West Nile <sup>†</sup>	4	-
Botulism:	foodborne	9	11	Hansen disease (leprosy) <sup>†</sup>	40	39
	infant	33	52	Hantavirus pulmonary syndrome <sup>†</sup>	7	5
	other (wound & unspecified)	10	6	Hemolytic uremic syndrome, postdiarrheal <sup>†</sup>	82	63
Brucellosis <sup>†</sup>	,	44	62	HIV infection, pediatric <sup>†§</sup>	98	88
Chancroid		35	23	Plague	-	2
Cholera		4	2	Poliomyelitis, paralytic	-	-
Cyclosporiasis <sup>†</sup>		82	64	Psittacosis <sup>†</sup>	12	7
Diphtheria		1	1	Q fever <sup>†</sup>	19	12
Ehrlichiosis:	human granulocytic (HGE) <sup>†</sup>	102	50	Rabies, human	1	1
	human monocytic (HME) <sup>†</sup>	48	44	Streptococcal toxic-shock syndrome <sup>†</sup>	42	52
	other and unspecified	2	3	Tetanus	9	22
Encephalitis:	California serogroup viral <sup>†</sup>	7	6	Toxic-shock syndrome	68	70
	eastern equine <sup>†</sup>	1	-	Trichinosis	9	10
	Powassan <sup>†</sup>	-	-	Tularemia <sup>†</sup>	27	54
	St. Louis <sup>†</sup>	-	-	Yellow fever	1	-
	western equine <sup>†</sup>	-	-			

-:No reported cases.

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

<sup>†</sup>Not notifiable in all states.

<sup>§</sup> 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 June 30, 2002.

### **MMWR**

								Escherie	chia coli	
	AID	s	Chlan	nvdia⁺	Cryptos	poridiosis	015	7:H7	Shiga Toxi Serogrour	n 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	20,967	20,376	386,540	401,933	1,045	1,070	983	1,046	38	45
NEW ENGLAND	802	731	13,879	11,565	63	51	78	111	8	20
Maine N.H.	19 19	20 15	782 849	642 708	2 14	4	3 7	12 12	-	- 3
Vt.	8	10	344	315	14	13	3	5	-	-
R.I.	62	51	1,487	4,400	13	25	5	6	-	-
Conn.	317	234	4,681	3,937	5	4	21	17	4	12
Upstate N.Y.	4,702 359	5,358 782	39,721 8,606	43,383 6,943	117 35	146 42	74 61	82 48	-	-
N.Y. City	2,554	2,968	15,057	15,965	55	60	4	8	-	-
Pa.	977	689	12,673	13,401	20	37	N	N	-	-
E.N. CENTRAL	2,241	1,404	66,447	74,241	267	358	252	241	1	3
Ind.	433 306	232 163	18,027 8,711	19,216 8,236	70 24	56 32	56 24	59 37	1	2
III. Mich	1,029	670	16,866	22,290	40 54	40	76	61	-	-
Wis.	109	78	6,253	8,599	79	157	40 56	57	-	-
W.N. CENTRAL	330	449	20,973	20,752	115	98	154	125	4	2
lowa	72 47	81 47	4,987 2,724	4,181 2,540	50 13	32 25	54 40	47 20	-	-
Mo. N. Dak	138	209	7,640	7,319	16	20	23	23	N	N
S. Dak.	2	18	1,150	950	5	5	17	8	1	1
Nebr. Kans.	31 39	47 46	589 3,414	1,867 3,336	16 9	12	9 8	15 11	-	1
S. ATLANTIC	6,499	6,108	75,501	77,235	167	170	100	90	15	13
Del. Md	114 961	115 753	1,426 7 796	1,550 8 141	1	1 27	4	1	-	-
D.C.	321	460	1,694	1,810	3	9	-	-	-	-
va. W.Va.	488 50	541 47	8,887 1,244	9,365 1,258	4 2	9 1	24	24 3	1	2
N.C.	456	376	12,797	11,286	23	17	17	26	-	-
Ga.	1,087	750	13,981	16,326	80	68	34	16	9	7
FIA.	2,567	2,728	20,643	19,100	43	36	14	12	5	4
Ky.	150	201	4,578	4,730	1	3	14	23	-	-
Tenn. Ala	404 173	271 224	8,459 8 157	7,812 7,509	38 28	4	21 7	18 8	-	-
Miss.	192	257	5,244	6,468	4	7	5	3	-	-
W.S. CENTRAL	2,181	2,021	55,305	57,264	15	35	13	112	-	-
La.	508	458	9,943	9,438	4	7	-	3	-	-
Okla. Tex.	119 1,405	106 1,353	5,485 36,550	5,795 37,932	6	6 19	10	13 92	-	-
MOUNTAIN	678	711	23,795	23,583	75	59	98	104	6	3
Mont.	6 15	12 15	1,143 1 324	1,155	4 17	5	9 7	6 14	- 2	- 2
Wyo.	4	1	467	431	6	1	2	4	1	-
Colo. N. Mex.	133 51	153 59	7,096 3,234	6,512 3,181	20 9	18 11	34 5	44 6	1 1	1
Ariz.	284	279	7,334	7,797	10	3	12	12	1	-
Nev.	150	130	2,060	2,671	3	3	10	6	-	-
PACIFIC	2,615	2,641	64,481	67,391	155	132	167	129	4	4
Oreg.	264 196	284 110	7,495 3,604	7,251 3,866	24 21	15	20 45	29 21	4	4
Calif. Alaska	2,090 12	2,205 14	49,414 1 860	52,738 1 443	109	114	78 4	69 2	-	-
Hawaii	53	28	2,108	2,093	1	3	20	8	-	-
Guam PB	2	8 579	- 1 576	221	-	-	Ν	Ν	-	-
V.I.	60	2	30	94		-		-	-	-
C.N.M.I.	2	U	117	U U	-	U	U -	U	-	U

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending July 13, 2002, and July 14, 2001 (28th Week)\*

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 June 30, 2002.

(28th Week)*							Haemophilus influenzae,					
							Inva	isive				
	Escheri	chia coli	-					Age <5	Years			
	Shiga Toxi Not Sero	n Positive, arouped	Giardiasis	Gono	rrhea	All All Se	Ages, rotypes	Serot B	уре			
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001			
UNITED STATES	17	4	7,198	163,406	182,299	881	878	12	15			
NEW ENGLAND	-	1	741	3.987	3.118	63	56	-	1			
Maine	-	-	82	62	70	1	1	-	-			
N.H.	-	-	25	64	82	5	-	-	-			
Vt. Mass	-	1	57	45	1 2 2 2	5	2	-	-			
R.I.	-		68	474	378	9	2	-	-			
Conn.	-	-	158	1,562	1,217	13	18	-	-			
MID. ATLANTIC	-	-	1,608	18,235	20,642	153	125	3	3			
Upstate N.Y.	-	-	554	4,409	4,293	69	39	2	-			
N.Y. City	-	-	641	6,133	6,648	34	34	-	-			
N.J. Pa	-	-	144 269	2,829 4 864	3,232	19	28	- 1	- 3			
	0	0	1 201	01,001	0,100	144	154	0	1			
Ohio	8	2	410	9 598	10,354	55	154	2	1			
Ind.	-	-	-	3,776	3,418	31	28	1	-			
III.	-	-	304	9,119	12,007	43	52	-	-			
Mich.	-	-	398	7,265	9,274	9	8	1	-			
VVI5.	-	-	209	1,920	3,033	0	10	-	-			
W.N. CENTRAL	-	-	847	8,157	8,550	33	38	-	1			
lowa	-	-	119	602	648	20	- 20	-	-			
Mo.	Ν	Ν	243	4,406	4,336	9	12	-	-			
N. Dak.	-	-	11	27	19	-	4	-	-			
S. Dak. Nobr	-	-	35	138	146	-	-	-	-			
Kans.	-		78	1.390	1.448	3	1	-	-			
			1 275	42,620	47.054	220	221	1	1			
Del.	-	-	26	859	887	-	-	-	-			
Md.	-	-	50	4,339	4,625	52	56	1	-			
D.C.	-	-	20	1,408	1,560	-	-	-	-			
va. W Va	-	-	20	5,375	4,997	16	18	-	- 1			
N.C.	-	-	-	8,535	8,790	21	31	-	-			
S.C.	-	-	35	4,212	6,242	11	4	-	-			
Ga.	-	-	497	7,615	8,801	67	59 45	-	-			
	_	_	310	10,734	10,024	47	40	-	_			
E.S. CENTRAL	1	1	1/2	15,029	16,969	37	56	1	-			
Tenn.	-	-	78	4.821	5.137	20	27	-	-			
Ala.	-	-	94	5,250	5,804	9	25	1	-			
Miss.	-	-	-	3,136	4,193	5	2	-	-			
W.S. CENTRAL	-	-	89	24,322	27,735	33	34	2	1			
Ark.	-	-	66 1	1,862	2,590	1	-	-	-			
∟a. Okla.	-		22	2.344	2.607	28	27	-	-			
Tex.	-	-	-	13,958	15,968	2	1	2	1			
MOUNTAIN	8	-	665	4,964	5,505	116	96	2	3			
Mont.	-	-	35	55	69	-	-	-	-			
Idaho	-	-	46	40	42	2	1	-	-			
Colo	8	-	219	1 704	1 657	21	26	-	-			
N.Mex.	-	-	77	623	519	18	14	-	-			
Ariz.	-	-	85	1,785	2,153	55	40	1	1			
Utan	-	-	123	107	80	14	5	- 1	- 2			
			480	10 400	14 640	80	00	1	2			
Wash	-	-	480	1 4 8 5	14,640	2	98	1	4			
Oreg.	-	-	198	434	615	42	30	-	-			
Calif.	-	-	-	10,859	11,910	12	44	-	4			
Alaska Hawaii	-	-	48	327	202	1	3	-	-			
	-	-	49	301	228	20	20	-	-			
Guam PR	-	-	- 11	- 237	24	- 1	- 1	-	-			
V.I.	-	-	-	17	14	-	-	-	-			
Amer. Samoa	U	U	U	U	U	U	U	U	U			
A INTIVIT	-		-	11		-	U	-				

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending July 13, 2002, and July 14, 2001

N: Not notifiable. U: Unavailable. - : No reported cases. \* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

	На	emophilus ir	<i>fluenzae</i> , Invas	sive							
		Age <	<5 Years		Hepatitis (Viral, Acute), By Type						
	Non-Se	rotype B	Unknown	Serotype		A		В	C; Non-A	A, Non-B	
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	
UNITED STATES	138	148	12	17	4,420	4,766	3,367	3,648	1,704	2,215	
NEW ENGLAND	7	10	-	-	181	269	113	70	18	27	
Maine	-	-	-	-	6	5	4	5	-	-	
Vt.	-	-	-	-	1	6	3	5	11	6	
Mass.	4	7	-	-	82	108	59	13	7	21	
R.I.	-	-	-	-	27	12	17	12	-	-	
	3	3	-	-	55	131	750	20	-	-	
MID. AI LANTIC	21	20	-	3	549 108	625 145	750 79	68	31	643 18	
N.Y. City	6	5	-		228	227	415	346	-	-	
N.J.	4	3	-	-	64	150	146	147	759	587	
Pa.	3	6	-	2	149	103	110	158	16	38	
E.N. CENTRAL	20	28	-	1	615	574	424	431	58	109	
Ind.	5 7	8	-	1	32	45	58 18	62 24	-	1	
III.	7	11	-	-	168	178	40	54	8	9	
Mich.	-	_	-	-	125	178	308	268	44	92	
VVIS.	1	5	-	-	94	42	-	23	-	-	
W.N. CENTRAL	2	2	3	2	186	204	114	114	476	680	
lowa	-	-	-	-	20 46	19	11	12	1	-	
Mo.	-	-	2	2	51	45	65	66	467	672	
N. Dak.	-	1	-	-	1	2	4	-	-	-	
S. Dak. Nebr	-	-	-	-	3	1 27	- 14	1 14	-	- 3	
Kans.	-	-	-	-	54	94	12	10	2	3	
S ATI ANTIC	33	30	2	5	1 328	887	873	665	89	36	
Del.	-	-	-	-	9	4	7	13	5	2	
Md.	1	4	-	1	163	129	67	72	6	4	
D.C.	-	-	-	-	49	22	10	9	-	-	
w.va.	-	4	1	-	10	7	13	16	2	6	
N.C.	3	1	-	4	131	77	134	110	14	10	
S.C.	4	1	-	-	42	34	56	15	4	4	
Ga. Fla	16	14	- 1	-	561	484	282	203	23 34	- 10	
	8	11	1	2	156	195	185	248	106	140	
Kv.	-	-	-	1	35	48	28	240	2	5	
Tenn.	5	5	-	-	60	74	75	125	20	40	
Ala.	3	5	1	1	23	58	40	51	3	2	
IVIISS.	-	1	-	-	38	15	42	45	01	93	
W.S. CENTRAL	6	4	-	-	67	542	214	430	22	457	
la	- 1	-	-	-	∠5 16	38 58	28	50 66	4 14	5 103	
Okla.	5	4	-	-	25	82	15	66	4	4	
Tex.	-	-	-	-	1	364	110	242	-	345	
MOUNTAIN	24	12	5	1	334	408	259	268	52	38	
Mont.	-	-	-	-	9	6	3	2	-	1	
Wvo.	-	-	-	-	20	40	9	0	7	4	
Colo.	2	-	-	-	55	40	49	60	23	5	
N.Mex.	4	6	1	1	9	18	44	71	-	11	
Ariz. Litah	12	4	3	-	35	210	94 23	87 15	3	9	
Nev.	1	-	1	-	29	48	32	24	17	6	
PACIFIC	17	31	1	3	1.004	1.062	435	703	77	85	
Wash.	1	-	-	1	97	55	33	67	13	16	
Oreg.	4	5	-	-	49	69	78	_88	13	10	
Jaska	9	24 1	1	1	850	916	318	530	51	59	
Hawaii	2	1	-	1	1	10	3	14	-	-	
Guam	-	-	-	-	-	1	-	-	-	-	
P.R.	-	1	-	-	58	95	47	147	-	1	
V.I.					-						
C.N.M.I.	U -	U	U -	U	-	U	31	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).

626

<u></u>	Legionellosis		Listeriosis		Lyme	Lyme Disease		Malaria		Measles Total	
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	
Reporting Area	2002	2001	2002	2001	2002	2001	2002	2001	2002	2001	
UNITED STATES	383	473	217	276	3,399	5,005	564	704	10†	84 <sup>§</sup>	
NEW ENGLAND	22	19	25	28	369	1,222	34	45	-	5	
N.H.	4	4	2	1	52	26	5	2	-	-	
Vt.	3	4	.1	-	4	4	1	-	-	1	
Mass. R I	9	5	15 1	15 1	244	578	13	21	-	3	
Conn.	4	4	4	11	23	491	11	16	-	1	
MID. ATLANTIC	91	102	39	47	2,407	2,731	121	185	5	12	
Upstate N.Y.	30	28	18	13	1,488	802	21	24	-	4	
N.Y. City	18 10	11	11	13	162	43	76 13	114 26	5	2	
Pa.	33	56	7	13	680	879	11	21	-	5	
E.N. CENTRAL	89	131	26	41	29	409	66	94	1	10	
Ohio	39	56	9	8	24	10	12	13	1	3	
Ind.	8	10 17	4	4	5	6 23	3 17	12	-	4	
Mich.	30	27	9	13	-	2	27	19	-	-	
Wis.	12	21	3	3	U	368	7	11	-	-	
W.N. CENTRAL	24	29	8	6	84	89	41	21	-	4	
Minn. Iowa	2	7	- 1	-	48 14	49 16	14	6	-	2	
Mo.	10	9	5	3	18	20	11	7	-	2	
N. Dak.	-	1	1	-	-	-	1	-	-	-	
S. Dak. Nebr	2	2	-	- 1	-	- 2	- 5	- 2	-	-	
Kans.	-	1	1	2	4	2	8	3	-	-	
S. ATLANTIC	91	74	38	32	413	423	158	149	1	4	
Del.	5	2	-	1	54	59	1	1	-	-	
Ma. D.C	15	21	5	4	229	266	44	64 9	-	3	
Va.	8	11	3	5	25	66	12	30	-	-	
W.Va.	N	N	-	4	5	8	2	1	-	-	
S.C.	5	3	5	2	52 5	2	9 5	6 4	-	-	
Ga.	10	8	10	7	1	-	55	21	-	1	
Fla.	38	22	12	6	30	5	23	13	1	-	
E.S. CENTRAL	12	37	8	10	25	21	9	15	-	2	
ry. Tenn.	1	9 16	2	4	12	7	2	4	-	2	
Ala.	4	8	3	3	6	4	3	3	-	-	
Miss.	-	4	-	-	-	3	2	2	-	-	
W.S. CENTRAL	3	16	4	23	2	57	3	49	-	1	
Агк. La.	- 1	-	-	-	- 1	- 4	2	3	-	-	
Okla.	2	3	4	1	-	-	-	2	-	-	
Tex.	-	7	-	21	1	53	-	40	-	1	
MOUNTAIN	17	28	18	25	12	6	27	29	-	1	
Idaho	-	1	2	- 1	2	3	-	2	-	- 1	
Wyo.	1	2	-	1		1	-	-	-	-	
Colo.	4	11	2	5	3	-	14	15	-	-	
Ariz.	3	8	9	6	2	-	5	3	-	-	
Utah	6	2	3	1	3	-	4	2	-	-	
Nev.	1	2	-	5	1	2	3	2	-	-	
PACIFIC	34	37	51	64	58	47	105	117	3	45	
Oreg.	N	N	3	4	8	6	5	8	-	2	
Calif.	31	26	39	56	49	38	81	97	3	22	
Alaska Hawaii	-	1 4	- 5	- 1	1 N	2 N	2	1 7	-	-	
Guam	_	- -	-	-	-	-	_	-	-	-	
P.R.	-	2	1	-	N	N	-	3	-	-	
V.I.		-									
Amer. Samoa C N M I	U -	U	U -	U	U -	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).

 † Of 10 cases reported, three were indigenous and seven were imported from another country.

 § Of 84 cases reported, 41 were indigenous and 43 were imported from another country.

<u> </u>	Meningo Dise	ococcal ase	Mu	mps	Per	tussis	Rabies, Animal		
Reporting Area	Cum. 2002	Cum. 2001	Cum.	Cum. 2001	Cum.	Cum. 2001	Cum.	Cum. 2001	
UNITED STATES	942	1.501	153	122	3.325	2.692	2.818	3.670	
NEW ENGLAND Maine N.H.	63 4 8	72 1 9	7-4	-	325 5 6	254 - 14	409 23 11	332 36 6	
Vt. Mass. R.I.	4 30 4	4 43 2	2	- - -	56 248 4	24 200 2	60 140 31	37 118 30	
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	94 32 13 12 37	158 45 25 27 61	14 2 1 1 10	- 14 2 8 - 4	158 112 7 3 36	200 103 33 8 56	521 316 10 75 120	603 368 14 98 123	
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	143 54 23 27 27 12	210 57 23 51 48 31	17 3 1 6 6 1	17 1 12 2 1	411 224 22 65 32 68	318 166 24 36 28 64	38 10 8 8 12	44 14 5 17 7	
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr.	85 22 12 34 - 2 10	99 15 21 35 5 4 10	11 3 - 3 1 -	5 2 - - - 1	314 109 107 61 5 4	121 31 15 55 3 3 3	213 16 33 21 11 32	200 19 43 18 24 29 4	
Kans. S. ATLANTIC Del. Md. D C	5 164 6 4	9 229 3 32	4 17 - 3	2 17 - 4	28 209 21 1	14 121 - 18 1	100 1,211 24 165	63 1,276 22 262	
Va. W.Va. N.C. S.C. Ga. Fla	28 - 19 15 24 68	28 8 55 22 34 47	3 - 1 2 4 4	2 - 1 7 2	88 12 20 28 16 21	12 1 40 21 16 12	262 95 360 43 132 130	228 67 318 71 202 106	
E.S. CENTRAL Ky. Tenn. Ala. Miss.	60 10 24 16 10	97 17 41 29 10	11 4 2 2 3	- 3 1 - 2	102 39 36 20 7	57 13 25 16 3	89 16 49 24	145 12 106 27	
W.S. CENTRAL Ark. La. Okla. Tex.	54 20 17 16 1	235 13 57 21 144	11 - 1 - 10	9 - 2 - 7	764 339 4 41 380	252 11 4 9 228	64 _ 64 _	727 - 5 43 679	
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah	62 2 3 - 20 3 19 4	71 3 7 4 27 8 11 7	12 - 1 - 2 1 1 4	8 - 1 2 2 1 1	452 2 46 7 181 82 89 27	907 10 165 - 171 50 461 39	132 7 8 13 20 4 76 2	137 20 2 20 - 5 87 2	
Nev. PACIFIC Wash. Oreg. Calif. Alaska Hawaii	11 217 42 34 134 1 6	4 330 43 39 238 2 8	3 53 - N 43 - 10	1 49 1 26 1 21	18 590 264 102 213 4 7	11 462 76 30 331 2 23	2 141 2 115 24	1 206 - 168 38	
Guam P.R. V.I. Amer. Samoa C.N.M.I.	3 - - -	- 4 - U U	- - - U	- - - U U	, 1 - U 1	- - - U U	43 U	62 U U	

N: Not notifiable. -: No reported cases. \* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

, ,				R					
	Rocky M Spotte	lountain d Fever	Rul	pella	Conge Rub	enital ella	Salmonellosis		
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	
UNITED STATES	339	215	5	15	2	-	15,304	16,861	
NEW ENGLAND	-	2	-	-	-	-	926	1,228	
Maine	-	-	-	-	-	-	72	110	
N.H.	-	-	-	-	-	-	61	95	
VI. Mass	-	- 2	-	-	-	-	34 513	35 715	
R.I.	-	-	-	-	-	-	59	64	
Conn.	-	-	-	-	-	-	187	209	
MID. ATLANTIC	19	11	3	6	-	-	1.934	2.291	
Upstate N.Y.	5	-	2	1	-	-	693	516	
N.Y. City	2	1	-	4	-	-	621	627	
N.J.	3	2	1	1	-	-	192	539	
ra.	9	0	-	-	-	-	428	609	
E.N. CENTRAL	6	13	-	2	-	-	2,474	2,326	
Ind	4	1	-	-	-	-	682 211	680 230	
III.	-	11	-	2	_	-	770	649	
Mich.	1	-	-	-	-	-	440	402	
Wis.	-	-	-	-	-	-	371	365	
W.N.CENTRAL	47	30	-	3	-	-	1,142	980	
Minn.	-	-	-	-	-	-	264	304	
lowa	1	1	-	1	-	-	195	152	
MO. N Dak	46	27	-	1	-	-	423	237	
S Dak	-	2	-	-	-	-	44	70	
Nebr.	-	-	-	-	-	-	51	68	
Kans.	-	-	-	1	-	-	140	134	
S. ATLANTIC	195	89	-	3	-	-	3,721	3,645	
Del.	2	-	-	-	-	-	ິ 31	42	
Md.	25	15	-	-	-	-	382	374	
D.C.	-	-	-	-	-	-	40	39	
va. W Va	12	o -	-	-	-	-	401	53	
N.C.	102	44	-	-	-	-	528	517	
S.C.	32	13	-	2	-	-	210	360	
Ga.	18	6	-	-	-	-	813	672	
Fla.	3	3	-	1	-	-	1,270	1,006	
E.S. CENTRAL	34	44	-	-	1	-	1,061	955	
Ky. Tann	2	1	-	-	-	-	164	166	
Ala	24	35	-	-	-	-	203 305	249 276	
Miss.	-	4	-	-	-	-	329	264	
	28	10	1	_	_	-	614	2 003	
Ark.	-	4	-	-	-	-	315	2,003	
La.	-	1	-	-	-	-	118	347	
Okla.	28	14	-	-	-	-	179	147	
lex.	-	-	1	-	-	-	2	1,250	
MOUNTAIN	8	7	-	-	-	-	1,031	1,014	
Mont.	1	1	-	-	-	-	48	39	
Wvo	2	2	-	-	-	-	29	31	
Colo.	1	-	-	-	-	-	261	279	
N.Mex.	-	1	-	-	-	-	143	125	
Ariz.	-	-	-	-	-	-	290	270	
Utan Nev	-	2	-	-	-	-	92 108	87	
	-						0.01	0,110	
Mach	2	-	1	1	1	-	2,401	2,419	
Orea.	- 1	-	-	-	-	-	225	145	
Calif.	1	-	1	-	-	-	1,809	1,835	
Alaska	-	-	-	-	-	-	35	25	
Hawaii	-	-	-	1	1	-	131	188	
Guam	-	-	-	-	-	-		10	
P.R.	-	-	-	3	-	-	101	484	
v.i. Amer Samoa	-	-	-	-	-	-	-	-	
C.N.M.I.	-	Ŭ	-	Ŭ	-	Ŭ	21	Ŭ	

N: Not notifiable. - : No reported cases. \* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

	Shig	ellosis	Streptococ Invasive	cal Disease, , Group A	Streptococcu Drug Resist	<i>s pneumoniae,</i> ant, Invasive	Streptococcus	<i>pneumoniae</i> , <5 Years)
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001
UNITED STATES	6,978	8,130	2,443	2,286	1,313	1,827	136	265
NEW ENGLAND Maine N.H. Vt	127 3 5	132 5 2	119 14 25 9	159 10 N	8 - - 3	85 - - 7	1 - N 1	30 - N
Mass.	88	92	58	51	Ň	Ň	Ň	Ν
R.I. Conn.	7 24	8 22	13	8 81	5	- 78	-	2 28
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	399 97 189 48 65	839 316 226 152 145	414 209 103 71 31	411 179 121 73 38	76 68 U N 8	115 113 U N 2	43 43 U N	73 73 U N
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	720 356 39 194 76 55	1,372 693 125 268 152 134	393 145 29 30 189	547 138 43 178 140 48	124 N 119 2 3 N	126 N 126 - N	53 N 28 - N 25	68 N 38 30 N
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. S. Dak. Nebr.	595 130 62 81 15 149 104 54	812 251 238 139 13 84 41	169 87 - 37 - 9 13 23	222 80 - 55 7 7 7 28 45	146 48 N 6 1 1 23 67	85 40 N 9 4 3 9	33 33 N - - N N	31 24 N - 7 N N
S. ATLANTIC Del. Md. D.C. Va. W.Va. N.C. S.C. Ga. Fla.	2,756 11 482 34 493 4 155 46 894 637	1,122 5 58 30 106 5 203 144 146 425	496 1 83 5 50 12 93 28 129 95	393 2 N 3 60 16 107 7 131 67	806 3 N 42 N 34 N 128 249 350	973 2 N 3 N 36 N 199 278 455	1 N 1 U N U N	4 N N 3 N 1 U N U N U N
E.S. CENTRAL Ky. Tenn. Ala. Miss.	681 75 33 348 225	808 295 50 146 317	68 12 56 -	50 18 32	91 10 81 -	174 18 155 1	N N N	N N N
W.S. CENTRAL Ark. La. Okla. Tex.	408 110 63 234 1	1,502 374 144 20 964	39 5 - 33 1	218 - - 31 187	34 5 29 N N	239 13 196 N N	2 - 1 -	59 - 59 -
MOUNTAIN Mont. Idaho	301 2 2	423 19	413 - 5	248 - 4	28 N	29 - N	3 - N	- N
Wyo. Colo. N. Mex. Ariz. Utah Nev.	3 59 57 139 23 16	2 86 64 193 27 32	7 147 68 177 9 -	7 99 53 82 3	9 - 19 - - -	5 - 22 - - 2	- N 3	N
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	991 70 48 844 2 27	1,120 97 59 933 4 27	332 36 N 260 - 36	38 N - - 38	N N	1 - N - 1		N N N
Guam P.R. V.I.	- 5	31 12	- N	1 N	- - -	-	N	N
Amer. Samoa C N M I	U 14	U	U	U	-	-	U	U

N: Not notifiable. -: No reported cases. \* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

(20th Week)		Syr	ohilis			Typhoid			
	Primary &	Secondary	Cong	enital	Tubero	ulosis	Fe	ver	
Reporting Area	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	
UNITED STATES	3,234	3,009	160	279	5,846	6,926	129	166	
NEW ENGLAND Maine	65	27	-	3	207 5	245 10	10	8 1	
N.H. Vt	1	1	-	-	7	11 4	-	1	
Mass.	47	15	-	2	104	117	8	5	
K.I. Conn.	2 14	3 6	-	- 1	27 64	38 65	2	- 1	
MID. ATLANTIC Upstate N.Y. N.Y. City	364 19 204	260 10 149	25 3 11	40 2 20	1,092 158 577	1,173 165 601	36 5 19	57 13 21	
N.J. Pa.	68 73	49 52	10 1	18	247 110	269 138	9 3	20 3	
E.N. CENTRAL Ohio Ind. III. Mich.	562 75 42 150 287	522 49 95 159 202	24 - - 18 6	41 2 5 27 4	548 95 58 270 119	694 134 48 349 126	13 4 2 1 3	20 2 2 9 4	
Wis.	8	17	-	3	6	37	3	3	
W.N. CENTRAL Minn. Iowa Mo. N. Dak.	52 18 2 16	43 20 3 9		5 1 - 3 -	282 122 17 81 1	266 116 18 59 3	4 3 - 1	6 2 - 4 -	
S. Dak. Nebr. Kans.	- 4 12	- 1 10	-	- - 1	9 9 43	8 21 41	- -	-	
S. ATLANTIC	860 8	1,067	38	72	1,227 7	1,338 9	16	21	
Md.	103	138	5	2	140	114	3	6	
D.C. Va.	48 41	15 61	1	2 4	- 93	37 127	-	- 6	
W.Va.	- 158	- 249	- 14	- 8	12 167	16 180	-	- 1	
S.C.	67	145	3	18	102	115	_	-	
Ga. Fla.	152 283	178 272	1 13	14 24	201 505	261 479	7 6	6 2	
E.S. CENTRAL	287	322	10	21	385	440	4	-	
Ky. Tenn.	52 110	25 179	2	- 13	71 147	70 157	4	-	
Ala.	97	58	4	4	120	145	-	-	
WISS.	28 433	369	39	4	47 713	1 109	-	- 11	
Ark.	12	22	1	5	71	73	-	-	
La. Okla	66 36	70 37	- 2	-	- 69	65 74	-	-	
Tex.	319	240	36	39	573	897	-	11	
MOUNTAIN	145	111	9	16	191	259	9	6	
Idaho	2	-	1	-	8	3	-	-	
Wyo.	-	-	-	-	2	1	-	-	
N. Mex.	25	10	-	1	25 21	34	5	-	
Ariz.	100	76	7	14	101	99	-	1	
Nev.	4	3	-	-	16	42	1	4	
PACIFIC	466	288	15	34	1,201	1,402	37	37	
Wash. Oreg	26 7	32	1	-	124	124	4	3	
Calif.	428	243	13	34	925	1,119	31	29	
Alaska Hawaii	- 5	-	-	-	32 70	24 83	-	- 2	
Guam	-	2	-	-	-	37	-	2	
P.R. V.I.	126	134	10	2	33	53	-	-	
Amer. Samoa C.N.M.I.	U 13	U U	U	U U	U 27	U U	U	U U	

N: Not notifiable. - : 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 July 13, 2002 (28th 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&l⁺ Total
NEW ENGLAND	511	350	101	35	13	12	32	S. ATLANTIC	1,120	687	266	108	32	26	79
Boston, Mass.	192	116	47	12	9	8	8	Atlanta, Ga.	74	37	23	10	4	-	2
Bridgeport, Conn.	31	24	5	2	-	-	-	Baltimore, Md.	167	97	45	14	8	3	11
Cambridge, Mass.	20	13	5	2	-	-	1	Charlotte, N.C.	103	60	23	11	4	4	5
Hartford Conn	42	30	0	- 	- -	- U	4	Miami Fla	80	45	32 16	20 12	2	2	21
Lowell Mass	28	22	4	2	-	-	5	Norfolk Va	70	40	18	8	2	2	5
Lynn, Mass.	17	13	3	1	-	-	1	Richmond, Va.	63	41	14	5	2	1	8
New Bedford, Mass.	24	21	2	1	-	-	-	Savannah, Ga.	55	33	12	5	-	5	2
New Haven, Conn.	31	23	6	1	-	1	2	St. Petersburg, Fla.	53	39	11	1	1	1	6
Providence, R.I.	U	U	U	U	U	U	U	Tampa, Fla.	169	119	35	12	1	2	11
Somerville, Mass.	6	3	2	1	-	-	-	Washington, D.C.	102	60	25	10	4	3	3
Springfield, Mass.	40	30	5	4	1	-	6	Wilmington, Del.	17	5	12	-	-	-	-
Worcester Mass	32 48	20	9 9	6	2	2	4	E.S. CENTRAL	673	452	138	55	15	10	57
					-	-		Birmingham, Ala.	153	111	28	10	1	-	18
MID. ATLANTIC	2,118	1,478	437	132	39	32	103	Chattanooga, Tenn.	91	61	22	6	2	-	4
Albany, N.Y.	53	35	12	5	-	1	5	Knoxville, Ienn.	84	51	17	13	2	1	2
Ruffalo N Y	19	82	21	5	-	3	15	Memphis Tenn	00	50	21	Ú,	2		11
Camden NJ	21	14	-	3	-	4	-	Mobile Ala	48	31	10	3	3	1	1
Elizabeth, N.J.	28	19	9	-	-	-	-	Montgomery, Ala.	49	36	10	2	1	-	7
Erie, Pa.	26	19	7	-	-	-	1	Nashville, Tenn.	160	104	30	14	4	8	19
Jersey City, N.J.	53	39	11	1	1	1	-	W.S. CENTRAL	1 406	911	280	121	62	31	105
New York City, N.Y.	1,264	876	264	86	25	13	48	Austin. Tex.	83	52	23	6	1	1	2
Newark, N.J.	64	32	21	4	6	1	1	Baton Rouge, La.	48	29	6	5	5	3	2
Paterson, N.J.	28	15	6	3	1	3	1	Corpus Christi, Tex.	55	35	12	4	3	1	5
Pittsburgh Pa §	51	30	16	1	3	1	4	Dallas, Tex.	181	106	45	20	7	3	8
Reading Pa	22	17	3	1	-	1	3	El Paso, Tex.	89	60	19	8	1	1	6
Rochester, N.Y.	147	112	27	6	-	2	13	Ft. Worth, Iex.	116	/6	18	9	5	8	12
Schenectady, N.Y.	21	19	1	1	-	-	4	Houston, lex.	360	223	/1	33	24	8	34
Scranton, Pa.	37	31	6	-	-	-	-	New Orleans La	44	24	11	5	3	1	-
Syracuse, N.Y.	97	72	13	8	3	1	3	San Antonio. Tex.	216	147	38	19	10	2	13
Irenton, N.J.	34	22	(	4	-	1	-	Shreveport, La.	74	54	13	5	-	2	8
Yonkers, N.Y.	24	12	6 4	3	-	-	3	Tulsa, Ökla.	140	105	24	7	3	1	15
E.N. CENTRAL	1,405	946	265	90	40	28	82		832	538	178	68	28	20	50
Akron, Ohio	U	U	U	U	U	U	U	Boise Idaho	90 60	41	19	21	2	4	0
Canton, Ohio	40	27	12	1	-	-	4	Colo, Springs, Colo,	58	44	13	1	-	-	3
Chicago, III.	U	U	U	U	U	U	U	Denver, Colo.	118	68	29	13	5	3	6
Cincinnati, Onio	101	77	0	14	0	0	0	Las Vegas, Nev.	189	118	50	11	6	4	8
Columbus Ohio	176	122	32	14	4	7	13	Ogden, Utah	U	U	U	U	U	U	U
Davton. Ohio	114	78	26	6	3	1	5	Phoenix, Ariz.	U	U	U	U	U	U	U
Detroit, Mich.	128	78	29	10	7	4	7	Pueblo, Colo.	19	15	3	1	-		3
Evansville, Ind.	60	47	8	4	1	-	3	Tucson Ariz	154	115	20	9 7	7	1	12
Fort Wayne, Ind.	83	60	16	6	1	-	6				27				12
Gary, Ind.	1/	13	1	1	1	-	-	PACIFIC Barkalay Calif	1,671	1,148	338	112	43	30	95
Indiananolis Ind	33 175	20	9 18	1	1	2	11	Erespo Calif	20 113	80	10	10	-	-	4
Lansing Mich	48	35	10	1	1	1	3	Glendale Calif	23	20	2	1	-	_	-
Milwaukee. Wis.	119	80	28	5	4	2	7	Honolulu, Hawaii	78	61	14	2	-	1	3
Peoria, III.	73	35	15	16	6	1	5	Long Beach, Calif.	91	60	20	6	3	2	7
Rockford, III.	U	U	U	U	U	U	U	Los Angeles, Calif.	365	248	73	27	11	6	-
South Bend, Ind.	46	32	9	4	1	-	-	Pasadena, Calif.	24	15	3	4	1	1	1
Toledo, Ohio	102	76	18	5	1	2	5	Portland, Oreg.	112	75	28	4	2	3	4
Youngstown, Ohio	70	56	10	1	1	2	3	Sacramento, Calif.	207	145	41	17	5	1	18
W.N. CENTRAL	432	292	98	30	8	4	39	San Erancisco Calif	174	107	40	17 11	11	4	21
Des Moines, Iowa	81	60	18	2	1	-	12	San Jose Calif	126	87	29	10	-	-	8
Duluth, Minn.	26	18	5	3		-	2	Santa Cruz. Calif.	31	22	6	3	-	-	2
Kansas City, Kans.	U	U	U	U	U	U	U	Seattle, Wash.	147	96	31	8	8	4	6
Lincoln Nebr	/5 20	42	21	9	3	-	Ю	Spokane, Wash.	52	43	6	2	1	-	4
Minneanolis Minn	29 60	42	12	<u>ک</u> ۸	1	- 1	- 6	Tacoma, Wash.	108	78	19	8	2	1	9
Omaha, Nebr.	100	65	22	9	2	2	6	TOTAL	10.168 <sup>¶</sup>	6.802	2,101	751	280	193	642
St. Louis, Mo.	Ŭ	Ŭ	U	Ŭ	Ū	Ū	Ŭ			-,	_,				
St. Paul, Minn.	61	42	17	1	-	1	7								
Wichita, Kans.	U	U	U	U	U	U	U								

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 <sup>1</sup> 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: 2002-733-100/69044 Region IV