



#### MORBIDITY AND MORTALITY WEEKLY REPORT

- 793 Knowledge About Folic Acid and Use of Multivitamins Containing Folic Acid Among Reproductive-Aged Women — Georgia, 1995
- **796** Bull Riding-Related Brain and Spinal Cord Injuries Louisiana, 1994–1995
- 798 Human Ehrlichiosis Maryland, 1994802 Asthma Surveillance Programs
  - in Public Health Departments United States
- 804 Notice to Readers

## Knowledge About Folic Acid and Use of Multivitamins Containing Folic Acid Among Reproductive-Aged Women — Georgia, 1995

Neural tube defects (NTDs) are serious birth defects that affect an estimated 4000 pregnancies each year in the United States (1). However, women can substantially decrease the risk for this birth defect by consuming 400  $\mu$ g (0.4 mg) of folic acid per day before conception and during early pregnancy. In September 1992, the Public Health Service (PHS) recommended that all women of childbearing age who are capable of becoming pregnant consume 400  $\mu$ g of folic acid daily (2). To characterize knowledge about the benefits of folic acid and use of multivitamins containing folic acid among Georgia women, the Division of Public Health, Georgia Department of Human Resources (GDHR), analyzed data from the 1995 Georgia Women's Health Survey (GWHS)—a comprehensive study of women's health that included questions about folic acid. This report summarizes the survey findings regarding knowledge and use of folic acid, which indicate that only 20% of Georgia women aged 15–44 years consumed a multivitamin containing ≥400  $\mu$ g of folic acid per day, and 71% did not know that folic acid can prevent some birth defects.

GDHR conducted the GWHS during January–July 1995. GWHS was a random-digit–dialed telephone survey of a probability sample of 4005 Georgia women aged 15–44 years; 3130 (78%) women responded (3). Data for households with more than one eligible woman or multiple residential phone numbers were weighted to adjust for the unequal probability of selection. The sample was highly representative of all childbearing-aged women in Georgia (3).

Survey respondents were asked, "During the past 30 days, how often have you taken multivitamins?"; responses were "every day," "several times a week," "once a week," "less than once a week," and "don't know." Respondents also were asked "What brand of multivitamins do you or did you take most often?" and "Have you heard or read that taking a vitamin called folic acid can help prevent some birth defects?" The amount of folic acid women consumed was estimated based on the amount in the multivitamin brand they reported using.

Overall, 20% (95% confidence interval [CI]=19%–21%) of respondents reported consuming a multivitamin containing ≥400 μg of folic acid per day, 5% (95% CI=4%–6%) reported consuming a multivitamin containing ≥400 μg of folic acid several times a week, and 29% (95% CI=27%–30%) reported they had heard folic acid can help

Folic Acid — Continued

prevent some birth defects. Of those who had heard folic acid can help prevent some birth defects, 30% (95% Cl=27%–32%) reported consuming a multivitamin containing  $\geq$ 400  $\mu g$  of folic acid per day, and 6% (95% Cl=5%–8%) reported consuming a multivitamin containing  $\geq$ 400  $\mu g$  of folic acid several times a week. Of the 71% (95% Cl=70%–73%) who had not heard about folic acid, 16% (95% Cl=15%–18%) reported consuming a multivitamin containing  $\geq$ 400  $\mu g$  of folic acid per day, and 4% (95% Cl=3%–5%) reported consuming a multivitamin containing  $\geq$ 400  $\mu g$  of folic acid several times a week.

Prevalence of knowledge about folic acid varied directly by respondents' educational and income levels. Women with a college degree were more likely to have heard about folic acid than were those with only some high school (45% [95% Cl=41%–49%] versus 12% [95% Cl=9%–15%]), and women with incomes above 150% of poverty level were more likely than women with incomes below 150% of poverty level (31% [95% Cl=29%–33%] versus 18% [95% Cl=15%–21%]).\* Women with higher educational levels were more likely to consume a multivitamin containing ≥400 μg of folic acid per day than were less educated women (some high school education [10% (95% Cl=7%–13%)], high school diploma [20% (95% Cl=17%–23%)], some college education [23% (95% Cl=20%–25%)], and college or postgraduate degree [27% (95% Cl=24%–30%)]), and women with incomes above 150% of poverty level were more likely than women with incomes below 150% of poverty level (22% [95% Cl=20%–23%] versus 14% [95% Cl=11%–17%]).

For each educational level, women who reported knowledge of folic acid were more likely to have consumed a multivitamin containing ≥400 μg of folic acid per day than women who had not heard about folic acid. Among women who had heard about folic acid, the prevalence of consuming a multivitamin containing ≥400 μg per day was 16% (95% Cl=8%–24%) for those with some high school education; 32% (95% Cl=26%–38%), with a high school diploma; 32% (95% Cl=27%–37%), with some college education; and 29% (95% Cl=25%–34%), with a college degree.

Reported by: F Serbanescu, MD, R Rochat, MD, Office of Perinatal Epidemiology, Epidemiology and Prevention Br, V Floyd, MD, Family Health Br, KE Toomey, MD, State Epidemiologist, Div of Public Health, Georgia Dept of Human Resources. Birth Defects and Genetic Diseases Br, Div of Birth Defects and Developmental Disabilities, National Center for Environmental Health; Pregnancy and Infant Health Br, and Behavioral Epidemiology and Demographic Research Br, Div of Reproductive Health, National Center for Chronic Disease Prevention and Health Promotion. CDC.

**Editorial Note:** The findings in this report are subject to at least two limitations. First, folic acid consumption in the GWHS was measured on the basis of reported use of multivitamins only; no information was obtained about consumption of folic acid tablets or foods fortified with folic acid. Second, 22% of the sample did not participate in the survey, and the survey excluded households without telephones; therefore, prevalences of knowledge and use of folic acid may be overestimated.

In 1986 and 1995, nationwide surveys estimated that 20% and 25% of U.S. women, respectively, reported consuming a multivitamin containing  $\geq$ 400  $\mu$ g of folic acid per day (4,5); in South Carolina, 12% of the women who gave birth during October 1992–September 1994 reported consuming a multivitamin containing  $\geq$ 400  $\mu$ g per day (6).

<sup>\*</sup>Poverty statistics are based on a definition originated by the Social Security Administration in 1964, that was subsequently modified by federal interagency committees in 1969 and 1980, and prescribed by the Office of Management and Budget as the standard to be used by federal agencies for statistical purposes.

#### Folic Acid — Continued

These studies and the GWHS findings underscore that 75%–88% of the 60 million women of reproductive age in the United States may not obtain the amount of folic acid recommended by PHS to reduce the risk for spina bifida and other NTDs. In addition, GWHS and a recent survey by the March of Dimes (5) indicate a substantial percentage of reproductive-aged women remain unaware of the potential benefits of folic acid despite publication of the PHS recommendation in 1992.

The results of the survey in Georgia underscore the need for continuing efforts to increase consumption of and awareness about the benefits of folic acid among women of childbearing age. Convenient approaches for ensuring that women obtain adequate amounts of folic acid to reduce the risk for NTDs include daily consumption of either a vitamin supplement or a fortified breakfast cereal containing 400  $\mu g$  of folic acid. In March 1996, the Food and Drug Administration (FDA) required many enriched foods (e.g., most flours, corn meals, pasta, and rice) to be fortified with 140  $\mu g$  of folic acid per 100 g of cereal grains by January 1, 1998 (7); this mandate will increase daily consumption of folic acid on average by 100  $\mu g$ . FDA also issued a regulation that permits the labels of products containing sufficient amounts of folate to claim the products may reduce the risk for having a pregnancy with NTDs (8). The use of health claims on folic acid-containing products and folate-rich foods (e.g., orange juice and green leafy vegetables) will assist in increasing awareness about the benefits of folic acid.

Because women who know about the benefits of folic acid are more likely to consume daily a multivitamin containing 400  $\mu$ g of folic acid, the design and implementation of health education programs for women of childbearing age will be important in educating them about these benefits at the earliest possible time before they become pregnant.

#### References

- 1. CDC. Surveillance for an encephaly and spina bifida and the impact of prenatal diagnosis— United States, 1985–1994. In: CDC surveillance summaries (August 25). MMWR 1995;44 (no. SS-4).
- 2. CDC. Recommendations for the use of folic acid to reduce the number of cases of spina bifida and other neural tube defects. MMWR 1992;41(no. RR-14).
- 3. Serbanescu F, Rochat R. Georgia Women's Health Survey, 1995: preliminary report, September 1996. Atlanta, Georgia: Georgia Department of Human Resources, Division of Public Health, Epidemiology and Prevention Branch, Office of Perinatal Epidemiology (in press).
- 4. Moss AJ, Levy AS, Kim I, Park YK. Use of vitamin and mineral supplements in the United States: current users, types of products, and nutrients. Hyattsville, Maryland: US Department of Health and Human Services, Public Health Service, CDC, National Center for Health Statistics, 1989.
- 5. CDC. Knowledge and use of folic acid by women of childbearing age—United States, 1995. MMWR 1995;44:716–8.
- 6. CDC. Prevention program for reducing risk for neural tube defects—South Carolina, 1992–1994. MMWR 1995;44:141–2.
- 7. Food and Drug Administration. Food standards: amendment of standards of identity for enriched grain products to require addition of folic acid. Federal Register 1996;61:8781–97.
- 8. Food and Drug Administration. Food labeling: health claims and label statements—folate and neural tube defects. Federal Register 1996;61:8752–80.

# Bull Riding-Related Brain and Spinal Cord Injuries — Louisiana, 1994–1995

Rodeos are popular sporting events in the southern and western United States, and bull riders sustain 37% of all rodeo-related injuries—more than participants in any other rodeo event (1,2). During 1994–1995 in Louisiana, five cases of central nervous system trauma associated with riding bulls in rodeo events were identified through the Louisiana Central Nervous System Injury Registry, a statewide, population-based surveillance system addressing brain and spinal cord injury incidence, etiology, and outcome. To further characterize these injury events, the Office of Public Health, Louisiana Department of Health and Hospitals, conducted chart reviews and follow-up telephone interviews with the five injured persons or their parents and interviewed rodeo organizations about rules, regulations, and membership. This report summarizes the investigations of these five cases and recommends use of protective equipment to reduce the risk for such injuries.

In November 1995, the Louisiana division of the National High School Rodeo Association (NHSRA) listed 67 high school students who were registered to compete as bull riders in Louisiana (F. Hinton, Louisiana division, NHSRA, personal communication, November 1995). Because other rodeo associations exist and riders frequently have membership in multiple associations, the number of bull riders cannot be accurately estimated.

Case 1. A 28-year-old man with 15 years' riding experience was thrown to the ground while riding a bull and suffered a fracture of the fifth and sixth cervical vertebrae and an incomplete\* spinal cord injury. He had not been wearing any protective equipment (i.e., mouth guard, helmet, or protective vest). Emergency medical service (EMS) was not present at the event; the time between the call for an ambulance and its arrival was 45 minutes. He was hospitalized for 9 days; at discharge from acute care, he was unable to function independently in activities of daily living (e.g., eating, dressing, and walking) and was considered to have a severe disability. He had impaired movement below the level of the injury.

Case 2. A 14-year-old boy who had ridden a bull three times previously was thrown to the ground while riding; he struck his head and was then trampled by the bull. He sustained a brain stem contusion and an incomplete C2 spinal cord injury and was unconscious for 16 days. No information was available about the use of protective equipment or EMS response. He remained in a persistent vegetative state (i.e., dependent and no meaningful responsiveness) on discharge from the reporting acutecare facility 24 days after he was injured.

**Case 3.** A 26-year-old man with 2 years' riding experience struck his head against a bull's head while riding. He sustained a concussion with brief loss of consciousness, multiple facial bone fractures, and a trimalleolar fracture of his leg. He was wearing a protective vest. EMS was not present; the patient was transported to a hospital in a private vehicle by a family member and was in acute care for 2 days. He recovered with no reported functional limitations.

**Case 4.** A 15-year-old boy with 2 years' riding experience was thrown from and then trampled by a bull. He sustained an incomplete T10–T11 spinal cord injury, multiple rib

<sup>\*</sup>A spinal cord injury resulting in any preserved motor or sensory function below the level of the injury.

Bull Riding-Related Injuries — Continued

fractures, a tension pneumothorax, and a splenic injury. He was not wearing protective equipment. The time between the EMS call and arrival was 10 minutes. Although at the time of discharge from acute care 17 days after he was injured he was reported to have no major deficits, he is no longer able to do heavy manual labor or compete in athletic events.

**Case 5**. A 17-year-old boy with 3 years' riding experience struck his head against a bull's head while riding. He sustained a brain injury and multiple nasal fractures and was unconscious for 5 days. He was not wearing protective equipment. EMS was present at the rodeo. After 40 days in acute care, he had pronounced cognitive and behavioral impairments.

Reported by: LI Gibbs, MPH, DW Lawrence, MPH, BA Reilley, Disability Prevention Program, Injury Research and Prevention Section, Office of Public Health, Louisiana Dept of Health and Hospitals. Div of Acute Care, Rehabilitation Research, and Disability Programs, National Center for Injury Prevention and Control, CDC.

**Editorial Note**: In competitive bull riding, the rider holds with one hand a length of braided rope wrapped around the bull's midsection. The rope is not tied in any way; only the force of the rider's grip on the rope keeps the rider on the bull. Riders must remain on the bull for 8 seconds, during which their free hand cannot touch the bull, themselves, or the rope (3,4). Because riders and bulls are matched by random draw, injuries are more likely to occur when a younger, less experienced rider draws a high-spirited bull. Bull-riding schools for experienced riders exist but are not widely used. For developing basic skills, riders practice on mechanical bulls, calves or young steers, and barrels suspended from ropes (K. Henry, Professional Bull Riders Association [PBR], personal communication, January 1996), although mechanical bull riding also has been associated with injuries (5).

The findings in this report document severe bull riding-associated brain and spinal cord injuries and permanent disability among young males. The number of such injuries may increase directly with the popularity of rodeo sports—from July 1992 to July 1995, membership in the Louisiana division of the NHSRA increased 47% (F. Hinton, NHSRA, personal communication, November 1995).

Protective head gear designed for bull riding has not been developed or recommended by rodeo organizations. Protective vests designed for bull riding are required for youth competition but not for professional competition (3,4,6). Use of protective head gear recommended to prevent horseback-riding-associated traumatic brain injuries (7) may decrease the risk for brain injury in bull riding but has not been assessed for that use. Potential barriers to using protective equipment include cost and a perception that some protective equipment detracts from the desired rugged, western appearance (K. Henry, PBR, personal communication, January 1996; T. Corfield, National Intercollegiate Rodeo Association [NIRA], personal communication, November 1995).

Timely transport by EMS providers to definitive care should decrease the severity and improve the outcome of injuries (8). EMS availability depends on which rodeo organization, if any, sponsors the event. For example, the Professional Rodeo Cowboys Association requires the onsite presence of an emergency medical technician and an ambulance; if the ambulance leaves to transport an injured rodeo participant, the rodeo is to be suspended until another ambulance arrives (4). Rodeos sponsored by college and high school associations require the presence at all times of an

Bull Riding-Related Injuries — Continued

emergency medical technician with a suitable conveyance (3; T. Corfield, NIRA, personal communication, November 1995). At least three of the five injuries described in this report occurred at nonsanctioned rodeos.

The cases described in this report indicate the need for assessing the effectiveness of existing equipment, recommendations for its use in bull riding, and the need for new equipment; graduated competition; and matching the bulls with the skill levels of riders. To reduce the impact of injuries, adequate emergency medical care and transportation should be required for all rodeo events. The Louisiana Office of Public Health is working with the Louisiana Sports Medicine and Safety Advisory Committee (a group initially formed in 1990 to address spinal cord injuries among high school football players), the Tulane Institute of Sports Medicine, the Louisiana Sports Medicine Alliance, and the Louisiana High School Rodeo Association to increase participant awareness of the risk for injury related to bull riding and to develop prevention strategies.

#### References

- Justin Sportsmedicine Program. Ten year injury report. Grapevine, Texas: Justin Sportsmedicine Program, 1995:1–9.
- 2. Griffin R, Peterson KD, Halseth JR, Reynolds B. Injuries in professional rodeo: an update. Physician and Sportsmedicine 1987;15:105–15.
- 3. National High School Rodeo Association. Rules, constitution and by-laws, 1995–1996. Denver, Colorado: National High School Rodeo Association, 1995:72–5.
- 4. Professional Rodeo Cowboys Association. Rulebook. Colorado Springs, Colorado: Professional Rodeo Cowboys Association, 1995:R10.6.1–R10.6.7.
- 5. McConnell RY, Rush GA. Mechanical bull syndrome. South Med J 1982;75:681-6.
- 6. National Little Britches Rodeo Association. Official rule book, 1995–1996. Colorado Springs, Colorado: National Little Britches Rodeo Association, 1995:58–9.
- 7. Anonymous. American Academy of Pediatrics Committee on Sports Medicine and Fitness: horseback-riding and head injuries. Pediatrics 1992;89:512.
- 8. McSwain NE Jr. Emergency medical services. In: McSwain NE Jr, Kerstein MD. Evaluation and management of trauma. Norwalk, Connecticut: Appleton-Century-Crofts, 1987:43–53.

## Human Ehrlichiosis — Maryland, 1994

Ehrlichiosis is an emerging tickborne infectious disease caused by obligate intracellular, gram-negative rickettsia that infect leukocytes. Human monocytic ehrlichiosis (HME) is caused by *Ehrlichia chaffeensis* and is believed to be transmitted by *Amblyomma americanum* (the Lone Star tick). Most HME cases have been reported in southeastern and south-central states. During May–July 1994, five cases of serologically confirmed HME were identified among residents of Maryland. All five persons lived near the Chesapeake Bay and had antecedent histories of tick exposure. This report summarizes the clinical and epidemiologic features of these cases and the results of serologic testing at CDC of specimens from Maryland residents with suspected tickborne infection.

Case 1. On May 17, 1994, a 35-year-old man had onset of fever, headache, malaise, fatigue, myalgia, and back pain. His illness progressed to include anorexia, nausea, vomiting, diarrhea, and a nonproductive cough. On May 22, he was admitted to a hospital for evaluation with a white blood cell count (WBC) of 7.2 X 10<sup>6</sup>/L (with

63% neutrophils and 20% band forms) and a temperature of 100.3 F (37.9 C), and rales were noted in the right lung base. Other laboratory abnormalities included thrombocytopenia (platelet count: 118 X 10<sup>6</sup>/L [normal: 150–400 X 10<sup>6</sup>/L]) and elevated aspartate aminotransferase (AST) (87 IU/L [normal: 8-20 IU/L]) and lactate dehydrogenase (LDH) (303 IU/L [normal: 45–90 IU/L]). The patient's hospital course included persistent fever despite intravenous (IV) treatment with a third-generation cephalosporin, hypotension, and progressive confusion and somnolence. Bacterial cultures of blood, cerebrospinal fluid (CSF), and stool and serologic tests, including an antibody titer for E. chaffeensis, were negative. He was placed in intensive care for pharmacologic support of his blood pressure. Analysis of CSF indicated lymphocytic pleocytosis. Because he did not improve within 48 hours, the antibiotic regimen was empirically changed to IV ciprofloxacin and doxycycline, and symptoms began to resolve within 24 hours. He was discharged on May 30. A serum specimen obtained at discharge was positive for E. chaffeensis antibody by immunofluorescent assay (IFA) (titer of 1:4096). The patient reported a history of extensive tick exposure associated with his job as a surveyor and at his residence on a farm in Kent County.

Case 2. On June 3, 1994, a 41-year-old man had onset of fever, chills, severe headache, malaise, fatigue, myalgia, and back pain. His illness progressed during the next week, and he was evaluated as an outpatient. On June 10, he was admitted to a hospital because of continuing fever and progression of symptoms. Physical examination was normal except for a temperature of 101 F (38.3 C). Laboratory tests included a WBC of 3.4 X 10<sup>6</sup>/L (with 12% atypical lymphocytes); AST, 268 IU/L; LDH, 517 IU/L; alkaline phosphatase (AP), 150 IU/L (normal: 20-70 IU/L); 1+ protein, ketones, and bilirubin in the urine; and CSF lymphocytic pleocytosis. An initial serologic test for E. chaffeensis antibodies and other infectious agents and bacterial cultures of blood were negative. Because the patient's physician was aware of case 1 and recognized clinical similarities to that case, E. chaffeensis infection was suspected, and he was treated with IV ciprofloxacin and doxycycline. Although the patient's fever resolved in 3 days, headache, myalgia, and lethargy persisted. He was discharged on June 16. Analysis of a serologic specimen obtained at discharge detected a titer to E. chaffeensis of >1:1024; a follow-up titer to E. chaffeensis obtained 2 months after the onset of his illness was <1:16. Diplopia attributed to a palsy of the sixth cranial nerve developed late in the course of illness but subsequently resolved. The patient reported frequent exposure to ticks in the vicinity of his residence in a small town and while hiking and biking in the neighboring woods of Kent County.

Case 3. In July 1994, a 45-year-old construction worker who lived near Annapolis and worked in Aberdeen had gradual onset of fatigue, fever, headache, myalgia, and malaise. He sought care from his physician on July 20 and received trimethoprim-sulfamethoxazole for suspected sinusitis. However, he developed nausea, vomiting, diarrhea, and jaundice, and on July 27 his physician prescribed doxycycline and obtained a serum sample for Lyme disease (LD) serology (antibody to *Borrellia burgdorferi*). On about August 1, the physician notified the patient to discontinue the doxycycline because his LD test was negative (titer <1:75). On August 8, the patient was hospitalized because of continuation and progression of his symptoms. Clinical and laboratory findings included an elevated temperature, petechial rash, leukopenia, thrombocytopenia, and modestly elevated levels of serum alanine aminotransferase (ALT) and AST. IV doxycycline and cefotaxime were initiated for treatment of the

unexplained fever and severe headache. When analysis of CSF, an abdominal ultrasound, and a computerized axial tomography of the brain were normal, the cefotaxime was discontinued. Analysis of a blood specimen obtained August 10 included an indeterminate IFA for Rocky Mountain spotted fever (RMSF) and an *E. chaffeensis* titer of 1:1024. Symptoms began to resolve within 3–4 days after initiation of IV doxycycline, and monocytic inclusion bodies were detected in a peripheral blood smear obtained August 15. The patient reported that on some days he removed 25–30 ticks from his clothes and that 2 weeks before onset of symptoms, he removed a partially engorged tick from his hip approximately 36 hours after attachment.

Case 4. On July 27, 1994, a 63-year-old woman began a camping trip to Virginia, North Carolina, South Carolina, and Tennessee. On August 6, she removed an engorged tick attached to her back, which she believed had become attached 24-48 hours earlier during a hike in the mountains of eastern Tennessee. On August 8, she had onset of a backache followed by fever, headache, myalgia, abdominal pain, fatigue, and confusion. She was admitted to a hospital in Maryland on August 15 because of progression of her symptoms. Laboratory abnormalities on admission included pancytopenia—which progressed over a 24-hour period to a WBC of 2.6 X 10<sup>6</sup>/L, a red blood cell count of 3.5 X 10<sup>6</sup>/L, and a platelet count of 88 X 10<sup>6</sup>/L—and increased levels of AP (245 IU/L) and AST (201 IU/L). Atypical pneumonia and hepatitis were suspected, and IV erythromycin was initiated. IV doxycycline subsequently was added to the regimen when a consulting physician suspected RMSF or ehrlichiosis. Because of persistent abdominal pain and tenderness with mildly elevated bilirubin, ALT, and AST, an ultrasonogram of the gall bladder was performed. The wall appeared thickened, and on August 16 she underwent a cholecystectomy; complications included extensive bleeding. Analysis of a blood sample obtained August 15 was negative for ehrlichiosis and RMSF; however, an IFA titer to E. chaffeensis was 1:1024 in a sample obtained August 23. Administration of doxycycline was continued, and she was discharged on September 11.

Case 5. On June 20, 1994, a 38-year-old man who worked at a golf course had onset of fatigue, "a feverish feeling," myalgia, arthralgia, mild headache, and generalized weakness. On June 23, he was examined by a physician who diagnosed atrial fibrillation; neutropenia (1.2 X 10<sup>6</sup>/L) with a lymphocytosis (59%) was detected. Digoxin was initiated for treatment of atrial fibrillation. On June 27, he was hospitalized to evaluate his persistent fever. Findings included a temperature of 104 F (40 C), headache, facial flushing, generalized mild lymphadenopathy, and enlarged erythematous tonsils. Although his WBC had increased to 4.2 X 10<sup>6</sup>/L, his platelet count had decreased (from 153 X 10<sup>6</sup>/L to 100 X 10<sup>6</sup>/L), and liver enzymes were slightly elevated (AST: 70 IU/L and ALT: 72 IU/L). Treatment with gentamicin and piperacillin was initiated for fever of uncertain origin. However, because rickettsial disease infection was suspected, on June 30 treatment was changed to include ampicillin and doxycycline. His clinical condition improved markedly within 48 hours. An IFA of a serum specimen obtained June 30 indicated a titer to E. chaffeensis of ≥1:512, and an enzyme immunoassay for LD indicated a titer of 1:435. The patient reported removing nonengorged ticks from his body approximately 2–3 weeks before the onset of his illness.

Serologic testing. The Shore Health Laboratory in eastern Maryland saved frozen aliquots of serum specimens from 91 patients submitted for RMSF serology by physicians practicing on the eastern shore of Maryland during 1993 and 1994. CDC

performed IFAs for *Rickettsia rickettsii* and *E. chaffeensis* antibodies on these specimens. Of the 12 persons who provided both acute- and convalescent-phase specimens, one was positive for *R. rickettsii* and two for *E. chaffeensis*; of the latter two, one had at least an eightfold increase in IFA titer, and the other had titers of 1:256 and 1:512 on serum samples drawn 6 weeks apart. Of the 79 patients with one blood specimen, no samples were positive for *R. rickettsii*; however, 11 (14%) had titers to *E. chaffeensis* of ≥1:128, which is considered to be consistent with recent infection.

Reported by: LE Silvers, DVM, S Watkins, GT Strickland, MD, School of Medicine, Univ of Maryland, Baltimore; M Clothier, J Grant, MD, Kent County Health Dept, Chestertown; E Hall, DVM, L Joe, MD, Anne Arundel County Health Dept, Annapolis; MA Thompson, Queen Anne's County Health Dept, Centreville; S Sullivan, MS, Talbot County Health Dept, Easton; J Ryan, MD, Queen Anne's and Talbot county health depts; P Shanahan, MD, CG Baumann, MD, Chestertown; M Shochet, MD, Glen Burnie; G Sprouse, MD, Chester; JT Nevins, MS, H McQuay, Shore Health Laboratories, Easton; E Israel, MD, DM Dwyer, MD, State Epidemiologist, Maryland Dept of Health and Mental Hygiene. Viral and Rickettsial Zoonoses Br, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases, CDC.

**Editorial Note**: The findings of this investigation and results of IFAs for *E. chaffeensis* conducted by CDC on serum specimens from Maryland residents indicate that cases of HME occurred in Maryland at least as early as 1988 and that the incidence of HME may be increasing (Table 1). In addition, these findings are consistent with other reports indicating that the incidence of HME is equal to or greater than that of RMSF (1–3). Although no cases of human granulocytic ehrlichiosis (HGE) have been confirmed in Maryland, the clinical features of HGE are identical to those of HME (4), and its suspected vector, *Ixodes scapularis*, is present throughout the eastern and central part of the state. The IFAs for HME and HGE usually do not crossreact, and each test must be performed independently if ehrlichiosis is suspected in patients potentially exposed in areas where both vectors are present (4,5).

The cases described in this report underscore that serologic testing for ehrlichiosis often is negative during the acute phase of infection. Therefore, therapy with a tetracycline antibiotic or with chloramphenical should be initiated based on clinical suspicion before the diagnosis is serologically confirmed (1–3). The responses of the cases in Maryland are consistent with previous reports (1,2), which indicate that IV therapy with large doses of third-generation cephalosporins—a practice often used for treating fevers of unknown origin—is not effective for treating ehrlichiosis, and

TABLE 1. Results of immunofluorescent assays for *Ehrlichia chaffeensis* antibody conducted by CDC on serum specimens, by year — Maryland, 1985–1994

Year	Negative	Positive*	Total
1985	2	0	2
1986	1	0	1
1987	0	0	0
1988	10	2	12
1989	11	1	12
1990	10	0	10
1991	10	0	10
1992	14	1	15
1993	8	1	9
1994	27	8 <sup>†</sup>	35

<sup>\*</sup>Titers ≥1:128.

<sup>&</sup>lt;sup>†</sup>Includes all five cases described in this report.

treatment with doxycycline generally is associated with clinical improvement within 24–48 hours.

The cases in Maryland also reflect the spectrum of illness caused by HME. HME, HGE, and RMSF should be considered in the differential diagnosis of febrile patients with generalized illness who reside, work, or vacation in tick-endemic areas and who have histories of tick exposure (1–3). These tickborne infections may be associated with thrombocytopenia, elevated hepatic enzymes, and CSF pleocytosis, and should be included in the differential diagnosis of patients with suspected influenza, viral hepatitis, aseptic meningitis, and cholecystitis.

Because HME is transmitted by ticks, persons who work outdoors, participate in outdoor activities, or reside in tick-endemic areas should take precautions to reduce tick exposures. These include wearing long pants and pulling socks over the pants cuffs when walking in woods or grassy areas, using insect repellent, and carefully checking for and removing ticks found on clothing and skin.

The CDC surveillance case definition for ehrlichiosis requires a clinically compatible history with a minimum antibody titer of ≥1:64 or a fourfold or greater change in antibody titers to *E. chaffeensis* using the IFA. Serum samples from persons with clinically suspected cases should be sent to CDC through the state health department or, in Maryland, to the Shore Health Laboratory at the Easton Memorial Hospital or to the Clinical Microbiology Laboratory at the Johns Hopkins Medical Systems in Baltimore.

#### References

- 1. Fishbein DB, Kemp A, Dawson JE, Greene NR, Redus MA, Fields DH. Human ehrlichiosis: prospective active surveillance in febrile hospitalized patients. J Infect Dis 1989;160:803–9.
- 2. Harkess JR, Ewing SA, Crutcher JM, Kudlac J, McKee G, Istre GR. Human ehrlichiosis in Oklahoma. J Infect Dis 1989;159:576–9.
- 3. Eng TR, Harkess JR, Fishbein DB, et al. Epidemiologic, clinical, and laboratory findings of human ehrlichiosis in the United States, 1988. JAMA 1990;264:2251–8.
- 4. Walker DH, Dumler JS. Emergence of ehrlichioses as human health problems. Emerging Infectious Diseases 1996;2:18–29.
- 5. Dawson JE, Fishbein DB, Eng TR, Redus MA, Greene NR. Diagnosis of human ehrlichiosis with the indirect fluorescent antibody test: kinetics and specificity. J Infect Dis 1990;162:91–5.

# Asthma Surveillance Programs in Public Health Departments — United States

Although asthma affects more than 14 million persons in the United States (1,2), there have been no nationally coordinated efforts to assist state health departments in developing asthma surveillance programs. To characterize asthma surveillance and control programs in public health departments in the United States, during March and April 1996, the Council of State and Territorial Epidemiologists and CDC conducted a survey of state and territorial epidemiologists. This report presents the results of that survey, which indicate that most states lack the funding and data necessary to develop asthma surveillance programs.

Questionnaires were sent to the 54 state and territorial epidemiologists who were asked to identify the appropriate person to respond to questions about asthma programs in the state. Responses were received from 48 states and three territories. Of the 51 respondents, 43 reported no state- or territorial-level asthma-control program.

Asthma Surveillance — Continued

Based on a priority ranking scale with five items suggesting reasons states might not have an asthma-control program, the two most important reasons included lack of funds and shortage of staff. In an open-ended response, 10 states reported that asthma was not a public health priority in their state. However, 37 (86%) of the 43 states/territories expressed an interest in starting an asthma-control program.

Potential data available for characterizing asthma included hospital discharge records (42 [82%]), emergency department visits (16 [31%]), use of public or private health-care services for asthma care (10 [20%]), first-time visitors to a health-care provider (four [8%]), and survey data about the quality of life for persons with asthma (four [8%]). Only Wisconsin maintained a surveillance system to monitor trends in asthma.

Of the 42 states/territories with hospital discharge data, 14 previously had analyzed the data for asthma morbidity. Reasons for inability to use hospital discharge data included restricted access to the data because of legislative constraints and incompatible data formats.

Although no state or territory maintains an asthma-control program, 26 state health departments have been associated with efforts to control asthma in selected communities in their state, including environmental control measures (22), public education (14), patient education (14), education of health-care providers (12), and legislation (five).

Reported by: HA Anderson, MD, WR Forrester, MPA, DM Perrotta, PhD, Council of State and Territorial Epidemiologists, Atlanta. Air Pollution and Respiratory Health Br, Div of Environmental Hazards and Health Effects, National Center for Environmental Health, CDC.

**Editorial Note**: During 1985–1990, the estimated medical costs of asthma care in the United States increased from \$4.5 billion to \$6.2 billion, and in 1985 these costs represented approximately 1% of total U.S. health-care costs (3). In the United States, asthma is the most common chronic disease of childhood and affects approximately 5 million children aged <18 years; asthma is the fourth leading cause of disability in children (4). Rates for asthma prevalence, hospitalization, and death are highest among children residing in inner cities, and important risk factors for asthma-related mortality include being poor or black (1,5,6).

National health objectives for the year 2000 regarding asthma prevention are to establish and monitor state-based plans to define and track sentinel respiratory diseases triggered by environmental factors, reduce hospitalizations, reduce the proportion of persons with activity limitations, and increase the proportion of persons with asthma that get formal patient education (objectives 11.16, 11.1, 17.4, and 17.14b) (7).

The findings in this report indicate that states lack the funding, staff, and data necessary to develop asthma surveillance programs. Although 84% of respondents reported the availability of hospital discharge data, most state and territorial health departments have not used the data because of barriers to its access such as negotiating its use with a private entity, legal barriers, or incompatible data systems. Other potential sources for obtaining state-specific data about asthma include adding state-specific questions about asthma to the Behavioral Risk Factor Surveillance System, designating asthma a performance measure in the Health Plan and Employer Data and Information Set (HEDIS), and monitoring Medicaid data over time.

#### Asthma Surveillance — Continued

Despite the need for state-specific data and the need to develop surveillance systems to monitor trends in asthma, approximately half of the responding health departments have been associated with efforts to reduce the impact of asthma in selected communities in their state. State and territorial health departments need to determine the local burden of asthma and should explore approaches for eliminating barriers that prevent the use of existing data. Collaboration between CDC and other federal agencies, managed-care organizations, academic institutions, and states and territories to design and implement comprehensive community-based asthma surveillance systems will better characterize the burden of asthma in the United States and will enable states to target areas where asthma-prevention programs should be implemented.

#### References

- 1. CDC. Asthma mortality and hospitalization among children and young adults—United States, 1980–1993. MMWR 1996;45:350–3.
- 2. Adams PF, Marano MA. Current estimates from the National Health Interview Survey, 1994. Vital Health Stat 1995;10:94.
- 3. Weiss KB, Gergen PJ, Hodgson TA. An economic evaluation of asthma in the United States. N Engl J Med 1992;326:862–6.
- 4. CDC. Disabilities among children aged ≤17 years—United States, 1991–1992. MMWR 1995;44: 609–13.
- 5. CDC. Asthma—United States, 1982-1992. MMWR 1995;43:952-5.
- 6. Weiss KB, Wagener DK. Changing patterns of asthma mortality. JAMA 1990;264:1683-7.
- 7. Public Health Service. Healthy people 2000: national health promotion and disease prevention objectives. Washington, DC: US Department of Health and Human Services, Public Health Service, 1991; DHHS publication no. (PHS)91-50213.

#### Notice to Readers

### Satellite Videoconference on HIV/AIDS Prevention for Teens

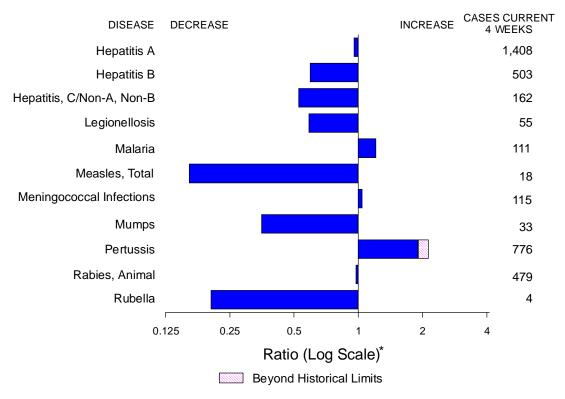
"HIV/AIDS Prevention for Teens," a satellite videoconference, will be broadcast live to sites nationwide from the Massachusetts Corporation for Educational Telecommunications (MCET) through a cooperative agreement with CDC on December 12, 1996, from 3 p.m. to 4:30 p.m. eastern standard time. The course is aimed at teachers of students in grades 6–12, health educators, community leaders, counselors, and administrators. Participants will receive an overview of HIV/AIDS education and guidance in targeting prevention strategies for youth.

Additional information, registration forms, and coordinates for down-link sites are available from MCET, telephone (800) 556-4376. The deadline to register down-link sites and participants is November 5.

### Erratum: Vol. 45, No. 34

In the article "HIV Testing Among Women Aged 18–44 Years—United States, 1991 and 1993" in the third paragraph, on page 734, the first sentence should read, "From 1991 to 1993, the proportion of women aged 18–44 years who had ever been tested for HIV increased 69% (from 18.8% to 31.8%) (Table 1)."

FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending September 14, 1996, with historical data — United States



<sup>\*</sup>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 — provisional cases of selected notifiable diseases, United States, cumulative, week ending September 14, 1996 (37th Week)

	Cum. 1996		Cum. 1996
Anthrax Brucellosis Cholera Congenital rubella syndrome Cryptosporidiosis* Diphtheria Encephalitis: California* eastern equine* St. Louis* western equine* Hansen Disease Hantavirus pulmonary syndrome*†	60 2 1 1,392 1 44 1 - 73	HIV infection, pediatric*§ Plague Poliomyelitis, paralytic¶ Psittacosis Rabies, human Rocky Mountain spotted fever (RMSF) Streptococcal toxic-shock syndrome* Syphilis, congenital** Tetanus Toxic-shock syndrome Trichinosis Typhoid fever	195 1 28 1 487 13 225 20 99 15

<sup>-:</sup> no reported cases

<sup>-:</sup> no reported cases

\*Not notifiable in all states.

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

§ Updated monthly to the Division of HIV/AIDS Prevention, National Center for HIV, STD, and TB Prevention (NCHSTP), last update August 27, 1996.

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

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

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending September 14, 1996, and September 16, 1995 (37th Week)

	AIDS*		AIDS* Chlamydia		richia 157:H7 PHLIS <sup>§</sup>	Gono	rrhea		atitis A,NB	Legionellosis	
Reporting Area	Cum. 1996	Cum. 1995	Cum. 1996	NETSS <sup>†</sup> Cum. 1996	Cum. 1996	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995
UNITED STATES	45,416	50,257	257,673	1,722	908	200,026	277,220	2,361	2,795	597	842
NEW ENGLAND	1,849	2,388	12,017	243	55	5,067	5,354	81	95	34	21
Maine N.H.	31 58	75 75	635 397	20 28	30	38 80	66 81	- 7	12	2 2	5 1
Vt.	14	21	U	16	15	42	44	28	9	3	-
Mass. R.I.	873 123	999 179	4,796 1,387	121 10	10 -	1,573 369	1,881 364	40 6	69 5	18 9	12 3
Conn.	750	1,039	4,802	48	-	2,965	2,918	-	-	Ň	Ň
MID. ATLANTIC	12,627	13,055	31,096	150	38	23,096	31,484	205	324	149	140
Upstate N.Y. N.Y. City	1,672 7,052	1,707 6,555	N 15,097	106 8	12 -	4,393 7,762	6,730 12,543	161 1	158 1	53 5	38 4
N.J.	2,402	3,090	3,286	36	5	3,649	3,163	-	134	9	21
Pa.	1,501	1,703	12,713	N	21	7,292	9,048	43	31	82	77
E.N. CENTRAL Ohio	3,616 810	3,791 808	44,210 13,640	422 108	284 57	30,113 10,058	55,251 17,258	328 25	226 8	156 67	252 119
Ind.	462	379	7,080	62	39	4,589	6,448	7	2	34	58
III. Mich.	1,579 570	1,521	17,415	179 73	84 56	12,555	14,099	52 244	67 149	9 33	22 23
Wis.	195	816 267	U 6,075	/3 N	48	U 2,911	12,691 4,755	244	149	33 13	30
W.N. CENTRAL	1,060	1,179	20,020	383	197	8,827	14,479	90	63	33	55
Minn. Iowa	189 69	242 68	2,702 2,866	163 86	115 55	U 741	2,176 1,090	1 40	2 12	3 9	2 17
Mo.	541	559	2,000 8,767	47	-	5,865	8,159	31	17	6	17
N. Dak.	10	4	2	12	13	100	21	-	5	-	3
S. Dak. Nebr.	9 74	14 80	721 1,801	13 33	3	102 670	141 855	- 5	1 14	2 10	1 12
Kans.	168	212	3,161	29	11	1,449	2,037	13	12	3	7
S. ATLANTIC	11,216	12,603	38,190	93	51	67,754	76,530	182	171	101	138
Del. Md.	215 1,324	239 1,621	1,148 4,812	- N	1 7	1,039 10,018	1,565 9,108	- 1	- 7	10 20	2 24
D.C.	799	740	N	-	-	3,144	3,161	-	-	8	4
Va. W. Va.	795 83	961 83	7,881 1	N N	22 2	6,550 365	7,845 497	10 9	10 41	13 1	18 3
N.C.	603	712	ΰ	23	12	12,727	16,888	34	45	7	30
S.C. Ga.	586	673	7,990	8 27	7	8,038	8,668	21 U	16	4 3	28 14
Ga. Fla.	1,651 5,160	1,639 5,935	16,358	27 25	-	13,243 12,630	14,193 14,605	107	15 37	35	15
E.S. CENTRAL	1,563	1,614	21,070	40	37	22,499	28,900	424	756	36	48
Ky. Tenn.	272 580	196 665	4,709 9,309	8 18	4 30	2,982 8,198	3,368 9,810	21 323	24 730	3 18	9 23
Ala.	431	410	5,969	9	3	9,530	12,007	323 4	2	3	23 6
Miss.	280	343	U	5	-	1,789	3,715	76	U	12	10
W.S. CENTRAL	4,562	4,589	30,539	38	10 3	22,697	38,836	331	211	18 2	17
Ark. La.	186 1,046	209 713	4,962	11 5	3 4	2,489 5,336	3,736 8,035	7 142	5 131	1	5 2
Okla.	189	206	5,463	8	1	3,497	3,887	69	34	5	4
Tex. MOUNTAIN	3,141 1,325	3,461 1,515	20,114 11,688	14 140	2 68	11,375 5,127	23,178 6,760	113 418	41 334	10 29	6 89
Mont.	23	1,515	11,000	140	-	24	51	14	11	1	4
Idaho	29 3	37	1,106	27	10	80	107	92	43	-	2
Wyo. Colo.	362	12 493	410	8 53	2 31	25 1,077	39 2,093	134 40	133 51	3 7	8 33
N. Mex.	118	123	2,705	8	-	576	754	54	39	1	4
Ariz. Utah	370 127	392 111	4,768 1,108	N 19	17 -	2,589 213	2,608 173	53 22	32 10	13 2	9 12
Nev.	293	331	1,591	11	8	543	935	9	15	2	17
PACIFIC	7,597	9,523	48,843	213	168	14,846	19,626	302	615	41	82
Wash. Oreg.	508 339	664 324	6,588 U	65 62	71 36	1,437 412	1,885 560	41 6	156 33	5	19
Calif.	6,594	8,292	36,799	83	52	12,446	16,278	106	394	32	58
Alaska Hawaii	23 133	53 190	832 872	3 N	2 7	296 255	479 424	3 146	1 31	1 3	- 5
Guam	4	190	168	N	-	31	81	146	5	2	5 1
P.R.	1,524	1,828	N	13	U	210	426	77	170	-	-
V.I. Amer. Samoa	17	27	N	N N	U U	-	19	-	-	-	-
C.N.M.I.	1	-	N	N	Ü	11	45	-	5	-	-

N: Not notifiable U: Unavailable

<sup>-:</sup> no reported cases

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

<sup>\*</sup>Updated monthly to the Division of HIV/AIDS Prevention, National Center for HIV, STD, and TB Prevention, last update August 27, 1996.

†National Electronic Telecommunications System for Surveillance.

§Public Health Laboratory Information System.

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending September 14, 1996, and September 16, 1995 (37th Week)

	Lyı Dise		Mal	aria	Mening@ Dise			hilis Secondary)	Tubero	culosis	Rabies,	Animal
Reporting Area	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995
UNITED STATES	8,108	7,888	969	903	2,344	2,221	7,547	11,727	13,372	14,565	4,287	5,598
NEW ENGLAND Maine	2,676 24	1,573 16	39 7	37 5	97 12	102 7	120	269 2	292 4	356 11	517 70	1,127 21
N.H. Vt.	29 15	19 8	2	1 1	3	18 6	1	1	9 1	9 2	48 117	115 136
Mass.	205	94	12	11	37	36	57	46	152	200	85	339
R.I. Conn.	388 2,015	251 1,185	6 10	4 15	10 32	4 31	1 61	3 217	24 102	35 99	33 164	247 269
MID. ATLANTIC	4,572	5,156	233	247	202	280	295	610	2,418	3,095	519	1,463
Upstate N.Y. N.Y. City	2,673 189	2,618 342	59 113	48 135	63 30	76 38	51 94	63 264	291 1,239	359 1,768	264	867 -
N.J. Pa.	571 1,139	1,367 829	46 15	47 17	53 56	70 96	77 73	126 157	499 389	524 444	101 154	260 336
E.N. CENTRAL	54	347	99	121	328	315	933	2,015	1,433	1,390	74	80
Ohio Ind.	35 17	23 13	9	9 15	124 51	90 46	336 160	629 241	211 120	195 128	11 5	10 12
III.	2	16	35	64	87	83	313	788	766	705	18	13
Mich. Wis.	Ū	5 290	30 11	13 20	34 32	57 39	U 124	203 154	261 75	300 62	27 13	33 12
W.N. CENTRAL	109	74	38	18	192	137	272	579	334	430	404	269
Minn. Iowa	39 18	5 9	17 2	3 2	25 39	23 25	51 15	34 36	78 44	105 48	21 183	14 95
Mo. N. Dak.	22	37	9 1	6 1	79 3	51 1	175	472	144 6	163 3	16 52	25 24
S. Dak.	-	-	-	1	9	5			15	15	103	73
Nebr. Kans.	2 28	4 19	3 6	3 2	17 20	12 20	12 19	11 26	13 34	20 76	3 26	5 33
S. ATLANTIC	485	513	215	171	487	364	2,660	2,941	2,448	2,584	1,977	1,498
Del. Md.	78 274	37 343	3 58	1 46	2 53	6 31	30 464	10 326	20 213	43 291	52 451	74 307
D.C. Va.	3 32	2 40	7 32	15 38	10 44	4 48	109 300	77 454	98 201	71 167	9 417	11 294
W. Va.	11	21	3	2	11	8	3	9	45	54	76	88
N.C. S.C.	58 4	44 14	20 9	15 1	60 46	64 47	715 293	815 438	329 254	316 226	508 69	354 100
Ga. Fla.	1 24	9 3	23 60	23 30	118 143	72 84	479 267	550 262	448 840	478 938	217 178	200 70
E.S. CENTRAL	51	52	23	20	133	148	1,686	2,397	1,230	1,021	153	212
Ky. Tenn.	12 17	12 20	3 11	2 7	21 16	36 54	100 594	130 630	172 297	218 323	34 56	22 72
Ala. Miss.	6 16	7 13	3	8 3	56 40	29 29	406 586	473	596	296 184	60 3	111 7
W.S. CENTRAL	84	83	22	38	274	268	1,118	1,164 2,318	165 1,578	1,941	285	526
Ark. La.	21 1	7 4	- 4	2	30 47	26 39	121 381	354 743	127 59	146 188	15 13	33 24
Okla.	13	35	-	1	27	28	139	141	134	146	23	28
Tex. MOUNTAIN	49 6	37 7	18 44	31 43	170 132	175 161	477 109	1,080 166	1,258 416	1,461 446	U 108	441 130
Mont.	-	-	6	3	4	2	-	4	14	10	19	38
Idaho Wyo.	2	3	4	1 -	19 3	8 7	4 2	-	6 5	11 1	23	1 23
Colo. N. Mex.	- 1	- 1	20 2	18 4	28 22	40 30	23 1	92 5	54 55	38 60	30 5	9
Ariz.	-	-	6	7	34	47	66	32	177	224	25	37
Utah Nev.	2 1	1 2	4 2	5 5	12 10	13 14	2 11	4 29	39 66	19 83	3 3	11 6
PACIFIC	71	83	256	208	499	446	354	432	3,223	3,302	250	293
Wash. Oreg.	12 11	8 13	17 17	16 13	77 87	73 80	5 10	11 18	182 75	187 84	6	9 1
Calif. Alaska	47	62	212 3	167 2	326 6	283 6	338	402 1	2,793 48	2,850 52	236 8	276 7
Hawaii	1	-	7	10	3	4	1	-	125	129	-	-
Guam P.R.	-	-	-	1 1	1 5	2 18	3 97	8 200	35 63	83 120	- 32	- 35
V.I. Amer. Samoa	-	-	-	2	-	-	-	-	-	3	-	-
C.N.M.I.	-	-	-	1	-	-	1	5	-	30	-	

N: Not notifiable

U: Unavailable

-: no reported cases

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending September 14, 1996, and September 16, 1995 (37th Week)

-	H. influ		Сертен	Hepatitis (vir	al), by type			Measles	(Rubeol	a)	
		sive		A	E		Ind	igenous	Imported <sup>†</sup>		
Reporting Area	Cum. 1996*	Cum. 1995	Cum. 1996	Cum. 1995	Cum. 1996	Cum. 1995	1996	Cum. 1996	1996	Cum. 1996	
UNITED STATES	808	837	18,744	20,405	6,639	7,045	-	397	3	39	
NEW ENGLAND	22	32	255	196	140	171	-	10	-	4	
Maine N.H.	8	3 8	14 12	21 8	2 10	7 17	-	-	-	-	
Vt. Mass.	1 11	2 10	6 134	5 80	10 44	5 64	-	1 8	-	1 3	
R.I.	2	3	13	25	9	8	-	-	-	-	
Conn.	-	6	76	57	65	70	-	1	-	-	
MID. ATLANTIC Upstate N.Y.	131 41	120 33	1,131 309	1,248 296	963 248	1,000 272	-	23	-	5 -	
N.Y. City N.J.	25 40	28 14	399 245	615 173	425 184	315 260	-	9 3	-	3	
Pa.	25	45	178	164	106	153	-	11	-	2	
E.N. CENTRAL	126	144	1,597 583	2,344 1,309	705 95	796 82	-	5	3 3	7	
Ohio Ind.	77 7	73 18	232	128	120	150	-	2	-	3 -	
III. Mich.	30 7	35 16	352 311	484 266	173 270	207 301	-	2	-	1 3	
Wis.	5	2	119	157	47	56	-	1	-	-	
W.N. CENTRAL Minn.	41 25	62 34	1,639 94	1,405 141	323 41	465 43	-	21 16	-	2 2	
lowa	5	3	265	63	66	34	-	-	-	-	
Mo. N. Dak.	7 -	18 -	762 80	1,008 22	153 2	325 4	-	4	-	-	
S. Dak.	1 1	1	41	37	4 31	2	-	-	-	-	
Nebr. Kans.	2	3 3	159 238	38 96	26	23 34	-	1	-	-	
S. ATLANTIC	185	166	900	796	1,042	894	-	6	-	9	
Del. Md.	2 47	- 55	12 155	9 155	7 217	6 183	-	1 2	-	2	
D.C. Va.	5 6	- 21	23 121	18 150	28 99	15 82	-	-	-	3	
W. Va.	7	7	13	17	18	40	-	-	-	-	
N.C. S.C.	22 4	25 1	102 42	85 35	253 61	203 37	-	3	-	1 -	
Ga. Fla.	73 19	52 5	90 342	51 276	10 349	62 266	-	-	-	2 1	
E.S. CENTRAL	22	8	999	1,257	594	628	_	2	_	-	
Ky.	4	2	22	35	39	54	-	-	-	-	
Tenn. Ala.	9 8	5	676 139	1,035 64	353 48	495 79	-	2	-	-	
Miss.	1	1	162	123	154	-	U	-	U	-	
W.S. CENTRAL Ark.	31 -	53 5	3,882 369	2,880 375	876 61	948 43	-	26	-	2	
La. Okla.	3 25	1 20	109 1,670	84 719	84 59	152 118	-	-	-	-	
Tex.	3	27	1,734	1,702	672	635	-	26	-	2	
MOUNTAIN	78	92	3,021	2,932	776	596	-	152	-	5	
Mont. Idaho	1	2	89 156	83 244	9 71	19 70	-	1	-	-	
Wyo. Colo.	35 11	5 14	26 330	86 378	33 98	17 87	-	1 4	-	3	
N. Mex.	9	12	292	609	269	226	-	16	-	-	
Ariz. Utah	9 7	22 9	1,246 704	830 533	189 73	88 48	-	8 117	-	2	
Nev.	6	28	178	169	34	41	-	5	-	-	
PACIFIC Wash.	172 2	160 8	5,320 346	7,347 599	1,220 66	1,547 139	-	152 51	-	5 -	
Oreg. Calif.	22 144	22 125	612 4,277	1,907 4,678	50 1,085	92 1,294	-	4 33	-	2	
Alaska	2	1	32	32	10	10	-	63	-	-	
Hawaii	2	4	53	131	9	12	-	1	-	3	
Guam P.R.	1	3	2 80	6 78	261	4 456	U -	6	U -	-	
V.I. Amer. Samoa	-	-	-	6 6	-	14	U U	-	U U	-	
C.N.M.I.	10	11	1	23	5	16	Ü	-	Ŭ	-	

N: Not notifiable

U: Unavailable

-: no reported cases

 $<sup>^{*}</sup>$ Of 188 cases among children aged <5 years, serotype was reported for 42 and of those, 12 were type b.

<sup>&</sup>lt;sup>†</sup>For imported measles, cases include only those resulting from importation from other countries.

TABLE III. (Cont'd.) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending September 14, 1996, and September 16, 1995 (37th Week)

	Measles (Rub		Г	iibci i	0, 1995	(07 ()	TVVCCK	<i>'</i>	T			
		tal		Mump	s		Pertussi	s	Rubella		a	
Reporting Area	Cum. 1996	Cum. 1995	1996	Cum. 1996	Cum. 1995	1996	Cum. 1996	Cum. 1995	1996	Cum. 1996	Cum. 1995	
UNITED STATES	436	268	11	459	613	152	3,335	2,906	-	195	106	
NEW ENGLAND	14	8	-	1	11	8	677	382	-	25	44	
Maine N.H.	-	-	-	-	4 1	-	19 66	22 28	-	-	- 1	
Vt.	2	-	-	-	-	6	55	58	-	2	-	
Mass. R.I.	11 -	2 5	-	1 -	2 1	-	489 25	259 2	-	20	7 -	
Conn.	1	1	-	-	3	2	23	13	-	3	36	
MID. ATLANTIC Upstate N.Y.	28	12 1	-	60 19	92 24	15 14	264 144	237 108	-	8 4	13 3	
N.Y. City	12	5	-	14	13	1	23	35	-	2	8	
N.J. Pa.	3 13	6	-	2 25	14 41	-	11 86	16 78	-	2	2	
E.N. CENTRAL	12	14	3	83	105	21	339	345	_	3	3	
Ohio Ind.	5	1	3	38 6	32 7	7 1	166 33	102 24	-	-	-	
III.	3	2	-	18	31	12	108	67	-	1	-	
Mich. Wis.	3 1	5 6	-	20 1	35 -	1	27 5	56 96	-	2	3	
W.N. CENTRAL	23	2	-	13	37	16	224	178	_	1	-	
Minn. Iowa	18	-	-	5 1	2 9	15	172 9	78 7	-	- 1	-	
Mo.	4	1	-	4	21	1	28	46	-	-	-	
N. Dak. S. Dak.	-	-	-	2	1	-	1 4	8 10	-	-	-	
Nebr.	-	-	-	-	4	-	6	8	-	-	-	
Kans.	1	1	-	1	-	-	4	21	-	- 01	-	
S. ATLANTIC Del.	15 1	11 -	6	81 -	90	33	410 11	237 9	-	91 -	9 -	
Md. D.C.	4	1	-	21	27	12	145	31 5	-	- 1	1	
Va.	3	-	-	12	19	12	55	15	-	2	-	
W. Va. N.C.	4	-	2	- 19	16	-	2 75	84	-	- 77	1	
S.C. Ga.	2	2	- 1	5 3	9	3	29 17	20 18	-	1	-	
Fla.	1	8	3	21	13	6	76	55	-	10	7	
E.S. CENTRAL	2	-	-	19	7	1	68	255	-	2	1	
Ky. Tenn.	2	-	-	1	-	-	26 17	17 203	-	-	1	
Ala.	-	-	- U	3	4 3	1 U	17 8	34	- N	2 N	- N	
Miss. W.S. CENTRAL	28	23	1	15 24	40	2	79	1 229	-	3	7	
Ark.	-	2	1	2	6	2	9	31	-	-	-	
La. Okla.	-	18	-	12 -	9	-	7 8	13 22	-	1 -	-	
Tex.	28	3	-	10	25	-	55	163	-	2	7	
MOUNTAIN Mont.	157 -	68	-	22	26 1	14 8	310 25	467 3	-	6	4	
ldaho	1	-	-	-	2	1	99	88	-	2	-	
Wyo. Colo.	1 7	26	-	2	1	-	5 78	1 69	-	2	-	
N. Mex.	16	31	N	N	N	2	44	79	-	-	-	
Ariz. Utah	8 119	10	-	1 2	2 11	3	23 14	153 18	-	1 -	3 1	
Nev.	5	1	-	17	9	-	22	56	-	1	-	
PACIFIC Wash.	157 51	130 19	1 -	156 18	205 10	42 38	964 451	576 178	-	56 2	25 1	
Oreg.	4	1	-	-	-	2	31	37	-	1	-	
Calif. Alaska	35 63	108 -	-	113 2	176 12	-	458 2	319 -	-	50 -	19 -	
Hawaii	4	2	1	23	7	2	22	42	-	3	5	
Guam P.R.	6	3	U	5 1	3 2	U	1 1	2 1	U	-	1	
V.I.	-	-	U	-	3	U	-	-	U	-	-	
Amer. Samoa C.N.M.I.	-	-	U U	-	-	U U	-	-	U U	-	-	

N: Not notifiable

U: Unavailable

-: no reported cases

TABLE IV. Deaths in 121 U.S. cities,\* week ending September 14, 1996 (37th Week)

	All Causes, By Age (Years)						P&I <sup>†</sup>		All Cau	ıses, B	/ Age (Y	ears)		P&I <sup>†</sup>	
Reporting Area	All Ages	>65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	>65	45-64	25-44	1-24	<1	Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn. Cambridge, Mass. Fall River, Mass. Hartford, Conn. Lowell, Mass. Lynn, Mass. New Bedford, Mass. New Haven, Conn. Providence, R.I. Somerville, Mass. Springfield, Mass. Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J.	551 126 36 9 25 44 18 12 s. 24	414 91 26 7 7 22 30 14 7 7 20 23 60 4 36 27 47 1,464 34 25 65 22 23	69 15 7 1 2 8 1 2 1 6 7 1 8 1 9 444 8 9 105	42 15 2 1 4 1 3 2 4 4 1 2 - 3 2 15 5 3 4 4 1 2 4 4 1 2 1 5 3 3 4 4 4 4 1 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	15 3 - 1 2 2 - 1 2 - 2 5 5 0 - 1 1 3 3 - 1	11 2 1 - - 3 1 2 2 30 3 3	22 3 1 1 1 1 2 - 1 4 4 - 1 1 3 - - - - - - - - - - - - - - - -	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla. Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, Fla. Tampa, Fla. Washington, D.C. Wilmington, Del. E.S. CENTRAL Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala.	1,393 151 289 107 117 106 54 83 41 56 183 177 29 640 98	843 82 172 70 64 34 48 25 42 1199 18 415 63 29 41 40 65 56		162 20 43 14 13 10 4 10 3 1 12 32 5 5 1 8 3 4 10 8 3	41 6 13 1 1 2 5 1 1 - 7 4 - 20 3 - 3 2 5 3	43 6 10 1 3 1 4 1 3 4 5 5 -	66 2 19 10 2 4 4 2 4 2 16 5 7 9 10
Erie, Pa.§ Jersey City, N.J. New York City, N.Y. Newark, N.J. Paterson, N.J. Philadelphia, Pa. Pittsburgh, Pa.§ Reading, Pa. Rochester, N.Y. Schenectady, N.Y. Scranton, Pa.§ Syracuse, N.Y. Trenton, N.J. Utica, N.Y. Yonkers, N.Y. E.N. CENTRAL	39 35	33 25 732 23 12 201 43 7 84 16 21 64 10 19 25	6 5 250 16 7 62 14 2 12 1 4 20 4 2	139 14 22 4 1 4 1 6 2 1 2	22 3 12 1 1 2 - 1 4 -	1 10 2 - 3 1 1 2 - 2 1 - 1	2 3 47 1 18 3 4 4 2 1 4 1 2 5	Nashville, Tenn.  W.S. CENTRAL Austin, Tex. Baton Rouge, La. Corpus Christi, Tex. Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La. San Antonio, Tex. Shreveport, La. Tulsa, Okla.	155 1,475 70 33	963 38 22 32 104 61 71 210 44 83 158 44 96	41 288 18 3 10 28 17 23 78 14 28 44 10 15	13 144 10 6 3 16 6 13 41 6 15 15 8 78	4 44 2 1 8 5 2 11 5 3 1 6	1 34 2 2 6 2 7 10 1 1 3	7 81 4 3 1 6 1 6 33 2 10 7 8
Akron, Ohio Canton, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Gary, Ind. Grand Rapids, Micl Indianapolis, Ind. Madison, Wis. Milwaukee, Wis. Peoria, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	58 34 391 167 137 199 123 214 43 57 136 U 124 34 59 97 63 738 70 29 19 87 87 87 87 87 87 87 87 87 87	1,342 244 229 1222 88 141 88 120 34 35 50 40 72 28 48 49 49 48 49 19 10 10 10 10 10 10 10 10 10 10 10 10 10	8 8 8 9 2 8 8 2 8 3 2 0 5 9 6 1 6 U 9 2 6 U 2 1 5 9 9 9 1 2 1 4 1 3 5 5 6 4 4 4 8 2 5 1 5 7 1 2	1032 422 112 115 115 115 115 115 115 115 115 1	33 1 12 2 3 7 3 7 · 2 U · 3 U 3 · 3 · 5 2 2 7 1 · 1 4 · 6 2 5 5 3	3 1 1 17 3 6 6 6 3 5 5 - - - U 3 3 3 U 1 1 2 2 3 3 - - - - - - - - - - - - - - - -	1543166513U53U635452 5664 4343651	Albuquerque, N.M. Colo. Springs, Colo Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, Utah Tucson, Ariz. PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawaii Long Beach, Calif. Los Angeles, Calif. Pasadena, Calif. Portland, Oreg. Sacramento, Calif. San Diego, Calif. San Francisco, Calif. San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash.	94 97 119 21 184 192 143 1,598 16 61 9 62 69 516 31 144 U	66 373 76 15 115 57 100 1,100 5 500 47 339 24 97 U 91 20 93 46 U	15 71 15 30 4 32 3 15 31 284 5 15 28 11 100 31 31 31 4 5 7 0 7 0	12 8 11 6 1 15 15 9 135 2 2 1 7 52 4 11 U 13 15 12 2 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 - 2 - 6 - 1 - 1 - 3 - 4 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1	4 6 6 - 7 7 - 2 3 3 3 3 2 3 3 9 1 - U 8 3 3 5 - 2 U 257	1 35 8 1 12 3 10 108 2 3 1 11 9 4 7 U 17 11 17 1 6 U

U: Unavailable -: no reported cases

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

†Pneumonia and influenza.

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

Total includes unknown ages.

## Contributors to the Production of the MMWR (Weekly)

## Weekly Notifiable Disease Morbidity Data and 121 Cities Mortality Data

Denise Koo, M.D., M.P.H.

Deborah A. Adams

Timothy M. Copeland

Patsy A. Hall

Carol M. Knowles

Sarah H. Landis

Myra A. Montalbano

# **Desktop Publishing and Graphics Support**

Jolene W. Altman

Morie M. Higgins

Peter M. Jenkins

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 on Friday of each week, send an e-mail message to lists@list.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/ or from CDC's file transfer protocol server at ftp.cdc.gov. 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 (404) 332-4555.

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.

Director, Centers for Disease Control and Prevention David Satcher, M.D., Ph.D. Deputy Director, Centers for Disease Control and Prevention Claire V. Broome, M.D.

Director, Epidemiology Program Office Stephen B. Thacker, M.D., M.Sc.

Editor, MMWR Series Richard A. Goodman, M.D., M.P.H. Managing Editor, MMWR (weekly) Karen L. Foster, M.A. Writers-Editors, MMWR (weekly) David C. Johnson Darlene D. Rumph Person Caran R. Wilbanks Editorial Assistant, MMWR (weekly) Teresa F. Rutledge

☆U.S. Government Printing Office: 1996-733-175/47027 Region IV