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Multistate Outbreak of Listeriosis — United States, 2000

Since May 2000, 29 illnesses caused by a strain of *Listeria monocytogenes* (LM) have been identified in 10 states: New York (15 cases); Georgia (three); Connecticut, Ohio, and Michigan (two each); and California, Pennsylvania, Tennessee, Utah, and Wisconsin (one each). Dates of LM isolation ranged from May 17 through November 26 with 26 (90%) infections occurring since July 15. When subtyped, the LM isolates from these cases were indistinguishable by pulsed-field gel electrophoresis (PulseNet pattern numbers GX6A16.0014 by *Asc*1 and GX6A12.0017 by *Apa*1) and ribotyping (DUP-1053). This report summarizes the investigation, which linked these cases of listeriosis to eating deli turkey meat.

Eight perinatal and 21 nonperinatal cases were reported. Among the 21 nonperinatal case-patients, the median age was 65 years (range: 29–92 years); 13 (62%) were female. The 29 cases have been associated with four deaths and three miscarriages/stillbirths.

A case-control study conducted by five state and two local health departments and CDC implicated eating deli turkey meat as the probable source of infection. Thirteen (76%) of 17 case-patients and five (21%) of 24 controls ate deli turkey meat during the 30 days before illness onset (Mantel-Haenszel weighted odds ratio=8.0; 95% confidence interval=1.2–43.3). State health and agriculture departments investigated 13 stores and delicatessens where 11 patients reported purchasing turkey; these stores and delicatessens carried turkey meat produced by at least 27 federally inspected establishments. Two establishments were linked to 10 of 11 patients; one of these establishments produced turkey meat for the second establishment.

On December 8, investigators from the Food Safety and Inspection Service, U.S. Department of Agriculture (USDA) began investigating the implicated establishments. On December 12, Cargill Turkey Products, Inc. (Waco, Texas) stopped shipping ready-to-eat foods and, on December 14, voluntarily recalled processed turkey and chicken delimeat that might have been contaminated.

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Editorial Note: LM infection causes an estimated 2500 serious illnesses and 500 deaths in the United States each year. Infected pregnant women may experience only a mild, influenza-like illness; however, infections during pregnancy can lead to premature delivery, miscarriage, stillbirth, or serious infection of the newborn. Other persons at increased risk for infection are those aged ≥65 years, persons with cancer, diabetes, kidney disease, acquired immunodeficiency syndrome, or who take immunosuppressive medications. Manifestations of illness include meningitis and sepsis. Healthy persons aged <65 years rarely are affected.

The risk for a person developing *Listeria* infection after eating a contaminated product is very small. Persons who have eaten a recalled product but do not have symptoms do not require tests or treatment even if they are in a high-risk group. However, persons in a high-risk group who have eaten contaminated product and become ill within 2 months with fever or signs of serious illness should consult a physician.

Guidelines for preventing listeriosis are similar to those for preventing other foodborne illnesses. The general recommendations are 1) cook thoroughly raw food from animal sources (e.g., beef, pork, or poultry); 2) wash raw vegetables thoroughly before eating; 3) keep uncooked meats separate from vegetables and from cooked foods and ready-to-eat foods; 4) avoid raw (unpasteurized) milk or foods made from raw milk; and 5) wash hands, knives, and cutting boards after each handling of uncooked foods. Persons at high risk for listeriosis may choose to 1) avoid soft cheeses (i.e., feta, Brie, Camembert, blue-veined, and Mexican-style cheese such as queso fresco). Hard cheeses, processed cheeses, cream cheese, cottage cheese, or yogurt need not be avoided; 2) cook leftover foods or ready-to-eat foods (e.g., hot dogs) until steaming hot; and 3) avoid foods from deli counters (e.g., prepared salads, meats, and cheeses) or thoroughly reheat cold cuts before eating.

Cases of listeriosis with onset since October 1, 2000, should be reported to state and local health departments; information about the recall is available at http://www.fsis.usda.gov/OA/recalls/rec_actv.htm*. Consumers who have recalled meat products, even if they have been stored in freezers, should discard or return them to the point of purchase. High-risk consumers who have processed turkey or chicken deli meat but are uncertain of the brand should call the place of purchase to find out if it might be a recalled product, or discard it. Answers to meat-safety questions are available at the USDA meat and poultry hotline, (800) 535-4555. Listeriosis information is available at http://www.cdc.gov/ncidod/dbmd/diseaseinfo/listeriosis_g.htm.

^{*}References to sites of non-CDC organizations on the World-Wide Web 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 pages found at these sites.

Foodborne Outbreak of Group A Rotavirus Gastroenteritis Among College Students — District of Columbia, March-April 2000

On March 31, student health services at a university in the District of Columbia (DC) notified the DC health department that an increased number of students had become ill with acute gastroenteritis beginning March 29. Some ill students reported eating tuna or chicken salad sandwiches from dining hall A on campus. On March 31, the DC health department initiated an outbreak investigation. This report summarizes results of the investigation, which indicated that group A rotavirus transmitted by food was the cause of the outbreak.

Telephone interviews were conducted with students who reported illness to student health services, with additional ill students who were identified during interviews, and with healthy controls selected randomly from the university registry of students residing on campus. A case of gastroenteritis was defined as three or more episodes of diarrhea and/or two or more episodes of vomiting within a 24-hour period in a student with onset on or after March 20. Controls and case-patients whose illness onset occurred during March 27–31 were questioned about food history, residence and dining hall, source of water, use of a public access computer or sports equipment at the university gym, and attendance at social or athletic events. Electronic records of student meal attendance were available for 49 case-patients with illness onset during March 27–31 and for 55 control subjects.

Twenty-three (79%) of 29 employees of dining hall A were interviewed to identify their work duties and determine whether they were ill. Stool specimens were collected during March 29–April 10 from six ill students and 21 dining hall A employees. Samples were screened for bacterial and parasitic pathogens at a commercial laboratory and for viral pathogens at CDC.

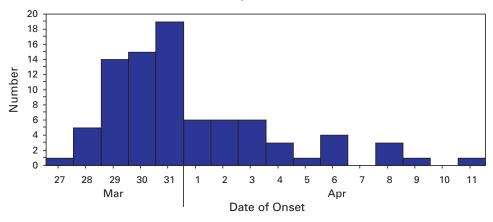
The outbreak among students began March 27 and peaked at 19 cases on March 31 (Figure 1). A total of 108 students (55 were identified by telephone interviews and 53 were self-reported) had gastrointestinal symptoms during March 26–April 11; 85 (79%) had illness that met the case definition. The attack rate among students residing on campus was 5% (77 of 1641), with no significant differences in attack rates by sex, occupancy of residence hall, or grade level. Eight case-patients resided off campus (attack rate: 0.02%). Among the 83 case-patients for whom a complete list of symptoms was reported, 77 (93%) had diarrhea, 75 (90%) abdominal pain or discomfort, 69 (83%) loss of appetite, 67 (81%) nausea, 64 (77%) fatigue, 56 (67%) vomiting, 49 (59%) headache, 48 (58%) chills, 48 (58%) subjective or low-grade fever, and 42 (51%) myalgia. Sore throat, cough, and/or congestion were reported by six case-patients with onsets on or after April 2. The median duration of illness was 4 days (range: 1–8 days). Nine (11%) case-patients received intravenous fluids to treat dehydration.

Of those who completed the telephone interview, 40 (91%) of 44 case-patients and 27 (68%) of 40 controls ate at least one deli sandwich from campus dining hall A during March 27–30 (p=0.017; odds ratio [OR]=4.8; 95% confidence interval [CI]=1.3–22.1). During March 27–30, four (8%) of 49 case-patients ate four or more meals at dining hall B compared with 18 (33%) of 55 controls (p=0.005; OR=0.2; 95% CI=0.04–0.6). Food histories of employees were not recorded; however, six employees reported illness.

Stool specimens of students and employees were negative for bacterial and parasitic pathogens and for Norwalk-like viruses. Using electron microscopy, enzyme immunoassay, and reverse transcriptase-polymerase chain reaction (RT-PCR), nine (33%) of 27

Rotavirus Gastroenteritis — Continued

FIGURE 1. Number* of gastroenteritis¹ cases among college students, by date of illness onset — District of Columbia, March 27–April 11, 2000



^{*} n=85.

specimens were positive for group A rotavirus. Rotavirus positive stool specimens from four students and three employees were identified as genotype combination P[4],G2 by RT-PCR. Two of the three P[4],G2-positive employees were line cooks who reported having symptoms of gastroenteritis on March 27 and April 2, respectively, while the third positive employee, a deli server, reported no illness.

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Editorial Note: Group A rotavirus is the most common cause of childhood diarrhea worldwide, infecting >90% of children by age 3 years (1). Because rotavirus immunity develops early in life, disease among older children and adults is uncommon (1). Although the role of rotavirus in diarrhea outbreaks in adults has not been well studied, it has been documented as the cause of adult diarrheal outbreaks in hospitals (2), nursing homes (3), isolated communities (4), and in travelers (5). Also, parents of children infected with rotavirus have been reported to experience acute gastroenteritis (6). However, the rotavirus G and P protein-type combinations, the proteins that elicit an immune response in humans, were not characterized in most of these reports.

The rapid increase and gradual decline of the campus outbreak suggest that the infection was foodborne during the first week and was spread person-to-person during the following week. During the first week, illness was associated with eating sandwiches at dining hall A and was associated inversely with eating frequently at dining hall B. The employee who prepared sandwich fillings did not report illness and tested negative for rotavirus. None of the three deli servers who assembled and served sandwiches reported illness; however, one was rotavirus P[4],G2 positive. It is unknown whether the deli server who tested positive was infected before the outbreak among students.

[†] A case of gastroenteritis was defined having three or more episodes of diarrhea and/or two or more episodes of vomiting within a 24-hour period in a student with onset on or after March 20.

Rotavirus Gastroenteritis — Continued

This rotavirus serotype G2 outbreak was unusual for two reasons; food was implicated as the source of infection and the adults affected should have been immune. During April 2000, a gastroenteritis outbreak among adults in Japan also was caused by foodborne transmission of group A rotavirus serotype G2 (7). These adults should not have been susceptible to severe rotavirus illness. G2 strains often are found combined with serotype P[4]1B (8). The G and P neutralization antigens of serotype G2 strains may allow G2 strains to escape immunity induced by the more common G1, G3, and G4 strains. In addition, G2 has been associated with more severe dehydration during diarrheal episodes in children than other common strains (9). These outbreaks of rotavirus gastroenteritis in adults in the United States and Japan raise questions about the persistence of immunity to rotavirus and the virulence of G2 strains. Investigators and clinicians should consider rotavirus as a possible cause of acute gastroenteritis in adults.

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Blood Lead Levels in Young Children — United States and Selected States, 1996–1999

Lead exposure adversely affects the cognitive development and behavior of young children (1). For children aged <6 years, CDC has defined an elevated blood lead level (BLL) as $\geq 10~\mu g/dL$, but evidence exists for subtle effects at lower levels (2). Data from CDC's Third National Health and Nutrition Examination Survey, Phase 2 (1991–1994) (NHANES) showed that average BLLs in children had decreased approximately 80% since the late 1970s but that elevated BLLs remained more common among low-income children, urban children, and those living in older housing (3,4). Although these data provide national estimates of the prevalence of elevated BLLs among children, they do not provide information at the state or local level. To target prevention efforts and monitor progress toward reducing BLLs at the state and local level, CDC's Childhood Blood Lead Surveillance (CBLS) program supports state blood lead surveillance programs on the basis of blood lead tests from public and private clinical laboratories. This report

summarizes data on BLLs in children aged 1–5 years from NHANES data collected in 1999 and children aged <6 years from state surveillance data provided to CDC by 19 state surveillance programs during 1996–1998. The findings indicate that, despite the decreases in mean BLL among children, the problem remains concentrated on a local level. Surveillance efforts should be used to target screening efforts to communities at highest risk.

NHANES is a continuous survey of the health and nutritional status of the U.S. civilian, noninstitutionalized population designed so that each year of data constitutes a nationally representative sample. Data in this report are from NHANES 1999, and NHANES III, Phase 2. A household interview and a physical examination were conducted for each survey participant. During the physical examination, blood was collected by venipuncture for all persons aged >1 year. Graphite furnace atomic absorption spectrophotometry was used to measure BLLs with detection limits of 0.3 μ g/dL (NHANES 1999) and 1.0 μ g/dL (NHANES III, Phase 2). Long-term quality-control data for these analyses, including similar standardized reference materials, were used in both surveys and showed that data from the two surveys can be compared. Because of limited sample size, NHANES 1999 analyses include only data on average BLLs and selected percentiles but not on the prevalence of elevated levels.

The analyses of CBLS data were based on reports from 19 of 28 states that provided blood lead data to CDC (Table 1). The 19 states were included because they received all blood lead test results of children from participating laboratories (regardless of level) and reported data from January 1, 1996 through December 31, 1998. These states accounted for 33% of all U.S. children aged <6 years.

An elevated BLL from CBLS is defined as a single blood lead test result $\geq 10 \ \mu g/dL$. If multiple tests were reported for a child during a calendar year, the highest BLL measured for that child was used. To estimate the proportion of children with elevated BLLs among those tested, the number of children with elevated levels was divided by the number of children tested at least once during a calendar year.

From NHANES III, Phase 2 (1991–1994) to NHANES 1999, the geometric mean BLL in children aged 1–5 years decreased from 2.7 (95% confidence interval [CI]=2.6–2.9) to 2.0 μ g/dL (95% CI=1.7–2.3), and the 50th percentile decreased from 2.6 (95% CI=2.4–2.8) to 1.9 μ g/dL (95% CI=1.6–2.1). The continued pattern of decline in BLLs between the two surveys also is indicated at the 10th, 25th, 75th, and 90th percentiles.

The CBLS data showed that the proportion of children tested with BLLs \geq 10 µg/dL decreased from 10.5% in 1996 to 7.6% in 1998 in the 19 states providing data (Table 1). The proportions of children with BLLs \geq 15 µg/dL and \geq 20 µg/dL also decreased.

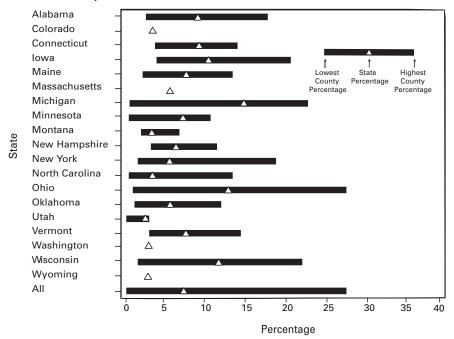
The percentage of children aged <6 years tested with BLLs \geq 10 μ g/dL in each state ranged from 2.7 to 14.9 (Figure 1). Within states, the proportion of children with elevated

TABLE 1. Percentage of children tested aged <6 years with elevated blood lead levels (BLLs), by year — selected states*, 1996–1998

		<u>% Childre</u>	% Children with elevated BLLs (μg/dL)						
Year	No. tested	≥10	≥15	≥20					
1996	1,220,596	10.5%	3.9%	1.9%					
1997	1,183,506	8.6%	3.2%	1.5%					
1998	1,256,907	7.6%	2.7%	1.2%					

^{*} Alabama, Colorado, Connecticut, Iowa, Maine, Massachusetts, Michigan, Minnesota, Montana, New Hampshire, New York, North Carolina, Ohio, Oklahoma, Utah, Vermont, Washington, Wisconsin, and Wyoming.

FIGURE 1. State-specific percentage of children aged <6 years tested with blood lead levels (BLLs) \geq 10 μ g/dL and highest and lowest percentage of elevated BLLs, by county — selected states, 1998*



^{*} Only counties with ≥200 children tested for BLL are included. Colorado, Washington, and Wyoming had <2 counties with 200 children tested, and Massachusetts did not report county of residence.

BLLs in counties with at least 200 children tested also varied considerably. For example, the proportion of children with elevated BLLs ranged from 1.3% to 27.3% in counties in Ohio. Across all 19 states, the county-specific proportions of children with elevated BLLs ranged from 0.5% to 27.3%, indicating a concentrated proportion of elevated BLLs in specific populations or geographic areas.

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Editorial Note: The findings in this report indicate that average BLLs of U.S. children aged 1-5 years have declined from the early 1990s to 1999. Because of the limited sample size of a single year of NHANES 1999 compared with that of the multiple years of NHANES III, additional data are necessary to confirm this trend. The dramatic decline in BLLs from the late 1970s through the early 1990s resulted primarily from the phaseout of leaded gasoline and the resulting decrease in lead emissions, although other exposures also decreased (3). Although air lead levels and lead emissions continued to decrease during the 1990s, most of this decline occurred before 1995 (5). The primary remaining sources of childhood lead exposure are deteriorated leaded paint and the soil and dust it contaminates in old housing. The construction of new housing and the demolition and rehabilitation of older housing may be contributing to a continued decline in BLLs. Data from NHANES III, Phase 2 showed that low-income children living in older housing had more than a 30-fold greater prevalence of BLLs ≥10 μg/dL than do middleincome children in newer housing (4). From 1993 to 1997, the number of low-income children living in pre-1940s and 1940-1974 housing declined by 31% and 14%, respectively. The number of low-income children living in post-1974 housing increased by 5% (6).

Despite the overall decline in average BLLs, CBLS data show that the risk for elevated BLLs in children tested remains high in some counties and varies greatly among and within states. This variation most likely reflects geographic variation in the prevalence of risk factors for elevated BLLs such as residence in older housing and poverty.

The findings in this report are subject to at least four limitations. First, the small NHANES 1999 sample does not permit observing risks in specific subgroups or geographic areas, but it provides a nationally representative estimate of BLLs in children. The CBLS data set provides local information but is limited to children who receive clinical or diagnostic blood lead testing. Second, because CDC guidelines recommend the use of blood lead data and census data to target screening efforts in populations at increased risk for lead exposure, the proportion of children with elevated BLLs is higher in CBLS data than would be expected in NHANES 1999. Third, the guidelines for testing children vary by state, and adherence to the guidelines varies by health-care provider. Finally, CBLS data include samples collected by fingerstick, which can slightly overestimate the blood lead result, and venous samples and results obtained by different laboratories. Despite these differences, the temporal trends in BLLs are similar between the CBLS and NHANES data sets.

One of the national health objectives for 2010 is the elimination of childhood lead poisoning (7). Data in this report document continued progress toward this goal but also show the ongoing need to target prevention efforts at communities and populations at highest risk. CDC recommends that state health agencies target screening efforts by using blood lead surveillance data, census data, Medicaid data, and other sources of information on risk factors such as housing age and poverty (8,9). Other federal agencies, including the U.S. Department of Housing and Urban Development and the U.S.

Environmental Protection Agency, also are implementing targeted strategies to prevent lead exposure. State blood lead surveillance systems play a key role in targeting and monitoring federal, state, and local prevention efforts. CDC encourages additional states to conduct surveillance for elevated BLLs in children.

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Notice to Readers

Public Health Service Recommendations for the Use of Vaccines Manufactured with Bovine-Derived Materials

The Center for Biologics Evaluation and Research (CBER), U.S. Food and Drug Administration (FDA) learned earlier this year that some vaccines were manufactured with bovine-derived materials obtained from countries in which bovine spongiform encephalopathy (BSE) or a substantial risk for BSE exists. A list of these countries is published by the U.S. Department of Agriculture (USDA).* This information was of concern because cases of variant Creutzfeldt-Jakob disease (vCJD) have been attributed to, among other possibilities, eating beef products from cattle infected with the agent of BSE. No evidence exists that cases of vCJD are related to the use of vaccines, and no cases of vCJD have been reported in the United States.

CBER assessed the risk for vCJD from vaccines manufactured with processes that use bovine materials potentially contaminated with the BSE agent. On July 27, 2000, CBER convened a joint meeting of the Transmissible Spongiform Encephalopathy Advisory Committee and the Vaccines and Related Biological Products Advisory Committee to review the results of these assessments and make recommendations about the use and manufacture of these vaccines. The committees concluded that the risk for vCJD

^{*9} CFR, part 94.

posed by vaccines in the scenarios presented was theoretical and remote. This conclusion was based on the inherent low risk of the bovine materials involved (e.g., type and amount of tissue[s] used, specific time and country, or herd of origin) and/or the dilutions of materials during manufacture. The committees concluded that the benefits of vaccination outweigh any remote risks for vCJD.

As a precautionary measure, the committees recommended that vaccines manufactured with bovine-derived materials from countries on the USDA list be replaced with bovine-derived materials from other countries. This recommendation, which is consistent with existing FDA guidance first issued in 1993 on the sourcing of bovine-derived materials, is intended to reduce even the remote risk for vCJD from vaccines. The committees also recommended that FDA provide information to the public about the safety of vaccines made with materials from countries in which BSE or BSE risk exists.

FDA has requested that manufacturers replace bovine-derived materials obtained from countries on the USDA list with materials obtained from countries not on the USDA list. All of the affected manufacturers have agreed to implement these changes or have already done so. FDA anticipates that most of these changes will be completed in 2001.

The Public Health Service (PHS) recommends that all persons continue to be vaccinated according to current schedules. PHS has no preference for using one licensed vaccine product over another based on the source of bovine-derived materials used in vaccine production. Failure to obtain the recommended vaccinations with licensed vaccines poses a risk for serious disease.

Additional information about BSE or vaccines manufactured with bovine-derived materials from countries on the USDA list can be obtained from the FDA World-Wide Web site, http://www.fda.gov/cber/BSE/BSE.htm[†], or from CBER's Office of Communications, Training and Manufacturers Assistance, telephone (800) 835-4709.

Notice to Readers

Availability and Use of Parenteral Quinidine Gluconate for Severe or Complicated Malaria

Since 1991, quinidine gluconate, a class 1a anti-arrhythmic agent, has been the only parenteral antimalarial available for use in the United States (1). It is indicated for the treatment of patients with life-threatening *Plasmodium falciparum* malaria (2), including those who cannot tolerate oral therapy, have high-grade parasitemia, or have complications (e.g., cerebral malaria or acute renal failure) (3,4).

The limited availability of and delays in obtaining quinidine gluconate have contributed to adverse patient outcomes (5–7). As newer anti-arrhythmics have replaced quinidine for many cardiac indications, some hospitals and other health-care facilities have dropped quinidine gluconate from their formularies and, as a result, fewer clinicians have had experience using the drug. Discussions among quinidine gluconate manufacturer Eli Lilly Company (Indianapolis, Indiana), CDC, the U.S. Department of Defense, and the U.S. Food and Drug Administration have resulted in the following recommendations

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to improve quinidine gluconate availability for acutely ill malaria patients in U.S. health-care facilities:

- Before an acute need arises, hospital drug services should consider maintaining or adding quinidine gluconate to formularies or should be able to immediately locate a nearby source.
- Pharmacists and clinicians requiring quinidine gluconate in hospitals in which an immediate source cannot be located should contact their local or regional distributor to request quinidine gluconate.
- 3. In clinical settings in which the need for the drug is more acute than can be met by the local or regional distributor, pharmacists and clinicians should contact Eli Lilly Company, telephone, (800) 821-0538 to arrange a rapid shipment of the drug. This telephone number, or an alternate number given to callers, is staffed 24 hours a day, 7 days a week.
- 4. If further assistance is needed in obtaining quinidine gluconate or in managing patients with malaria, contact CDC's malaria hotline, (770) 488-7788 (Monday–Friday, 8 a.m. to 4:30 p.m. eastern standard time). After business hours, weekends, and holidays, contact CDC's security station, telephone, (404) 639-2888 and ask to have the on-call person for malaria questions paged.

The following dosing recommendations for quinidine gluconate administration are provided for pharmacists and clinicians treating patients with severe or complicated malaria:

- Quinidine gluconate intravenous should be administered in a monitored setting.
 Prolongation of the QT interval as indicated by an electrocardiogram, ventricular arrhythmia, hypotension, and hypoglycemia can result from the use of this drug at treatment doses.
- Quinidine gluconate for malaria is administered as an initial intravenous loading dose of 10 mg/kg salt (equivalent to 6.25 mg/kg quinidine base) infused over 1–2 hours. Quinidine gluconate is administered subsequently as a continuous infusion of 20 μg/kg/min quinidine gluconate salt (equivalent to 12.5 μg/kg/min quinidine base) (2).
- An alternative regimen is an intravenous loading dose of 24 mg/kg quinidine salt (equivalent to 15 mg/kg quinidine base) infused over 4 hours, followed by a maintenance infusion of 12 mg/kg of quinidine gluconate salt (equivalent to 7.5 mg/kg quinidine base) infused over 4 hours every 8 hours, starting 8 hours after the loading dose (2). These regimens have been shown to be effective with or without concomitant exchange transfusion (2).
- The risk for serious ventricular arrhythmia associated with quinidine is increased by bradycardia, hypokalemia, and hypomagnesemia (2). When determining whether a patient should receive a bolus dose, previous administration of other drugs that can prolong the QT interval (e.g., quinine, halofantrine, and mefloquine) should be considered.
- No alternatives to quinidine exist for patients in the United States who require
 intravenous therapy for malaria. Acute cardiac events can be minimized by careful calculation of the loading dose and infusion rate. Consulting a cardiologist may
 be helpful when attempting to resume infusion in the patient who has experienced QT prolongation or hypotension associated with intravenous quinidine
 infusion.
- Consulting a physician with experience in treating malaria is advised.

References

- CDC. Treatment with quinidine gluconate of persons with severe *Plasmodium falciparum* infection: discontinuation of parenteral quinine from CDC drug service. MMWR 1991;40(no. RR-4):21–3.
- Quinidine gluconate injection [package insert]. Indianapolis, Indiana: Eli Lilly Company, February 2000.
- Zucker JR, Campbell CC. Malaria: principles of prevention and treatment. Infect Dis Clin No Am 1993;7:547–67.
- Miller KD, Greenberg AE, Campbell CC. Treatment of severe malaria in the United States with a continuous infusion of quinidine gluconate and exchange transfusion. N Engl J Med 1989;321:65–70.
- Rosenthal PJ, Petersen C, Geertsma FR, et al. Availability of intravenous quinidine for falciparum malaria [Letter]. N Engl J Med 1996;335:138.
- Humar A, Sharma S, Zoutman D, et al. Fatal falciparum malaria in Canadian travelers. Can Med Assoc J 1997;156:1165-7.
- CDC. Availability of parenteral quinidine gluconate for treatment of severe or complicated malaria. MMWR 1996;45:494–5.

Notice to Readers

Availability of MMWR Mirror Website in Spain

CDC, in collaboration with the Toxic Oil Syndrome Research Centre (CISAT) of the Institute of Health Carlos III, Madrid, Spain, has established a *MMWR* mirror website in Spain. The website was developed to reduce the delay caused by transoceanic electronic transfers of large documents and to increase access to information published in *MMWR* for European public health practitioners. The mirror website is updated simultaneously with the posting of new reports on the *MMWR* website (http://www.cdc.gov/mmwr). The address for the CISAT *MMWR* mirror website is http://cisat.isciii.es/mmwr. The website also hosts a mirror site from the Agency for Toxic Substances and Disease Registry (ATSDR). This mirror site can be found at http://cisat1.isciii.es/atsdr. Other features of the website include information on environmental health problems and rare diseases in Spanish.

CISAT is a part of the WHO Collaborating Centre for the Clinical Epidemiology of Environmental Diseases and has established agreements with CDC/ATSDR and the University of Pittsburgh. Support of the *MMWR* mirror website is part of a larger effort undertaken by CISAT to create a comprehensive environmental health information site.

Notice to Readers

Epidemiology in Action: Intermediate Methods

CDC and Emory University's Rollins School of Public Health will co-sponsor a course, "Epidemiology in Action: Intermediate Methods" during February 26–March 2, 2001, at Emory University. The course is designed for state and local public health professionals.

The course will review the fundamentals of descriptive epidemiology and biostatistics, analytic epidemiology and computers as used in epidemiology but will focus on midlevel epidemiologic methods directed at strengthening participants' quantitative skills, with an emphasis on up-to-date data analysis. Topics include field investigations, advanced measures of association, normal and binomial distributions, logistic regression, and additional statistical methods. Prerequisite is an introductory course in epidemiology, such as Epidemiology in Action, International Course in Applied Epidemiology or any other introductory class. There is a tuition charge.

Deadline for applications is January 15. Additional information and applications are available from Emory University, Rollins School of Public Health, International Health Dept. (PIA), 1518 Clifton Road, N.E., Room 746, Atlanta, GA 30322; telephone (404) 727-3485; fax (404) 727-4590; or email pvaleri@sph.emory.edu.

Notice to Readers

Epi Info 2000: A Course for Teachers of Epidemiologic Computing

CDC and Emory University's Rollins School of Public Health will co-sponsor a course, "Epi Info 2000: A Course for Teachers of Epidemiologic Computing" during March 12–15, 2001, at Emory University. The course is designed for teachers of epidemiologic computing with intermediate to advanced skills in computing.

The 4-day course covers hands-on experience with the new Windows® version of Epi Info, programming Epi Info software at the intermediate to advanced level, methods of teaching epidemiologic computing, computerized interactive exercises for teaching epidemiology, and computing. There is a tuition charge.

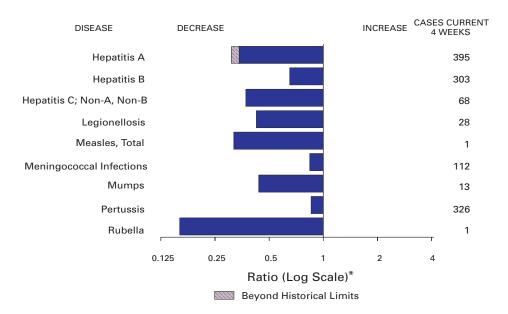
Deadline for applications is February 1. Additional information and applications are available from Emory University, Rollins School of Public Health, International Health Dept. (PIA), 1518 Clifton Road, N.E., Room 746, Atlanta, GA 30322; telephone (404) 727-3485; fax (404) 727-4590; or email pvaleri@sph.emory.edu.

Notice to Readers

Combined Issues of MMWR

A December 29, 2000, issue of *MMWR* will not be published. The next issue will be Volume 49, Numbers 51 and 52, dated January 5, 2001. It will include the figures and tables of notifiable diseases and deaths for the weeks ending December 23, 2000, and December 30, 2000.

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending December 16, 2000, 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 December 16, 2000 (50th Week)

	Cum. 2000		Cum. 2000
Anthrax Brucellosis* Cholera Cyclosporiasis* Diphtheria Ehrlichiosis: human granulocytic (HGE)* human monocytic (HME)* California serogroup viral* eastern equine* St. Louis* western equine*	- 61 2 38 2 178 98 109 2 3	Poliomyelitis, paralytic Psittacosis* Ofever* Rabies, human Rocky Mountain spotted fever (RMSF) Rubella, congenital syndrome Streptococcal disease, invasive, group A Streptococcal toxic-shock syndrome* Syphilis, congenital ¹ Tetanus Toxic-shock syndrome	10 21 2 416 6 2,619 70 257 26
Hansen disease (leprosy)* Hantavirus pulmonary syndrome*† Hemolytic uremic syndrome, postdiarrheal* HIV infection, pediatric** Plague	63 30 185 203 6	Trichinosis Tularemia* Typhoid fever Yellow fever	15 110 311

^{-:} 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).

⁵Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP). Last update November 26, 2000.

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

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

	S Ellulli	9		-, ====,	T	Cellibei	Escherichia coli O157:H7*						
	AID	s	Chlan	nydia⁺	Cryptosp	oridiosis	NET		PH				
Reporting Area	Cum. 2000 [§]	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999			
UNITED STATES	36,091	40,781	623,458	629,947	2,491	2,580	4,311	3,865	3,206	2,690			
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	1,884 38 31 37 1,137 95 546	2,070 75 46 16 1,319 96 518	20,396 1,368 1,004 507 8,586 2,409 6,522	20,437 1,009 944 469 8,588 2,251 7,176	104 20 23 27 30 4	185 31 19 36 71 6 22	383 31 39 36 163 19	401 39 35 32 176 27 92	367 28 35 34 168 18 84	363 - 34 21 187 26 95			
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	7,705 705 3,929 1,592 1,479	10,462 1,196 5,574 1,922 1,770	54,597 N 23,185 8,093 23,319	63,488 N 25,903 11,967 25,618	182 128 11 12 31	597 174 256 52 115	405 296 12 97 N	598 516 17 65 N	281 72 13 109 87	161 14 18 71 58			
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	3,442 546 352 1,693 652 199	2,810 462 317 1,345 552 134	102,284 23,724 12,648 27,498 25,179 13,235	106,620 28,285 11,595 31,316 21,439 13,985	795 260 58 7 96 374	627 66 41 87 52 381	986 272 133 188 137 256	975 250 104 497 124 N	589 220 83 21 104 161	527 219 67 89 83 69			
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	813 160 86 368 3 7 68 121	934 177 75 449 6 15 62 150	34,314 7,129 4,579 10,975 750 1,776 3,343 5,762	36,573 7,238 4,784 12,913 909 1,509 3,410 5,810	356 131 75 33 16 15 77 9	198 75 55 26 18 7 15	687 236 180 103 21 56 63 28	523 167 112 46 17 47 102 32	595 211 147 97 20 58 45 17	543 186 80 68 18 62 113			
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	10,157 199 1,197 785 764 60 667 755 1,117 4,613	11,255 158 1,339 636 777 64 741 917 1,585 5,038	122,481 2,760 12,946 3,067 15,053 1,534 20,654 9,032 25,728 31,707	131,464 2,674 12,533 N 13,392 1,736 21,234 17,998 31,300 30,597	467 6 13 20 18 3 28 - 170 209	373 - 17 7 27 3 34 - 136 149	369 1 32 1 77 15 90 21 42 90	332 6 42 1 75 16 74 20 35	270 1 1 U 61 13 68 14 36 76	188 3 4 U 62 11 52 14 3			
E.S. CENTRAL Ky. Tenn. Ala. Miss.	1,809 186 771 457 395	1,788 256 704 444 384	47,219 7,802 14,457 13,946 11,014	44,413 7,145 13,878 12,276 11,114	49 7 11 16 15	46 7 13 15 11	147 40 61 11 35	141 49 55 28 9	112 32 52 9 19	104 35 44 21 4			
W.S. CENTRAL Ark. La. Okla. Tex.	3,708 172 649 320 2,567	4,159 186 814 125 3,034	96,162 5,355 17,285 8,776 64,746	90,166 5,764 15,863 7,973 60,566	123 14 10 17 82	90 2 24 13 51	182 57 9 19 97	140 15 14 38 73	233 38 53 17 125	164 14 14 30 106			
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	1,322 14 20 9 300 140 427 137 275	1,605 13 22 11 290 82 816 141 230	34,774 1,385 1,816 769 8,490 4,279 12,190 2,150 3,695	31,714 1,496 1,713 757 5,998 4,843 11,799 2,085 3,023	174 10 23 5 72 21 11 28	100 13 8 1 14 42 13 N 9	431 31 74 21 162 23 54 52	328 25 68 16 112 13 36 35 23	283 	242 43 17 88 7 24 48 15			
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	5,251 480 171 4,479 22 99	5,698 336 208 5,047 14 93	111,231 12,606 5,140 88,299 2,343 2,843	105,072 11,612 5,857 82,700 1,817 3,086	241 N 21 220 -	364 N 98 266	721 222 156 298 30 15	427 164 68 180 1	476 200 114 150 1	398 180 69 136 1			
Guam P.R. V.I. Amer. Samoa C.N.M.I.	15 1,245 32 - -	18 1,180 35 - -	3,122 U U U	468 U U U U	- U U U	- U U U	N 7 U U	N 9 U U	U U U U U	U U U U			

U: Unavailable.

^{-:} No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

[†] Chlamydia refers to genital infections caused by *C. trachomatis*. Totals reported to the Division of STD Prevention, NCHSTP.

5 Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention. Last update November 26, 2000.

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

	Gono	rrhea	Hepati Non-A,	tis C;	Legione		Listeriosis	Ly	me ease
Reporting Area	Cum. 2000 [§]	Cum.	Cum.	Cum. 1999	Cum.	Cum.	Cum.	Cum.	Cum.
UNITED STATES	325,596	1999 347,165	2000 2,838	2,839	2000 913	1,004	647	2000 12,874	1999 15,127
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	5,722 84 103 63 2,346 611 2,515	6,414 78 111 50 2,381 572 3,222	16 2 - 4 4 6	16 2 - 7 4 3	51 2 3 5 16 8 17	78 3 8 14 27 12 14	56 2 4 3 27 1	4,313 62 38 1,098 590 2,525	4,465 41 22 23 780 499 3,100
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	34,181 6,862 10,140 5,443 11,736	38,472 6,558 11,827 7,565 12,522	611 65 - 510 36	123 59 - - 64	201 90 - 15 96	242 62 43 21 116	151 82 29 21 19	6,592 3,766 105 1,448 1,273	8,138 3,913 134 1,693 2,398
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	61,766 14,321 5,991 18,467 17,347 5,640	67,164 17,443 6,072 22,323 14,873 6,453	214 12 1 19 182	888 4 1 47 820 16	238 111 39 9 50 29	265 81 46 31 64 43	113 58 8 11 31 5	325 88 32 11 -	579 44 19 17 11 488
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr.	15,724 2,784 1,086 7,584 53 270 1,320	16,087 2,741 1,208 7,937 79 191 1,419	432 7 2 406 1 - 6	301 10 - 287 1 - 3	57 7 14 25 - 2 4	56 13 14 18 2 3 6	14 5 2 5 2	421 322 32 44 2 -	340 219 22 71 1 -
Kans.	2,627	2,512	10	-	5	-	-	17	16
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	90,154 1,671 9,003 2,666 9,891 494 16,844 11,071 16,814 21,700	101,279 1,615 9,635 3,462 9,153 548 18,893 14,479 21,560 21,934	124 - 18 3 3 16 20 3 3 3 58	156 - 21 1 11 17 33 22 1	189 10 64 6 33 N 16 6 7 47	148 19 35 4 40 N 15 11 3	104 2 22 - 8 5 - 9 21 37	974 140 530 11 146 34 46 17	1,288 159 874 6 118 18 74 6
E.S. CENTRAL Ky. Tenn. Ala. Miss.	33,995 3,411 11,469 10,959 8,156	35,316 3,250 11,120 10,857 10,089	427 35 99 8 285	331 25 117 1 188	37 20 12 4 1	50 22 22 4 2	20 3 13 4	47 12 28 6 1	99 18 57 20 4
W.S. CENTRAL Ark. La. Okla. Tex.	50,947 2,920 12,870 3,968 31,189	51,408 3,159 12,672 3,905 31,672	442 9 308 10 115	572 28 299 16 229	18 - 6 5 7	34 1 11 4 18	16 1 - 7 8	45 4 4 1 36	58 5 9 8 36
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	9,568 53 89 52 2,688 958 4,050 230 1,448	9,258 53 82 35 2,469 943 4,185 230 1,261	396 5 3 303 30 16 21 2 16	220 5 8 76 37 34 46 6 8	47 2 5 2 16 1 8 12	48 - 3 - 13 1 7 18 6	38 - 1 9 2 17 4 5	30 - 3 9 11 - - 3 4	16 - 3 3 3 1 2 2 2
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	23,539 2,290 766 19,772 335 376	21,767 2,047 857 18,129 292 442	176 32 27 115 - 2	232 21 21 190 -	75 19 N 56 -	83 21 N 60 1	135 7 6 119 - 3	127 9 15 101 2 N	144 10 15 119 - N
Guam P.R. V.I. Amer. Samoa C.N.M.I.	577 U U U	55 313 U U U	1 U U	2 U U U	1 U U	- U U	- - - -	N U U	N U U U

N: Not notifiable.

U: Unavailable.

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

		, 20001110		Jo, and D	Salmonellosis*						
	Mal	aria	Rabie	s, Animal	NE	TSS		HLIS			
Reporting Area	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999			
UNITED STATES	1,220	1,458	5,736	6,396	35,788	37,641	29,879	32,019			
NEW ENGLAND Maine N.H. Vt. Mass. R.I.	65 6 1 3 27 8	64 3 2 4 22 5	801 130 21 57 268 61	867 171 45 88 221	2,132 123 140 108 1,203 137	2,176 131 137 92 1,180 129	2,124 91 135 114 1,189	2,195 104 135 82 1,196 160			
Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	20 262 81 114 36 31	28 415 71 246 56 42	264 1,112 799 U 193 120	247 1,267 896 U 179 192	421 3,938 1,190 946 820 982	507 5,261 1,367 1,418 1,188 1,288	439 4,356 1,246 866 821 1,423	518 5,137 1,332 1,469 1,093 1,243			
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	120 22 6 46 32 14	167 18 22 76 41 10	162 52 15 22 67 6	167 36 13 10 87 21	5,036 1,530 623 1,383 869 631	5,274 1,267 526 1,566 967 948	3,375 1,350 567 176 898 384	4,590 1,066 472 1,531 948 573			
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	61 27 2 15 2 1 7 7	74 41 13 14 - 1 5	525 90 78 50 115 90 2 100	711 110 149 31 141 176 4 100	2,324 554 351 682 61 99 215 362	2,201 555 248 735 51 93 189 330	2,388 638 312 877 74 105 94 288	2,347 691 224 855 62 118 172 225			
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	327 5 117 16 50 4 36 2 30 67	342 1 97 18 71 4 33 15 29 74	2,325 49 401 554 112 551 155 344 159	2,078 56 389 - 561 108 430 133 231 170	7,989 110 803 63 983 171 1,120 740 1,477 2,522	8,542 163 831 72 1,226 168 1,295 650 1,508 2,629	5,265 137 729 U 839 148 1,072 540 1,549 251	6,340 154 871 U 1,019 150 1,282 505 1,663 696			
E.S. CENTRAL Ky. Tenn. Ala. Miss.	47 18 12 16 1	27 7 9 7 4	199 21 102 76	252 35 93 122 2	2,356 376 656 664 660	2,188 405 571 595 617	1,656 259 755 521 121	1,448 291 581 479 97			
W.S. CENTRAL Ark. La. Okla. Tex.	20 3 8 9	61 3 10 2 46	77 20 - 57 -	482 14 - 91 377	3,962 704 262 390 2,606	3,674 651 714 446 1,863	4,024 587 753 279 2,405	2,748 254 603 346 1,545			
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	52 1 4 - 25 - 9 6 7	44 4 3 1 18 3 7 4 4	247 65 9 55 - 21 78 10 9	215 59 5 44 1 9 81 8	2,797 95 128 68 692 235 841 489 249	2,941 82 127 69 710 361 874 527 191	2,154 - 97 51 654 182 719 451	2,555 1 97 59 693 287 804 565 49			
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	266 33 41 181	264 27 21 203 1	288 - 7 258 23	357 - 4 345 8	5,254 576 300 4,085 61 232	5,384 651 409 3,942 53 329	4,537 670 348 3,270 23 226	4,659 821 461 3,070 32 275			
Guam P.R. V.I. Amer. Samoa C.N.M.I.	5 U U U	1 1 U U	80 U U U	70 U U U	620 U U U	37 630 U U U	UUUU	U U U U			

N: Not notifiable. U: Unavailable. -: No reported cases.
* Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

wee	ks ending			000, and D	<u>ecember</u>	<u>18, 1999</u>	(50th Week)			
			llosis*			philis	l <u>-</u> .			
	NET Cum.	Cum.	Cum.	PHLIS Cum.	Cum.	& Secondary) Cum.	Cum.	crculosis Cum.		
Reporting Area	2000	1999	2000	1999	2000	1999	2000	1999		
UNITED STATES	20,051	16,099	10,291	9,725	5,733	6,433	12,302	14,967		
NEW ENGLAND Maine	383 10	863 5	369 12	833	72 1	58	409 12	420 20		
N.H.	6	18	8	17	2	1	17	16		
Vt. Mass.	4 267	6 736	247	4 715	47	3 35	4 256	3 231		
R.I.	29	31	36	28	4	3	31	42		
Conn.	67	67	66	69	18	16	89	108		
MID. ATLANTIC Upstate N.Y.	2,006 760	1,092 281	1,325 211	722 74	259 15	290 19	2,220 284	2,479 307		
N.Y. City N.J.	716 337	346 270	470 384	233 234	118 48	128 67	1,195 535	1,273 520		
Pa.	193	195	260	181	46 78	76	206	379		
E.N. CENTRAL	3,750	3,169	1,197	1,754	1,123	1,200	1,294	1,588		
Ohio Ind.	409 1,506	410 334	309 147	141 113	69 345	90 427	263 109	265 132		
III.	963	1,295	103	973	350	412	628	782		
Mich. Wis.	642 230	507 623	579 59	453 74	315 44	230 41	214 80	310 99		
W.N. CENTRAL	2,369	1,184	1,884	793	59	129	473	515		
Minn. Iowa	774 522	233 68	837 316	251 58	13 11	9 9	165 35	190 54		
Mo.	633	697	462	349	27	93	192	178		
N. Dak. S. Dak.	51 7	3 18	49 4	2 10	-	-	5 16	6 17		
Nebr. Kans.	142 240	85 80	84 132	68 55	2 6	6 12	23 37	16 54		
S. ATLANTIC	2,925	2,363	1,110	525	1,918	2,040	2,598	3,070		
Del.	23	15	23	10	8	. 8	14	26		
Md. D.C.	204 80	160 51	115 U	58 U	289 48	337 45	239 36	258 52		
Va.	445	130	331	64	126	150	258	268		
W. Va. N.C.	22 385	8 206	17 265	5 92	469	5 449	31 390	37 482		
S.C. Ga.	136 257	119 230	87 167	63 83	212 370	248 435	128 555	222 568		
Fla.	1,373	1,444	105	150	394	363	947	1,157		
E.S. CENTRAL	1,134 495	1,182 232	525 112	677 147	851 82	1,114 101	852 114	999 176		
Ky. Tenn.	340	659	357	457	513	631	305	346		
Ala. Miss.	98 201	117 174	49 7	62 11	122 134	202 180	296 137	295 182		
W.S. CENTRAL	2,893	2,602	2,606	1,162	802	1,022	1,022	1,766		
Ark.	203	74 217	52	26	94 204	85	159 74	166 245		
La. Okla.	138 125	514	191 43	134 160	126	300 182	130	176		
Tex.	2,427	1,797	2,320	842	378	455	659	1,179		
MOUNTAIN Mont.	1,286 8	1,118 10	732	756 -	225	230 1	471 17	516 13		
ldaho	45	27	25	12	1	i	13	15		
Wyo. Colo.	5 266	3 198	3 196	1 156	1 11	4	4 70	3 75		
N. Mex. Ariz.	168 594	145 573	99 329	105 410	21 185	11 206	36 214	61 219		
Utah	80	64	80	66	1	2	46	39		
Nev.	120	98	-	6	5	5	71	91		
PACIFIC Wash.	3,305 447	2,526 121	543 405	2,503 111	424 67	350 65	2,963 236	3,614 239		
Oreg. Calif.	163 2,648	94 2,274	105	86 2,268	6 349	7 274	25 2,481	104 3,034		
Alaska	. 8	4	3	4	-	1	96	59		
Hawaii	39	33	30	34	2	3	125	178		
Guam P.R.	32	19 138	U U	U U	159	144	119	70 178		
V.I. Amer. Samoa	U U	U U	U U	U U	U U	U U	U U	U U		
C.N.M.I.	ŭ	Ü	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ		

N: Not notifiable. U: Unavailable. -: No reported cases.
*Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

	H infl	uenzae.		epatitis (Vi			Measles (Rubeola)						
		sive	Α	cputitio (Vi	В	pc .	Indige	nous	Impo		Total		
Reporting Area	Cum. 2000†	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	2000	Cum. 2000	2000	Cum. 2000	Cum. 2000	Cum. 1999	
UNITED STATES	1,168	1,170	11,963	15,786	6,417	6,662	-	61	-	18	79	94	
NEW ENGLAND	102	97	352	340	97	139	-	3	-	4	7	11	
Maine N.H.	2 12	8 18	21 18	14 17	5 18	1 16	-	2	-	1	3	1	
Vt. Mass.	10 40	5 39	10 122	19 138	6 18	4 44	-	- 1	-	3	3 1	8	
R.I. Conn.	4 34	6 21	25 156	26 126	22 28	33 41	Ū	-	Ū	-	-	2	
MID. ATLANTIC	183	198	1,071	1,140	836	865	-	14	-	5	19	5	
Upstate N.Y. N.Y. City	98 42	81 57	220 375	264 385	137 428	176 275	-	9 5	-	4	9 9	2 3	
N.J. Pa.	33 10	53 7	100 376	145 346	57 214	135 279	-	-	-	- 1	1	-	
E.N. CENTRAL	150	194	1,497	2,867	693	687	_	9	_	-	9	4	
Ohio Ind.	53 30	59 25	264 119	640 102	101 46	90 42	-	2	-	-	2	2	
III. Mich.	54 10	83 20	623 478	816 1,235	110 435	52 473	-	4	-	-	4 3	1 1	
Wis.	3	7	13	74	1	30	-	-	-	-	-	- :	
W.N. CENTRAL Minn.	74 43	75 47	699 184	1,025 95	526 41	347 52	-	3	-	2 1	5 1	1 1	
lowa	1	2 11	65 301	141 667	32	41 215	-	2	-	-	2	:	
Mo. N. Dak.	18 4	1	4	3	381 2	2	-	-	-	-	-	-	
S. Dak. Nebr.	1 3	2 4	3 34	9 49	2 44	1 20	-	-	-	-	-	-	
Kans.	4	8	108	61	24	16	-	1	-	1	2	-	
S. ATLANTIC Del.	292	251	1,505	1,815 2	1,299	1,097 1	-	4	-	-	4	20	
Md. D.C.	75 -	68 5	221 35	299 59	120 34	145 25	-	-	-	-	-	-	
Va. W. Va.	39 9	22 7	154 55	175 40	162 21	97 23	-	2	-	-	2	18 -	
N.C. S.C.	23 15	36 6	149 86	160 46	246 23	212 63	-	-	-	-	-	-	
Ga. Fla.	70 61	68 39	289 516	452 582	222 471	166 365	-	2	-	-	2	2	
E.S. CENTRAL	51	67	383	394	456	465	_	-	_	_	-	2	
Ky. Tenn.	12 26	8 38	48 140	66 147	75 218	46 207	-	-	-	-	-	2	
Ala. Miss.	12 1	18 3	57 138	60 121	56 107	86 126	-	-	-	-	-	-	
W.S. CENTRAL	58	61	2.224	2.962	706	1.104		-		-	-	12	
Ark. La.	2 11	2 15	112 60	74 212	78 93	84 171	-	-	-	-	-	5	
Okla. Tex.	43 2	40 4	256 1,796	489 2,187	154 381	145 704	-	-	-	-	-	- 7	
MOUNTAIN	117	105	987	1,213	561	551		12		1	13	2	
Mont. Idaho	1 4	3	7 42	17 45	7 8	17 29	-	-	-		-	-	
Wyo.	i	1	45	9 215	38	14	-	-	-	-	-	-	
Colo. N. Mex.	21 24	14 19	207 70	51	110 113	98 174	-	2	-	1 -	3	-	
Ariz. Utah	49 11	54 9	474 64	670 63	210 27	131 34	-	3	-	-	3	1 -	
Nev.	6	4	78	143	48	54	-	7	-	-	7	1	
PACIFIC Wash.	141 8	122 8	3,245 279	4,030 387	1,243 113	1,407 76	-	16 2	-	6 1	22 3	37 5	
Oreg. Calif.	29 33	41 53	177 2,765	244 3,362	120 989	114 1,186	-	13	-	2	15	12 17	
Alaska Hawaii	45 26	9 11	11 13	13 24	10 11	16 15	-	1 -	-	3	1 3	3	
Guam	-	-	_	1	-	4	U	-	U	-	-	1	
P.R. V.I.	4 U	2 U	230 U	348 U	259 U	248 U	Ū	Ū	Ū	Ū	Ū	Ū	
Amer. Samoa C.N.M.I.	U	U U	U U	U	U	U U	U	U	U U	U U	U U	U U	

N: Not notifiable. U: Unavailable. -: No reported cases.
*For imported measles, cases include only those resulting from importation from other countries.
*Of 242 cases among children aged <5 years, serotype was reported for 105 and of those, 23 were type b.

TABLE III. (Cont'd) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending December 16, 2000, and December 18, 1999 (50th Week)

	Meninge Dise		Decer	Mumps	o, 1993	(301)	Pertussis		Rubella			
	Cum.	Cum.		Cum.	Cum.		Cum.	Cum.		Cum.	Cum.	
Reporting Area UNITED STATES	2000 1,966	1999 2,243	2000 4	2000 308	1999 364	2000 68	2000 6,368	1999 6,400	2000	2000 151	1999 249	
NEW ENGLAND	1,900	108	4	4	9	6	1,566	861	_	13	249 7	
Maine N.H.	8 12	5 12	-	-	2	-	45 125	96	-	2	-	
Vt.	3	5	-	-	1	4	244	86	-	-	-	
Mass. R.I.	72 9	63 7		1 1	4 2	2	1,085 22	609 38		9 1	7 -	
Conn. MID. ATLANTIC	18 189	16 222	U	2 24	44	U 2	45 616	32 1,020	U	1 9	- 35	
Upstate N.Y.	65	71	-	11	12	2	313	764	-	2	21	
N.Y. City N.J.	37 44	56 50	-	4 3	12 2	-	51 42	60 27	-	7 -	7 4	
Pa.	43	45	-	6	18	-	210	169	-	-	3	
E.N. CENTRAL Ohio	350 94	396 131	-	30 7	51 20	4	725 321	654 291	-	1 -	2	
Ind. III.	48 78	64 104	-	1 6	5 13	3 1	119 <i>7</i> 9	82 97	-	1	1 1	
Mich. Wis.	104 26	61 36	-	16	9 4	-	124 82	70 114	-	-	-	
W.N. CENTRAL	149	222	1	19	14	16	585	484	-	3	130	
Minn. Iowa	21 34	48 38	-	7	1 8	14	365 55	226 95	-	1 -	5 30	
Mo. N. Dak.	68 2	89 4	-	4	1 1	-	79 7	73 18	-	1 -	2	
S. Dak. Nebr.	6 8	11 11	-	4	-	-	7 32	7 9	-	- 1	- 92	
Kans.	10	21	1	4	3	2	40	56	-	-	1	
S. ATLANTIC Del.	308 1	380 10	2	48	49	11	503 9	430 6	-	95 1	37	
Md. D.C.	26	53	-	10	6 2	7	122 3	119 1	-		1	
Va. W. Va.	42 12	55 8	1	11	10	1	112 1	51 4	-	-	-	
N.C.	36	47	-	7	8	-	110	101	-	82	36	
S.C. Ga.	26 47	44 61	-	11 2	5 4	2	41 40	18 40	-	10	-	
Fla. E.S. CENTRAL	118 130	98 156	1 1	7 8	14 15	1 1	65 106	90 111	-	2 5	2	
Ky.	26	33	-	1	-	-	54	42	-	1	-	
Tenn. Ala.	55 36	64 36	1	2 3	11	1 -	32 19	45 21	-	1 3	2	
Miss.	13	23	-	2	4	-	1	3	-	-	-	
W.S. CENTRAL Ark.	131 14	207 35	-	31 5	46	1 -	338 36	215 25	-	6	15 5	
La. Okla.	35 28	66 35	-	4	11 3	1	12 41	9 41	-	1 -	1	
Tex.	54	71	-	22	32	-	249	140	-	5	9	
MOUNTAIN Mont.	165 6	137 4	-	26 1	26	13	796 35	774 2	-	2	16 -	
ldaho Wyo.	7 3	12 5	-	1 4	3	-	64 6	145 2	-	-	-	
Colo. N. Mex.	34 12	36 15	-	2 1	6 N	1	457 89	287 149	-	1 -	1	
Ariz. Utah	91 8	41 16	-	4 7	8	12	99 31	118 58	-	1	13 1	
Nev.	4	8	-	6	5	-	15	13	-	-	i	
PACIFIC Wash.	422 64	415 65	-	118 11	110 2	14 12	1,133 416	1,851 645	-	17 7	5	
Oreg. Calif.	75 266	76 259	N	N 86	Ñ 89	2	113 550	60 1,091	-	10	- 5	
Alaska Hawaii	9 8	259 7 8	-	7 14	3 16	-	22 32	5 5 50	-	-	-	
Guam	ō -	8 1	U	- 14	3	U	3 <u>/</u>	50 2	U	-	-	
P.R. V.I.	9 U	13 U	Ū	- U	Ū	Ū	7 U	31 U	Ū	- U	- U	
Amer. Samoa	U	U	U	U	U	U	U	U	U	U	U	
C.N.M.I.	U	U	U	U	<u>U</u>	U	U	U	U	U	U	

N: Not notifiable.

U: Unavailable.

-: No reported cases.

TABLE IV. Deaths in 122 U.S. cities,* week ending December 16, 2000 (50th Week)

December 16, 2000 (50th Week)															
	A	All Cau	ises, By	Age (Y	ears)		P&I⁺			All Cau	ses, By	Age (Y	ears)		P&I⁺
Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn Cambridge, Mass. Fall River, Mass. Hartford, Conn. Lowell, Mass. Lynn, Mass. New Bedford, Ma New Haven, Conn Providence, R.I. Somerville, Mass Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J.	542 150 . 45 . 14 . 30 . 50 . 25 . 13 .ss. 26 . 45 . U	367 699 388 30 21 322 20 10 244 311 U 2 422 23 45 30 31 37 66 22 22 22 22 24 24 24 25 26 27 27 28 28 28 28 28 28 28 28 28 28 28 28 28	4 2 5 10 4 2 2 11 U 8 2 13 462 5 2 6 4	53 32 2 2 4 5 1 - - - - - - - 3 1 3 1 3 1 - - - - - -	16 11 - - 2 - 1 U - 1 - 36 1 - - 2 - 1 - 1 - - 1 - - 1 - - 1 - - - 1 - - - 1 -	15 10 1 - - 1 - 2 U - 1 - - 2 2 - - 1 - - - - - - - - - - -	60 18 3 3 7 3 5 5 1 1 134 3 1 12 4	S. ATLANTIC Atlanta, Ga Baltimore, Md. Charlotte, N.C. Jacksonville, Fla Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, Tampa, Fla. Washington, D. Wilmington, D. E.S. CENTRAL Birmingham, Al Chattanooga, Te Knoxville, Tenn. Lexington, Ky. Memphis, Tenn Mobile, Ala. Montgomery, A	1,289 173 205 102 171 971 171 96 182 182 183 187 180 181 181 181 181 181 181 181 181 181	868 117 116 77 111 58 37 39 31 64 143 57 18 652 118 64 85 64 134 72 36	266 34 60 14 37 21 10 13 6 8 30 26 7 152 31 12 20 11 16 11	102 18 22 7 15 7 5 3 1 9 6 9 - 611 3 9 8 7 85 5	35 46 43 3 1 - 18 5 - 32 2 2 12 5 7	17 - 1 - 4 3 1 1 1 2 - 2 3 3 - 27 2 1 1 - 2 6 1 1 1	79 4 17 9 7 12 1 5 4 4 15 1 68 16 4 11 5 5
Erie, Pa.§ Jersey City, N.J. New York City, N.J. New York City, N.Y. 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.	50 52 Y. 1,191 66 32 282 46 34 117 28 29 92 3 25 U	40 30 833 27 19 198 36 26 92 19 26 61 3 22 U	7 17 259 18 9 63 5 4 14 7 2 22	2 3 74 13 3 13 2 9 1 1 3	1 13 4 1 4 2 - 1 1 1 - 4	1 1 10 4 - 3 - 2 1 - 2	3 - 55 5 3 20 1 5 7 3 2 4 3 3 U	Nashville, Tenn. W.S. CENTRAL Austin, Tex. Baton Rouge, La Corpus Christi, Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La San Antonio, Te Shreveport, La. Tulsa, Okla.	155 1,064 98 1. 41 Fex. 54 183 67 103 U 49 . 71 x. 257	97 708 68 25 42 122 45 70 U 25 42 174 12 83	39 206 16 7 28 11 23 U 17 9 55 2	13 89 11 4 1 25 7 5 U 2 11 15	2 31 3 1 2 3 1 1 U 3 4 9	4 28 4 2 5 3 4 U 2 3 4 1	13 71 10 - 4 14 3 2 U 2 14 10 4 8
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Gary, Ind. Grand Rapids, Mi Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohi W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans Kansas City, Kans Kansas City, Mo. Lincoln, Nebr.	205 28 115 50 62 U 100 o 79 838 96 50 . 25 103 83 n. 127	1,4833 32241 1181 11044 1177 1122 1311 252 422 10 10 54 1388 211 81 81 55 76 611 73 49 94 94 94 94 94 95 95 97 97 97 97 97 97 97 97 97 97 97 97 97	6 7 6 6 4 4 2 2 3 2 3 6 6 4 9 2 6 4 3 3 2 1 1 0 9 U 1 3 1 8 1 3 5 1 6 6 7 2 6 3 2 1	136 4 2 28 6 7 11 6 23 · 1 3 6 6 15 3 8 4 4 2 U 6 1 55 4 3 2 5 2 7 4	51 1 1 1 1 1 1 5 2 2 1 4 1 1 2 2 2 5 1 1 1 2 2 2 2 1 1 1 1 1 1 1	44 - 9 11 57 71 15 - 3 4 11 11 UU 11 11 18 11	143 3 6 25 12 5 9 12 9 7 6 2 5 18 1 12 2 2 D 5 2 19 3 2 13 2 6 9	MOUNTAIN Albuquerque, N Boise, Idaho Colo. Springs, C Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, U Tucson, Ariz. PACFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawa Los Angeles, Ca Pasadena, Calif. Portland, Oreg. Sacramento, Ca San Diego, Calif. San Francisco, C San Francisco, C Santa Cruz, Calif. Spokane, Wash. Tacoma, Wash.	31 31 31 31 31 32 32 32 32 32 32 32 32 32 32 32 32 32	646 U26 466 737 154 200 1333 U 77 117 632 U U U U 135 97 U U 135 97 U U 22 99 948 U U U 24 948 U U U 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	183 U 4 177 23 66 4 4 33 U 14 23 152 U U U 0 7 7 16 U 0 3 3 28 U 31 28 U 31 28 U 31 28 U 32 U 0 U 0 U 0 U 0 U 0 U 0 U 0 U 0	66 U - 13 5 16 2 13 U 9 8 41 U U U 2 4 U 1 3 U 12 9 U 1 4 5 U	32 U · 3 2 2 7 · 9 U · 5 6 6 17 U U U · 2 2 U · 4 U · 2 4 U · 4 1 U	28 U 1 2 7 1 1 1 U 6 - 16 U U U 1 1 U - 1 U 1 5 U - 7 - U	65 U 3 6 8 19 1 10 U 10 8 91 U U U 2 11 U 3 10 U V 18 U 4 10 6 U
Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	97 92 112 98	77 46 95 74	13 17 11	4 22 2 4	2 1 2 3	1 6 2 2	10 6 7	TOTAL	10,843 [¶]	7,556	2,046	742	269	217	770

U: Unavailable. -: No reported cases.

U: Unavailable. -:No reported cases.

*Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of ≥100,000. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included. ¹Pneumonia and influenza. ¹Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks. ¹Total includes unknown ages.

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