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Update: Investigation of Bioterrorism-Related Anthrax — Connecticut, 2001

CDC and state and local health departments continue investigating cases of bioterrorism-related anthrax. This report revises the number of suspected cases and updates the investigation of a 94-year-old Connecticut (CT) resident who died from inhalational anthrax.

As of December 5, a total of 22 cases of anthrax have been identified; 11 were confirmed as inhalational anthrax, and 11 (seven confirmed and four suspected) were cutaneous. A 54-year-old man who lived in Delaware and who worked at a postal facility in New Jersey (NJ) previously had been classified as having a suspected case of cutaneous anthrax. Additional laboratory findings indicate that the patient's illness no longer meets the CDC surveillance case definition for anthrax (1). Initially, he was classified as having a suspected case because of a lesion on his left hand and elevated levels of antibody (IgG) to the protective antigen component of anthrax toxin. Subsequent biopsies of the skin lesion did not reveal *Bacillus anthracis* in the tissue, and additional confirmatory antibody tests on serum specimens were negative.

The investigation in CT has not identified any additional cases of anthrax through prospective and retrospective surveillance. For prospective surveillance, hospitals, clinicians, postal facilities, and the state medical examiner have been asked to report daily any persons with clinical findings that might be related to anthrax, including sepsis and pneumonia. To date, 50 such patients have been reported. No evidence of anthrax was found in 43 patients and the remaining seven are being evaluated; preliminary investigations of the seven patients have not identified evidence of anthrax. Retrospective surveillance has included a review of all deaths since September 1 involving residents of Oxford and eight surrounding towns (Beacon Falls, Naugatuck, Ansonia, Derby, Woodbury, Shelton, Seymour, and Southbury [total population: 152,481]); 487 death certificates for persons who died during September-November 2001 have been reviewed. Of the 131 deaths attributed to sepsis, pneumonia, sudden death, respiratory arrest, cardiac arrest, or undetermined cause, 66 occurred in hospitals. Of these, 52 had no apparent anthrax disease. For 14 persons who died soon after arrival to the hospital, review of hospital records revealed no evidence of anthrax, but information in the hospital record was insufficient to determine the specific cause of death, and postmortem examinations were not conducted.

The source of exposure for the case of inhalational anthrax in a 94-year-old woman who lived in Oxford, CT, remains unknown. Multiple environmental samples collected

Investigation of Bioterrorism-Related Anthrax — Continued

from all places (e.g., the patient's home, church, voting place, restaurants, and cars in which she traveled) the patient was known to have visited during the 60 days preceding illness onset were negative for *B. anthracis* by culture. Nasal swab specimens were negative from 16 persons epidemiologically linked to the case (e.g., persons who worked in the home and assisted with shopping).

Environmental sampling was performed at the postal processing and distribution center in Wallingford, CT, that serves the towns of Oxford and Seymour and identified B. anthracis spores in three high-speed mail sorters. This facility receives mail from several postal distribution facilities known to have been contaminated by *B. anthracis* spores, including the postal center in Hamilton, NJ, which was the origination site for envelopes containing *B. anthracis* powder that were addressed to two U.S. senators. To evaluate potential cross-contamination of envelopes (i.e., an envelope contaminated from another B. anthracis-contaminated envelope or environmental surface), postal sorting records from the Wallingford facility are being examined to determine the timing and pathways of mail delivered to the CT patient and her local relatives and contacts. Sorting records in Hamilton indicated that an envelope addressed to a postal code adjacent to Oxford had been processed using the same automatic canceling machine at Hamilton <1 minute after one of the two *B. anthracis* powder-containing letters sent to a U.S. senator. This envelope was subsequently sorted at Wallingford and delivered to Seymour. The envelope was received at a residence 4 miles from the home of the CT patient; this envelope was recovered from the recipient and *B. anthracis* spores were detected on the outside of the envelope; none of the members of this household had clinical evidence of anthrax. No record of mail to the CT case-patient processed at Hamilton was found, and no *B. anthracis* spores have been recovered from envelopes found at her home.

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Editorial Note: As of December 5, a total of 11 inhalational anthrax cases have been identified; direct exposure to a *B. anthracis*-containing envelope was likely in the first nine cases (2). The source of exposure to *B. anthracis* for the inhalational anthrax cases in CT and New York City (NYC) remain under investigation by public health and law enforcement officials. No direct exposure to *B. anthracis*-containing envelopes has been identified for these cases. Similar to the first nine cases of inhalational anthrax, exposure to *B. anthracis* might have occurred through the mail from exposure to an envelope containing *B. anthracis* powder. No direct exposure to envelopes containing *B. anthracis* powder. No direct exposure to envelopes containing *B. anthracis* powder. No direct exposure to envelopes containing *B. anthracis* powder has been identified for the inhalational cases in CT and NYC. In the absence of definitive evidence indicating how transmission occurred, infection from a cross-contaminated envelope is one hypothesis being considered by investigators.

Cross-contamination could explain how *B. anthracis* spores were spread to some postal facilities that did not process the envelopes addressed to the U.S. senators.

Investigation of Bioterrorism-Related Anthrax — Continued

Approximately 85 million pieces of mail were processed on the days after the implicated envelopes passed through the NJ and the District of Columbia (DC) sorting facilities until they were closed. Both of these facilities had evidence of widespread environmental contamination with *B. anthracis*. Some of the pieces of mail that passed through these facilities could have been cross-contaminated and, in turn, could have contaminated mail processing equipment or other envelopes processed elsewhere. Despite the high volume of mail distributed to metropolitan areas around these facilities, active surveillance has not identified cases of inhalational anthrax among approximately 10.5 million residents in NJ, DC, Pennsylvania, Maryland, and Virginia or in postal workers since the initial cluster of cases associated with the processing of the implicated letters sent to the U.S. senators. The large population, the duration of active surveillance, and the absence of additional cases of inhalational anthrax indicate that if there is a risk for inhalational anthrax associated with exposure to mail cross-contaminated by the letters addressed to the U.S. senators, it is very low.

Despite this very low risk, persons remaining concerned about their risk may want to take additional steps such as not opening suspicious mail; keeping mail away from your face when you open it and not blowing or sniffing mail or mail contents; washing your hands after you handle the mail; avoiding vigorous handling of mail, such as tearing or shredding mail before disposal; and discarding envelopes after opening mail. However, the effectiveness of these steps in reducing any residual risk is not known.

Suspicious persons or situations should be reported to law enforcement authorities. Health-care providers should remain alert for persons with clinical presentations consistent with early anthrax (3), obtain appropriate diagnostic tests (e.g., blood cultures and chest radiograph) (4), and report suspicious illnesses to local or state public health authorities. Fatalities can be minimized by promptly initiating combination antimicrobial therapy (5). Recommendations for risk reduction for persons with potential occupational exposure are available (6). Public health surveillance for anthrax and research efforts to further define the risk associated with exposure to *B. anthracis* in the environment as a result of the bioterrorist attack is ongoing. CDC will continue to provide updates as new information becomes available.

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Public Health Dispatch

Update: Unexplained Deaths Following Knee Surgery — Minnesota, 2001

Since November 13, 2001, the Minnesota Department of Health (MDH), in collaboration with CDC, has been conducting an investigation of three patients who died unexpectedly within 1 week following knee surgery (1). Patient 1 had received a knee osteochondral allograft, and patients 2 and 3 had undergone total knee replacement surgery. Epidemiologic and microbiologic investigations have not linked the deaths of the three patients.

Blood cultures obtained from patient 1 before his death grew a clostridial species that was identified subsequently at MDH and CDC as *Clostridium sordellii* by biochemical and molecular typing. Blood cultures from patients 2 and 3 did not yield growth of any bacteria. Molecular and special studies have not identified any *Clostridium* species in autopsy tissues from patients 2 and 3, and the cause of death in these patients remains unexplained. On the basis of investigative findings, MDH lifted a moratorium on elective knee surgery on November 25.

As of December 4, neither surveillance in Minnesota by MDH nor enhanced case finding by CDC outside of Minnesota and follow-up of reports to CDC have identified any additional cases of *C. sordellii* infection associated with severe hemodynamic collapse or death in patients recently undergoing knee or large joint surgery. Because infection associated with contaminated graft tissue is a known but uncommon complication of allograft surgery (*2*), MDH, CDC, and the Food and Drug Administration have initiated an investigation to determine whether the osteochondral allograft might have been the source for the *C. sordellii* found in patient 1. Nonimplanted knee tissue from the same donor source as the allograft used in patient 1 was obtained by CDC from the same tissue bank. Preliminary cultures of this tissue have yielded growth of *Clostridium* species; biochemical and molecular testing to identify the species is under way. Reports of other allograft recipients infected with clostridial species have been received at CDC and are being investigated.

Clinicians should consider possible clostridial infection in patients with evidence of infection following allograft implantation. Clinical evaluation should include looking for symptoms and signs of sepsis, including fever, hemodynamic compromise, and/or abdominal pain. In some patients, only local symptoms (e.g., knee pain) may be present during the early course of infection. Diagnostic evaluation should include two sets of blood cultures for both aerobes and anaerobes; these cultures should be incubated for 7 days. If appropriate, other specimens (e.g., knee aspirate or tissue) should be obtained and cultured aerobically and anaerobically. If appropriate, health-care providers should consider expanding empiric therapy to include anaerobic coverage. Consultation with an infectious disease physician might be helpful.

Health-care providers should report cases of clostridial infection following allograft implantation to their state health department or CDC's Division of Healthcare Quality Promotion, telephone 800-893-0485.

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In the United States, approximately 50,000 knee surgeries are performed each year for repairing anterior cruciate ligament (ACL) injuries (1). Tissue allografts frequently are used for ACL reconstruction, and septic arthritis is a rare complication of such procedures. This report describes four patients who acquired postsurgical septic arthritis probably associated with contaminated bone-tendon-bone allografts used for ACL reconstruction. Effective sterilization methods that do not functionally alter musculoskeletal tissue are needed to prevent allograft-related infections.

Florida

On April 5, 2000, at a surgical center, a girl aged 16 years had ACL reconstruction using a bone-tendon-bone allograft. On April 21 at a local orthopedic clinic, she sought medical care for swelling and redness of the left knee. On examination, septic arthritis was diagnosed, and she was treated with joint irrigation, a 6-week course of intravenous antimicrobial therapy, and removal of the allograft and screw. Cultures from the left knee aspirate yielded *Pseudomonas aeruginosa, Staphylococcus aureus*, and *Enterococcus faecalis*.

On April 7 at a surgical center, a man aged 40 years underwent ACL reconstruction using a bone-tendon-bone allograft. On April 24, he sought medical care for drainage from the knee. On examination, septic arthritis was diagnosed; his treatment was an 8-week course of antimicrobials and screw removal. *P. aeruginosa* was cultured from the surgical site.

The allografts used for the two patients were supplied by a Texas tissue bank (tissue bank A) and were harvested from a common donor. Both patients' initial ACL reconstruction procedures were performed on different days by different surgeons using different arthroscopic instruments but at the same surgical center. The local health department conducted an onsite investigation of the center and identified no breaches in infection-control procedures. At tissue bank A, the implicated allografts had been irradiated and processed using standard quality-control procedures. All other allografts used during the preceding 4 years at this surgical center had been supplied by a tissue bank other than tissue bank A; no postoperative infections were detected by orthopedic surgeons at follow-up visits among approximately 1,000 ACL reconstructions performed at this center during the 4-year period. *P. aeruginosa* isolates cultured from the surgical site infections of the two patients had genotypic patterns that were indistinguishable from each another by pulsed-field gel electrophoresis.

Florida and Louisiana

On October 9 at a surgical center in Florida, a woman aged 55 years had ACL reconstruction using a bone-tendon-bone allograft. On October 17, she was taken to an orthopedic clinic for purulent drainage from the left knee. On examination, septic arthritis was diagnosed, and she was treated with joint irrigation and 12 weeks of antimicrobial therapy. On July 11, 2001, the patient required a total knee arthroplasty. *Citrobacter werkmanii/ youngae* and group B beta hemolytic streptococci grew from the knee aspirate.

On October 19 in Louisiana, a woman aged 29 years had ACL reconstruction using a bone-tendon-bone allograft at a local surgical center. On November 7 at an orthopedic clinic, she presented with a temperature of 103° F (39.4° C) and septic arthritis. She was treated with joint irrigation and 13 weeks of antimicrobial therapy. *Klebsiella oxytoca* and *Hafnia alvei* were cultured from the knee aspirate.

Septic Arthritis — Continued

Both patients received allografts from the same Florida tissue bank (tissue bank B), and the allografts were from a common donor. When tissue bank B conducted a traceback investigation and reviewed quality-control procedures, the implicated allografts had not received terminal sterilization with gamma irradiation. The same species of organisms isolated from the two recipients and *Serratia liquefaciens* were cultured from the donor allografts during tissue processing; other donor tissues were culture negative. No isolates from the donor or recipients were available for additional testing.

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Editorial Note: In the cases described in this report, clinicians suspected contaminated allografts because of the rarity of septic arthritis following arthroscopic interventions and the polymicrobial nature of these infections and worked with local public health authorities and tissue bank staff to link the infections to allografts of common donors. The epidemiologic and laboratory investigation related to tissue bank A indicated that the allografts were the source of the infection despite no apparent lapses in tissue processing. Cases related to tissue bank B were linked to allografts from a common donor that were released inadvertently before standard terminal sterilization procedures were conducted.

In 1999, U.S. tissue banks distributed approximately 750,000 allografts for transplantation (2). Transmission of infectious agents (e.g., fungi, bacteria, and human immunodeficiency virus [HIV]) caused by contaminated allografts has been described (3–5). The number of persons who develop septic arthritis caused by bacterially contaminated allografts is unknown. In addition, tissue banks, donors, and recipients often are located in different states, complicating detection of bacterial infections associated with contaminated allografts. The Food and Drug Administration (FDA) requires screening of tissue donors for HIV, hepatitis B and C, and other bloodborne pathogens. Reporting of infections resulting from contaminated allografts is not required. FDA has proposed regulations that would require reporting adverse reactions that involve the transmission of communicable diseases if fatal, life threatening, or results in permanent impairment.

The American Association of Tissue Banks (AATB) publishes quality standards for procuring and processing tissue, and provides guidelines on donor screening, time limits for retrieval of soft tissues, and procedures for preservation (e.g., freezing or freezedrying), sterilization, preparation, and evaluation, and labeling of tissue components (6). Gamma irradiation or ethylene oxide are used to sterilize allografts. Tissue banks use gamma irradiation for sterilization, but high doses of gamma irradiation may adversely affect the biomechanical properties of allografts (7). Ethylene oxide has limited ability to penetrate tissue and has been associated with adverse patient outcomes (8,9). Concern about possible sterilization-related complications has resulted in musculoskeletal tissues (e.g., bone-tendon-bone allografts) being processed aseptically but is not necessarily sterile. Although aseptic processing avoids contamination of tissue at the tissue bank, it does not eliminate contamination originating from the donor that might be inherent to the graft. AATB standards require that tissue banks establish a list of organisms which, when

Septic Arthritis — Continued

cultured from tissue, necessitate discarding, sterilization, or disinfection of harvested tissues (6). However, not all tissue is cultured, and AATB does not specify the organisms for which corrective actions should be taken (6).

According to the Office of the Inspector General, approximately 44% of tissue banks identified were not accredited by AATB or inspected by Florida or New York (the two states that require licensing and inspection of tissue banks) (2), and this probably represents an underestimate of the tissue banks that are unaccredited or unlicensed (10). Tissue banks that lack accreditation and licensure are not required to comply with external quality requirements beyond donor screening for HIV and hepatitis (2).

This report underscores the need for 1) standard practices for screening, disinfecting, sterilizing, or discarding potentially contaminated allografts; 2) mechanisms for certification and oversight of tissue banks and adherence to quality standards; 3) a system for reporting and investigating infections (bacterial, viral, or fungal) potentially transmitted through human tissues; and 4) the development of safe and effective sterilization methods for musculoskeletal tissue. When septic arthritis occurs after use of an allograft, allograft contamination should be suspected, especially when the infection is polymicrobial or associated with Gram-negative organisms. Clinicians should report infections involving allograft tissue to FDA's MedWatch system and through local and state health departments to CDC's Division of Healthcare Quality Promotion, National Center for Infectious Diseases, telephone 800-893-0485.

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Influenza Activity — United States, 2001–02 Season

In collaboration with the World Health Organization (WHO) and its collaborating laboratories, National Respiratory and Enteric Virus Surveillance System (NREVSS) collaborating laboratories, state and local health departments, and a network of sentinel physicians, CDC conducts surveillance to monitor influenza activity and to detect antigenic changes in circulating strains of influenza viruses. This report summarizes influenza activity in the United States* (1) during September 30–November 24, 2001, when the viruses isolated most frequently were influenza A (H3N2). These viruses were well matched antigenically by the 2001–02 influenza A (H3N2) vaccine. Vaccine supplies are plentiful and influenza vaccine should continue to be offered during December and later.

As of November 24, WHO and NREVSS collaborating laboratories in the United States tested 8,140 specimens for influenza viruses; 73 (0.9%) were positive. The percentage of positive influenza isolates identified each week is an indicator of the level of influenza activity, and for the weeks ending October 6 through November 24, the percentage of respiratory specimens testing positive for influenza viruses ranged from 0.4% to 1.7%. These percentages are low compared with the 24%–33% testing positive at the peak of the 1998–99, 1999–2000, and 2000–01 seasons. Of the 73 influenza isolates reported since September 30, 70 (96%) were influenza A viruses and three (4%) were influenza B viruses. Of the 70 influenza A viruses identified, 45 (64%) have been subtyped; 44 were influenza A (H3N2) viruses and one was an influenza A (H1N1) virus. Influenza A (H3N2) isolates were identified in Alaska, Arizona, Colorado, Florida, New York, North Carolina, North Dakota, Texas, Utah, and Wisconsin. The influenza A (H1N1) isolate was identified in Washington, and unsubtyped influenza A isolates were identified in Alabama, Alaska, Hawaii, Louisiana, Minnesota, New York, Washington, and Wisconsin. Influenza B isolates were identified in Louisiana, Michigan, and Texas. Thirty-nine (52%) of the 73 influenza viruses isolated were identified in Alaska.

CDC antigenically characterized 10 influenza isolates collected in September and 13 collected in October. They consisted of 20 influenza A (H3N2) viruses, two influenza A (H1N1) viruses, and one influenza B virus. The antigenically characterized influenza A (H3N2), influenza A (H1N1), and influenza B isolates were similar to the vaccine strains A/Panama/2007/99 (H3N2), A/New Caledonia/20/99 (H1N1), and B/Sichuan/379/99, respectively.

During September 30–November 24, the weekly percentage of patient visits for influenza-like illness (ILI)⁺ to approximately 650 U.S. sentinel physicians ranged from 1.0% to 1.4%. For the week ending November 24, the percentage of patient visits for ILI was 1.4%, which is less than the national baseline of 1.9%[§]. During the same week, influenza activity[¶], as reported by state epidemiologists, was regional in Alaska and

^{*}As of November 29, 2001.

[†] Temperature of >100.0° F (>37.8° C) and either cough or sore throat in the absence of a known cause.

[§] The national baseline was calculated as the mean percentage of visits for ILI during noninfluenza weeks plus two standard deviations. Because of wide variability in regional level data, to calculate region-specific baselines is not possible and to apply the national baseline to regional level data is not appropriate.

[¶] Levels of activity: 1) no activity, 2) sporadic—sporadically occurring ILI or laboratory-confirmed influenza with no outbreaks detected, 3) regional—outbreaks of ILI or laboratory-confirmed influenza in counties with a combined population of <50% of the state's population, and 4) widespread—outbreaks of ILI or laboratory-confirmed influenza in counties with a combined population of ≥50% of the state's population.

Influenza Activity — Continued

sporadic in 25 states (Alabama, Arizona, California, Colorado, Connecticut, Georgia, Indiana, Iowa, Kansas, Kentucky, Maine, Michigan, Missouri, Nevada, New Mexico, New York, North Carolina, Ohio, Tennessee, Texas, Utah, Vermont, West Virginia, Wisconsin, and Wyoming), New York City, and District of Columbia; 23 states reported no influenza activity, and one state did not report.

During the week ending November 24, the 122 Cities Mortality Reporting System attributed 6.1% of recorded deaths to pneumonia and influenza (P&I). This percentage was below the epidemic threshold** of 7.4% for that week. The percentage of P&I deaths has been below the epidemic threshold for each week since September 30.

In November, two virologically confirmed institutional outbreaks caused by influenza A viruses were reported to CDC. On November 14, an elementary school in Fort Collins, Colorado, reported elevated and increasing absenteeism among its students. Of 675 students, 53 (8%) were absent on November 14, 96 (14%) were absent on November 15, and 110 (16%) were absent on November 16. Baseline absenteeism on November 12–13 was 18–20 students. Two of the three specimens submitted to the state laboratory for viral culture were positive for influenza A (H3N2). The school remained open and a letter was sent to parents describing influenza symptoms and requesting that sick children be kept at home. Use of influenza antiviral agents was left to the discretion of the child's health-care provider and family. Nursing homes in the Fort Collins area were advised of influenza activity in the community and a broadcast facsimile outlining antiviral treatments available for influenza was sent to all primary-care providers.

On November 17, an influenza A outbreak was reported in a long-term–care facility with 160 residents located in the Hudson Valley region of New York; 14 residents and eight staff members had an influenza-like illness and four of six ill residents tested positive for influenza A by rapid antigen testing. On November 18, all residents began to receive antiviral medication and since then, no new cases of influenza-like illness in this facility have been reported. The facility received its order of influenza vaccine a week and a half before the outbreak and vaccinated residents on November 12–16.

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^{**} The expected baseline proportion of P&I deaths reported by the 122 Cities Mortality Reporting System is projected using a robust regression procedure in which a periodic regression model is applied to the observed percentage of deaths from P&I since 1983. The epidemic threshold is 1.654 standard deviations above the seasonal baseline. Before the 1999–2000 season, a new case definition for a P&I death was introduced. During the summer of 2000, the baseline and epidemic thresholds were adjusted manually to account for these changes in case definition. For the 2001–02 season, sufficient data have been collected using the new case definition to allow projection of the baseline using the regression procedure employed before the 2000–01 season.

Influenza Activity — Continued

Editorial Note: The four influenza surveillance system components indicated low levels of influenza activity in the United States during September 30–November 24. The number of influenza viruses isolated this season is relatively low and it is too early to determine which strain(s) will predominate. However, two influenza A outbreaks were detected in November and influenza activity is expected to increase during the next few weeks to months. The viruses isolated most frequently have been influenza A (H3N2) viruses. The 2001–02 influenza vaccine strains are well matched to the influenza isolates that have been characterized antigenically this season.

The best prevention against influenza is vaccination. Vaccine supplies are plentiful and are available for immediate shipment from the three U.S. licensed manufacturers. Manufacturers estimate that approximately 87 million doses of influenza vaccine will be produced this year compared with 76.8 million doses available during the 1999–2000 season and 70.4 million doses available during the 2000–01 season. By the end of November, approximately 74.2 million (85%) of the projected 87 million doses of vaccine will have been distributed. An additional 12.8 million doses are expected to be available in December.

Health-care providers should continue to offer influenza vaccine during December and later because persons can benefit from vaccination after influenza activity has been detected in their community (2). The most important persons to be vaccinated are those in groups at increased risk for complications from influenza (i.e., persons aged \geq 65 years and persons aged 6 months–64 years with certain underlying medical conditions [3]), and health-care providers. In addition, household contacts of high-risk persons, healthy persons aged 50–64 years, and anyone who wants to reduce the likelihood of becoming ill with influenza should be vaccinated.

CDC collects and reports U.S. influenza surveillance data during October–May. This information is updated weekly and is available through CDC voice information, 888-232-3228, fax information, 888-232-3299 (request document number 361100) or at http:// www.cdc.gov/ncidod/diseases/flu/weekly.htm.

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Notice to Readers

Use of Onsite Technologies for Rapidly Assessing Environmental *Bacillus anthracis* Contamination on Surfaces in Buildings

Environmental sampling to ascertain the presence of *Bacillus anthracis* spores in buildings is an important tool for assessing risk for exposure. Similar to diagnostic testing, culture with positive identification of *B. anthracis* (CDC culture method) is the confirmatory test. Laboratory-based polymerase chain reaction (PCR) methods for detecting genetic material of *B. anthracis* can be used in preliminary assessments and as adjuncts to microbiologic methods. Although these tests are consistent with culture results, PCR methods are not approved by the Food and Drug Administration, and results should not be the basis for clinical decisions.

Rapid-assay devices that can provide results within minutes are used for onsite detection of environmental contamination. Some of these devices are PCR-based assays, and others are immune-based assays for *B. anthracis*. CDC has not obtained validation data for rapid-assay devices. A recent CDC evaluation of *B. anthracis* contamination at the Brentwood postal facility in the District of Columbia included use of one onsite PCR-based device and CDC culture method. Of 107 samples analyzed using CDC culture method and the PCR-based device, 95 (89%) were negative by both methods. Of six samples identified as positive by CDC culture method, two were positive using the PCR-based device. Of eight samples identified as positive by the PCR-based device, two were positive by CDC culture method. Although these results indicate a poor agreement between results from the onsite PCR-based device and CDC culture method, this assessment was not intended as a formal validation test because of limited capacity to implement adequate quality-control measures and the small number of *B. anthracis* positive samples.

The apparently poor agreement of the onsite PCR-based device could be attributed to several factors such as the concentration of spores on contaminated surfaces, sample collection and preparation procedures, sample splitting, and the methods used for removing the sample from collection material. Furthermore, PCR- or immune-based tests do not distinguish viable from nonviable spores and can produce positive scores for samples that culture methods would define as negative. As a result, these methods are not useful for evaluating the success of disinfection techniques that do not remove non-viable spores.

Public health officials are urged to understand the limitations of onsite, rapid technologies for *B. anthracis* before using them for public health decision making. Until validation testing is complete and guidelines for effective use are developed, PCR- or immune-based assay results for *B. anthracis* should not be used alone, but should be confirmed with samples analyzed by culture methods to make public health decisions. Notice to Readers

CDC Recognition of Members of MMWR Distribution Partnership

The recent bioterrorist attacks represent a national emergency that requires action by all of those responsible for public health and safety. In October and November in response to these attacks, CDC developed guidelines for anthrax treatment, prophylaxis, and exposure management that required immediate dissemination to all health-care professionals. To expand its distribution, *MMWR* enlisted the assistance of various organizations, agencies, publications, and health-care plans in a distribution partnership. Participants in this partnership electronically distributed to their members and subscribers bioterrorism-related reports published in *MMWR*. As a result, millions of health-care professionals and the public were notified immediately about critical public health information within hours of its release by CDC. CDC appreciates this collective effort to protect public health and safety.

Following are members of the *MMWR* distribution partnership: Alliance of Community Health Plans American Academy of Family Physicians American Academy of Orthopaedic Surgeons American Academy of Pediatrics American Association of Health Plans American Association of Poison Control Centers American Association of Public Health Laboratories American College of Emergency Physicians American College of Physicians, American Society of Internal Medicine American Hospital Association American Medical Association, Office of Specialty Society Relations Association of American Medical Colleges Association of State and Territorial Health Officials **Blue Cross Blue Shield Association Council of State and Territorial Epidemiologists** Employers' Managed Health Care Association Environmental Protection Agency ePocrates, Inc. Federal Emergency Management Agency Federation of State Medical Boards International Association of Fire Chiefs Journal of the American Medical Association Kaiser Permanente MyDrugRep.com, Inc. National Association of County and City Health Officials New England Journal of Medicine National Institutes of Health, National Library of Medicine National Institute for Health Care Management Research and Education Foundation U.S. Department of State Washington Business Group on Health WebMD CDC invites other organizations and agencies to join this distribution partnership by contacting MMWR at mmwrq@cdc.gov.



FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending December 1, 2001, with historical data

- * No measles or rubella cases were reported for the current 4-week period yielding a ratio for week 48 of zero (0).
- [†] Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

		Cum. 2001		Cum. 2001
Anthrax		15	Poliomyelitis, paralytic	-
Brucellosis [†]		82	Psittacosis [†]	23
Cholera		3	Q fever [†]	22
Cyclosporiasis	5 [†]	131	Rabies, human	1
Diphtheria		2	Rocky Mountain spotted fever (RMSF)	565
Ehrlichiosis:	human granulocytic (HGE)†	206	Rubella, congenital syndrome	1
	human monocytic (HME) [†]	86	Streptococcal disease, invasive, group A	3,274
Encephalitis:	California serogroup viral [†]	99	Streptococcal toxic-shock syndrome [†]	44
	eastern equine ^Ť	8	Syphilis, congenital [¶]	190
	St. Louis	2	Tetanus	23
	western equine [†]	-	Toxic-shock syndrome	115
Hansen diseas	se (leprosy) [†]	78	Trichinosis	26
Hantavirus pu	Imonary syndrome [†]	6	Tularemia [†]	98
Hemolytic ure	mic syndrome, postdiarrheal [†]	139	Typhoid fever	263
HIV infection,	pediatric [™]	200	Yellow fever	-
Plague	•	2		

TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending December 1, 2001 (48th Week)*

-: No reported cases. *Incidence data for reporting year 2001 are provisional and cumulative (year-to-date).

[†] Not notifiable in all states.

⁵ Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV,

STD, and TB Prevention (NCHSTP). Last updated November 27, 2001. Updated from reports to the Division of STD Prevention, NCHSTP.

								Escherichia	<i>coli</i> 0157:H7	7†
	All	OS Cum	Chlam	ydia ^s	Cryptosp	oridiosis	NET	rss Cum	PH	LIS
Reporting Area	2001 [¶]	2000	2001	2000	2001	2000	2001	2000	2001	2000
UNITED STATES	37,411	35,685	656,773	639,742	3,247	2,829	2,893	4,276	2,156	3,518
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	1,403 44 37 15 704 95 508	1,863 38 30 37 1,128 91 539	21,429 1,239 1,246 579 9,149 2,719 6,497	21,706 1,344 1,032 493 9,291 2,486 7,060	123 18 16 31 50 8	130 20 22 26 34 3 25	224 27 35 14 115 17 16	366 31 35 35 161 20 84	225 27 30 8 112 11 37	371 28 38 36 167 18 84
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	9,346 945 5,253 1,607 1,541	7,605 676 3,919 1,554 1,456	75,103 13,575 27,353 10,980 23,195	60,869 3,037 24,310 9,855 23,667	270 106 99 13 52	359 120 159 19 61	207 154 12 41 N	420 281 23 116 N	181 136 11 34	338 76 18 116 128
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	2,812 538 343 1,255 500 176	3,411 533 347 1,692 648 191	107,979 22,404 13,985 30,330 27,833 13,427	110,537 28,959 12,499 30,660 23,306 15,113	1,407 161 79 408 175 584	930 253 57 119 91 410	752 214 81 153 93 211	1,041 255 119 188 139 340	495 153 42 128 82 90	734 223 83 156 104 168
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. S. Dak. Nebr. Kans.	808 133 85 405 2 23 68 92	809 160 83 367 3 7 68 121	33,234 6,709 4,558 11,674 827 1,682 2,206 5,578	36,363 7,523 4,881 12,392 810 1,696 3,445 5,616	498 176 78 44 13 7 177 3	345 123 74 29 15 15 80 9	540 256 82 61 18 42 59 22	635 185 179 108 19 55 61 28	449 212 62 89 32 41 13	605 216 148 96 21 58 48 18
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	11,517 231 1,698 782 911 95 845 645 1,528 4,782	10,027 198 1,192 784 745 57 644 737 1,118 4,552	123,700 2,511 11,078 2,704 16,519 2,158 18,949 10,248 27,394 32,139	120,423 2,651 12,700 2,945 14,625 1,979 20,148 8,991 25,791 30,593	312 6 38 11 24 2 27 7 128 69	449 6 9 17 18 3 26 - 170 200	227 4 28 - 49 10 54 17 33 32	354 34 1 70 15 87 21 40 83	139 7 1 U 39 8 43 11 15 15	280 1 2 66 13 68 16 38 76
E.S. CENTRAL Ky. Tenn. Ala. Miss.	1,671 315 540 415 401	1,781 185 748 455 393	44,785 7,882 13,180 13,316 10,407	47,046 7,431 13,609 14,251 11,755	46 4 13 16 13	49 7 11 15 16	125 58 42 17 8	141 40 54 10 37	110 49 46 6 9	113 32 52 9 20
W.S. CENTRAL Ark. La. Okla. Tex.	3,856 189 806 214 2,647	3,666 170 632 322 2,542	96,125 6,389 15,893 9,573 64,270	96,307 6,031 16,686 8,617 64,973	112 8 7 15 82	159 15 12 17 115	105 13 4 32 56	222 56 15 19 132	91 - 26 28 37	277 38 49 17 173
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	1,288 15 19 4 267 137 502 110 234	1,324 14 20 9 326 140 410 133 272	37,747 1,775 1,788 767 8,876 5,313 13,216 1,619 4,393	34,952 1,265 1,727 731 9,015 4,775 11,721 2,077 3,641	227 37 22 7 40 27 7 82 5	168 10 23 5 69 21 10 26 4	273 20 71 7 86 14 29 30 16	411 30 72 19 154 22 51 49 14	131 - 1 53 11 23 42 1	303 - 40 11 110 18 43 71 10
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	4,710 483 213 3,898 18 98	5,199 463 170 4,444 23 99	116,671 12,473 6,757 91,499 2,389 3,553	111,539 11,929 6,397 87,601 2,318 3,294	252 - 49 199 1 3	240 U 20 220	440 123 65 230 4 18	686 221 133 287 31 14	335 62 61 203 1 8	497 203 114 163 6 11
Guam P.R. V.I. Amer. Samoa C.N.M.I.	12 1,113 11 1 -	13 1,242 32 -	2,240 53 U 124	465 U - U U	- - U -	- - U U	N 1 - U -	N 7 - U U		

TABLE II. Provisional cases	of selected notifiable	diseases, United States,
weeks ending December	1, 2001, and Decembe	er 2, 2000 (48th Week)*

N: Not notifiable.
U: Unavailable.
·: No reported cases.
C.N.M.I.: Commonwealth of Northern Mariana Islands.
* Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).
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.*Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention. Last updated November 27, 2001.

	Gonorrhea		Hepatit Non-A, I	tis C; Non-B	Legione	llosis	Listeriosis	Lyme Disease	
Reporting Area	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2001	Cum. 2000
UNITED STATES	302,379	327,661	2,957	2,913	962	1,013	464	11,653	15,862
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	6,136 132 171 63 2,868 776 2,126	6,116 85 97 60 2,539 604 2,731	20 - 7 13 -	29 2 4 18 5	71 8 11 5 21 12 14	53 2 3 5 17 9 17	40 2 4 3 25 1 5	3,783 148 16 826 453 2,340	5,216 63 40 1,145 571 3,397
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	38,825 8,111 11,676 7,426 11,612	36,211 6,792 10,738 6,667 12,014	1,449 53 - 1,342 54	641 37 - 561 43	192 64 31 13 84	286 89 46 22 129	66 27 12 12 15	5,762 3,404 9 927 1,422	8,150 3,540 177 2,427 2,006
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	56,276 12,547 6,277 16,776 15,797 4,879	65,648 17,796 5,869 19,269 16,234 6,480	150 5 1 13 131 -	218 12 19 187	280 126 22 19 77 36	261 106 36 31 48 40	65 15 8 11 23 8	656 111 23 21 13 488	764 58 22 35 23 626
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr.	14,141 2,171 1,199 7,273 37 262 713	16,483 2,944 1,142 8,123 68 261 1,385	714 9 - 688 - - 6	563 5 2 544 - - 4	47 9 8 21 1 3 4	55 7 13 25 - 2 4	19 2 2 10 - 1	380 314 36 24 - - 4	422 322 33 45 1 - 4
Kans. S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	2,486 76,101 1,545 6,205 2,465 9,693 668 15,240 6,808 15,240 6,808 15,055 18,422	2,560 85,108 1,598 8,833 2,478 9,579 609 16,439 7,788 16,873 20,911	11 97 - - 9 19 6 1 46	8 101 2 12 3 3 15 17 3 3 43	1 185 12 35 8 23 N 11 13 10 73	4 182 65 6 33 N 15 6 7 40	4 70 - 13 5 5 5 14 14	2 803 49 522 16 116 13 39 5 - 43	17 1,051 167 10 143 32 44 14 - 34
E.S. CENTRAL Ky. Tenn. Ala. Miss.	29,240 3,168 8,883 10,352 6,837	33,839 3,248 10,842 11,188 8,561	172 9 59 4 100	427 35 94 10 288	53 11 27 13 2	37 20 10 4 3	20 5 8 7	60 22 29 8 1	49 12 28 6 3
W.S. CENTRAL Ark. La. Okla. Tex.	46,914 3,961 10,876 4,371 27,706	50,865 3,529 12,406 3,828 31,102	177 4 88 4 81	695 9 426 10 250	11 - 2 3 6	26 - 7 5 14	18 1 - 2 15	82 1 2 - 79	88 5 7 1 75
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	9,257 98 70 2,768 882 3,610 125 1,627	9,733 48 84 45 2,964 1,062 3,893 209 1,428	52 1 2 8 10 11 9 3 8	71 5 3 2 13 14 19 1 14	56 3 1 17 3 22 6 4	43 2 5 15 1 7 12 1	37 - 1 2 10 7 8 2 7	13 - 5 1 1 2 1 2	13 - 3 - - - 3 4
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	25,489 2,763 1,046 20,750 383 547	23,658 2,148 937 19,797 328 448	126 23 13 90	168 31 25 110 - 2	67 10 N 53 - 4	70 17 N 52 1	129 10 9 104 - 6	114 8 10 94 2 N	109 9 12 86 2 N
Guam P.R. V.I. Amer. Samoa C.N.M.I.	541 6 U 14	51 479 - U U	- 1 - U -	3 1 - U U	2 - U	- 1 - U U	- - - -	- N - U -	N U U

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States,
weeks ending December 1, 2001, and December 2, 2000 (48th Week)*

N: Not notifiable. U: Unavailable. - : No reported cases. * Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

						Salmo	nellosis†	
	Mal	aria	Rabies	, Animal	NET	TSS	Pł	ILIS
Reporting Area	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	2000
UNITED STATES	1,156	1,373	7,218	6,526	33,911	36,341	27,020	30,299
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	85 4 2 1 38 13 27	69 6 1 32 8 19	686 63 22 60 249 67 225	783 127 21 57 262 56 260	2,220 162 162 76 1,258 131 431	2,055 119 136 104 1,179 124 393	2,101 151 149 63 1,115 168 455	2,110 97 140 101 1,200 148 424
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	330 65 196 35 34	372 73 217 47 35	1,122 739 29 180 174	1,233 789 18 184 242	4,062 1,179 996 905 982	4,710 1,158 1,132 1,097 1,323	3,588 1,213 1,297 657 421	4,989 1,217 1,217 975 1,580
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	132 22 16 34 39 21	137 20 6 64 31 16	142 51 15 24 46 6	151 50 - 22 68 11	4,450 1,184 494 1,227 768 777	5,045 1,459 601 1,428 834 723	3,844 1,076 459 1,049 778 482	3,435 1,372 576 231 886 370
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	35 6 9 13 - 2 5	66 27 2 19 2 1 8 7	332 44 74 41 37 42 4 90	511 87 74 50 113 90 2 95	2,203 624 330 630 56 144 144 275	2,255 515 344 677 55 93 209 362	2,281 665 301 906 80 118 - 211	2,412 645 337 826 75 101 139 289
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	269 2 108 13 46 1 18 7 30 30	309 5 106 49 4 34 2 30 63	2,093 30 332 - 461 131 550 112 311 166	2,235 49 393 110 541 149 302 152	8,130 87 768 79 1,247 131 1,274 847 1,630 2,067	7,583 114 711 942 156 1,076 716 1,434 2,373	5,595 104 836 U 958 133 1,219 677 1,210 458	5,617 125 684 U 885 144 1,082 543 1,649 505
E.S. CENTRAL Ky. Tenn. Ala. Miss.	33 12 11 6 4	45 18 12 14 1	196 27 103 64 2	196 20 100 75 1	2,473 355 598 714 806	2,298 360 637 628 673	1,735 217 758 474 286	1,726 249 773 576 128
W.S. CENTRAL Ark. La. Okla. Tex.	12 3 5 3 1	70 3 12 9 46	2,082 20 3 59 2,000	852 20 4 54 774	3,607 866 413 454 1,874	4,721 691 851 369 2,810	2,537 92 952 375 1,118	2,892 563 720 286 1,323
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	57 3 - 22 3 13 4 9	50 1 - 24 - 9 6 6	231 38 28 20 - 14 115 15 15 1	262 64 9 56 - 20 94 10 9	2,031 72 134 552 270 588 209 151	2,579 90 121 67 664 222 701 462 252	1,666 4 52 566 235 594 192 23	2,377 109 58 646 201 723 459 181
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	203 13 14 166 1 9	255 32 39 174 10	334 3 294 37	303 7 268 28	4,735 505 227 3,617 45 341	5,095 554 276 3,988 56 221	3,673 491 298 2,526 28 330	4,741 630 339 3,511 33 228
Guam P.R. V.I. Amer. Samoa C.N.M.I.	4 - U	2 5 - U U	85 - - -	- 74 - U U	515 - U 14	26 652 U U	U U U U	U U U U

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending December 1, 2001, and December 2, 2000 (48th Week)*

N: Not notifiable. U: Unavailable. -: No reported cases. * Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

[†] Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

	NIC.	Shigel	losis [†]		Syl	aulaala		
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Culosis Cum.
Reporting Area	2001	2000	2001	2000	2001	2000	2001	2000
UNITED STATES	16,403	20,887	7,413	11,921	5,380	5,573	11,200	13,217
NEW ENGLAND Maine	257 6	383 10	274 3	362 11	59 1	80 1	373 3	390 16
N.H.	6	6	4	8	1	2	16	18
Mass.	194	268	184	245	35	57	219	222
R.I. Conn.	22 22	30 65	26 52	32 66	9 10	4 16	36 95	30 100
MID. ATLANTIC	1,157	2,579	713	1,634	444	259	2,137	2,107
N.Y. City	329	907	351	612	255	111	1,078	1,111
N.J. Pa.	185 185	489 457	184 65	421 390	129 36	65 73	458 264	501 195
E.N. CENTRAL	3,996	3,938	1,707	1,220	934	1,129	1,233	1,344
Ind.	2,727 216	382 1,481	42	305 151	147	331	243 100	133
III. Mich	497 286	1,125 633	288 215	130 579	318 376	394 294	571 243	652 228
Wis.	270	317	27	55	22	44	76	77
W.N. CENTRAL Minn.	1,857 435	2,318 752	1,255 440	1,931 842	80 28	62 15	423 214	484 153
lowa Mo	356 307	508 631	290 210	336 451	4 20	11 28	34 128	33 182
N. Dak.	21	42	34	49	-	-	3	2
S. Dak. Nebr.	583 86	/ 143	246	4 116	- 5	2	32	23
Kans.	69	235	35	133	23	6	-	75
Del.	2,327 15	2,784 24	12	1,109	1,826	1,862	2,315	2,658
Md. D.C.	144 53	182 77	91 U	109 U	236 34	291 36	211 51	232 35
Va.	440	434	175	339	102	123	241	240
N.C.	318	363	170	254	414	449	324	362
S.C. Ga.	244 368	134 247	120 130	89 181	212 348	210 360	164 437	238 561
Fla.	737	1,305	34	105	464	382	846	948
E.S. CENTRAL Ky.	1,505 699	1,126 488	570 300	550 111	607 43	812 80	749 109	834 113
Tenn.	97 201	337	110 130	362	309 125	485 115	273	307 279
Miss.	508	211	30	7	130	132	121	135
W.S. CENTRAL Ark	2,167 531	3,313 198	1,146 155	1,085 58	694 39	776 100	783 146	1,925 167
La.	145	277	166	182	160	199	-	213
Tex.	1,402	2,721	789	801	435	365	512	1,408
MOUNTAIN Mont.	906 8	1,175 7	675	814	216	215	467 14	473 17
Idaho	40	44	- 5	25	1	1	8	8
Colo.	233	252	255	205	22	8	112	74
N. Mex. Ariz.	115 383	157 514	79 275	108 326	17 159	16 183	24 210	40 199
Utah Nev.	58 66	76 120	53 8	81 66	8 8	1 5	33 63	41 90
PACIFIC	2,231	3,271	333	3,216	520	378	2,720	3,002
vvasn. Oreg.	202	434 163	167	399 109	43 13	60 11	103	235
Calif. Alaska	1,876 7	2,633 7	- 6	2,673	452	305	2,214 47	2,451 101
Hawaii	60	34	56	32	12	2	136	125
Guam P.R.	- 8	38 33	U U	U U	249	3 154	- 76	50 152
V.I. Amer. Samoa	Ū	Ū	U U	U U	Ū	Ū	Ū	Ū
C.N.M.I.	7	U	U	U	10	U	32	U

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending December 1, 2001, and December 2, 2000 (48th Week)*

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

 * Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

 * Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

	H. influ	ienzae,	Hepatitis (Viral), By Type				Measles (Rubeola)						
	Inva	asive	A		В		Indige	nous	Impo	rted⁺	Tota	l	
Reporting Area	Cum. 2001 [§]	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	2001	Cum. 2001	2001	Cum. 2001	Cum. 2001	Cum. 2000	
UNITED STATES	1,229	1,194	9,310	12,070	5,962	6,520	-	52	-	43	95	75	
NEW ENGLAND Maine N.H. Vt. Mass. R.I.	88 2 6 4 41 5	100 1 12 9 40 4	642 11 18 16 307 66	368 21 18 10 128 24	95 5 16 4 11 28	105 5 16 6 15 22		4 - 1 2 -		1 - - 1 -	5 - 1 3 -	6 - 3 - -	
Conn.	30	34	224	167	31	41	-	1	-	-	1	-	
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	179 71 46 42 20	217 93 59 38 27	886 256 281 159 190	1,421 240 479 273 429	917 121 397 169 230	1,082 123 527 168 264		5 1 3 - 1		11 4 1 1 5	16 5 4 1 6	21 10 10 - 1	
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	202 56 46 62 13 25	167 51 28 56 11 21	1,086 213 95 403 308 67	1,564 245 111 666 461 81	833 84 47 149 553	691 98 46 108 400 39				10 3 4 3 -	10 3 4 3 -	8 2 3 3 -	
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	64 39 - 15 7 - 2 1	74 42 - 22 2 1 3 4	385 41 37 103 3 32 166	622 171 63 248 3 2 32 32 103	201 28 25 103 1 1 25 18	274 35 32 134 2 1 44 26	- - - - -	4 2 - 2 - - - -		1 - - - - -	5 3 - 2 - - - -	2 1 - - - 1	
S. ATLANTIC	348	260	2,174	1,341	1,364	1,189	-	4	-	1	5	4	
Del. Md.	- 84	- 75	276	15 186	132	14 114	-	2	-	- 1	- 3	-	
D.C. Va. W. Va. N.C. S.C. Ga. Fla.	27 14 44 9 97 73	- 37 8 23 7 65 45	52 127 25 206 71 865 552	24 147 53 131 77 283 425	11 170 20 200 29 442 360	29 156 19 236 21 220 380		- - - 1 -		- - - - -	- - - 1 -	- 2 - - - 2	
E.S. CENTRAL Ky. Tenn. Ala. Miss.	70 2 40 26 2	47 12 21 12 2	366 123 145 71 27	369 47 133 48 141	390 41 216 79 54	439 73 204 60 102		2 2 - -		- - -	2 2 - -		
W.S. CENTRAL Ark. La. Okla. Tex.	47 1 6 39 1	63 2 16 43 2	1,273 66 61 113 1,033	2,272 127 90 245 1,810	661 95 45 106 415	1,026 92 144 148 642		- - - -		1 - - 1	1 - - 1		
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	133 2 - 37 22 54 8 10	123 1 4 1 31 24 45 11 6	687 11 57 7 86 37 365 68 56	870 7 33 4 196 69 424 59 78	449 3 11 3 99 128 136 26 43	496 6 3 95 131 184 24 47	-	2 - - - 1 - -			2 - - - - 1 - -	12 - - 2 - 3 7	
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	98 5 19 44 6 24	143 7 32 35 45 24	1,811 140 74 1,580 14 3	3,243 268 161 2,788 13 13	1,052 133 111 782 9 17	1,218 107 113 975 11 12		31 13 4 12 2		18 2 - 11 5	49 15 4 23 7	22 3 - 15 1 3	
Guam P.R. V.I. Amer. Samoa C.N.M.I.	- 1 - U	1 4 - U U	- 119 - U -	1 234 - U U	- 176 - U 35	10 275 - U U	- - U U	- - - U	- - U U	- - U -	- - U	2 - U U	

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending December 1, 2001, and December 2, 2000 (48th Week)*

N: Not notifiable. U: Unavailable. - : No reported cases. * Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date). [†] For imported measles, cases include only those resulting from importation from other countries. [§] Of 257 cases among children aged <5 years, serotype was reported for 121, and of those, 21 were type b.

	Mening Dis	jococcal ease	Mumps				Pertussis		Rubella		
Reporting Area	Cum. 2001	Cum. 2000	2001	Cum. 2001	Cum. 2000	2001	Cum. 2001	Cum. 2000	2001	Cum. 2001	Cum. 2000
UNITED STATES	2,041	2,009	3	201	299	82	4,394	6,600	-	20	165
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	106 4 13 6 54 6 23	118 8 12 3 68 9 18			4 - - 1 1 2	4 - 3 - 1	415 21 38 36 297 6 17	1,789 45 126 236 1,318 19 45			12 - 2 - 8 1 1
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	203 59 40 49 55	238 72 41 49 76	- - -	20 3 10 3 4	26 10 7 3 6	2 2 - -	265 133 44 18 70	654 326 82 30 216	- - -	5 1 3 1 -	9 1 8 -
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	295 75 37 70 66 47	362 87 41 82 110 42		19 1 3 11 4 -	22 7 1 6 2	7 2 1 2 2	605 235 80 71 132 87	774 318 111 113 111 121	- - - -	3 - 1 2 -	1 - - 1 -
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	145 22 29 49 6 5 20 14	146 21 34 66 2 5 7 11	1 - - - - 1	11 3 - 2 - 1 5	17 - 7 4 1 - 2 3	23 21 - - - 1	345 167 43 92 5 4 7 27	564 343 56 85 6 7 27 40		3 - 1 - - - 1	2 1 - - 1 -
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla	347 5 40 - 38 13 62 34 48 107	268 1 26 - 39 13 36 22 45 86	1 - - - - - -	37 - - 8 - 5 5 7 5	44 9 10 7 11 2 5	9 - - 8 - 1 - -	247 - 38 1 49 4 70 32 27 26	478 9 114 3 106 1 108 35 40 62		7 1 - - 2 1 3	112 1 - - 82 27 - 2
E.S. CENTRAL Ky. Tenn. Ala. Miss.	123 21 56 31 15	127 26 53 34 14	- - -	9 3 1 - 5	5 1 2 2	2 2 - -	149 52 57 36 4	108 55 32 18 3	- - -		6 1 1 4
W.S. CENTRAL Ark. La. Okla. Tex.	322 19 65 28 210	214 13 43 27 131	1 - - 1	14 1 2 - 11	32 3 5 - 24	14 - - 14	470 45 3 27 395	351 36 20 47 248	- - -	1 - - 1	8 1 - 6
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	88 4 7 5 34 10 13 8 7	94 4 7 1 32 11 29 7 3		13 1 1 3 2 1 1 3	21 1 1 1 1 4 6 6	19 - - 11 2 - - 6	1,248 37 170 1 283 137 509 76 35	755 35 61 453 88 75 24 15			2 - - 1 - 1 - -
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	412 60 41 295 3 13	442 56 66 303 9 8	- N - -	78 2 N 39 1 36	128 10 N 87 8 23	2 2 - - -	650 161 51 395 11 32	1,127 391 106 569 21 40	- - - -	1 - - - 1	13 7 6 -
Guam P.R. V.I. Amer. Samoa C.N.M.I.	- 4 - U	- 10 - U U	- - - U U	- - - U	16 - - U U	- - - U U	2 - U	4 9 - U U	- - - U U	- - - U	1 - - U U

TABLE III. (Cont'd) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending December 1, 2001, and December 2, 2000 (48th Week)*

N: Not notifiable. U: Unavailable. - : No reported cases. * Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

		All Cau	ises, By	Age (Ye	ears)		D9.11			All Cau	ises, By	Age (Y	ears)		D9.11
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. Springfield, Mass Waterbury, Conn.	417 U 34 18 30 U 19 16 ss. 42 . 54 67 7 . 27 24	327 28 14 28 14 28 U 12 9 38 42 42 6 18 22	56 U 4 3 2 U 6 2 3 5 12 12 4 1	23 U 2 1 - U - 5 1 4 6 - 2 1	8 U - - U 1 - 1 - 3	3 U - - U - - - - -	27 U 3 - 2 U - 1 4 4 - 5	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla. Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, F Tampa, Fla. Washington, D.C Wilmington, Del	1,304 143 131 142 203 163 61 61 47 180 . 180 . 100 . 12	851 79 83 96 132 116 43 35 34 45 124 55 9	280 34 28 29 39 28 11 16 8 10 40 34 3	115 19 17 12 18 17 3 2 4 11 9 -	26 4 1 2 8 1 1 2 1 1 3 2 -	32 7 2 3 6 1 3 5 2 1 2 -	84 3 13 15 20 12 1 4 5 7 -
Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Erie, Pa.§	79 2,556 51 21 94 42 16 51	62 1,699 33 17 62 28 11 40	13 532 11 4 23 8 5 5	1 227 5 - 6 5 - 6	2 62 1 - 2 1 -	1 33 1 - 1 - -	8 123 8 1 8 1 -	E.S. CENTRAL Birmingham, Ala Chattanooga, Te Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Al Nashville, Tenn.	937 a. 162 nn. 73 107 101 152 148 a. 40 154	645 109 55 78 66 103 96 31 107	200 37 13 20 26 32 34 7 31	50 8 3 5 3 9 12 2 8	21 3 2 3 - 7 4 - 2	20 4 - 1 6 1 2 - 6	75 18 3 7 13 14 6 7 7
Jersey City, N.J. New York City, 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.	29 (. 1,449 U 19 364 42 23 192 27 366 15 15 19 U	16 918 U 11 238 31 18 149 20 32 40 32 40 17 U	8 324 U 4 72 9 5 28 5 3 13 3 2 U	4 149 U 3 35 1 - 7 2 1 2 1 - U	- 37 U - 14 - - - 1 - U	1 18 U 1 5 1 - 2 1 - U	51 U 21 21 21 21 2 15 2 10 1 U	W.S. CENTRAL Austin, Tex. Baton Rouge, La Corpus Christi, 1 Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La. San Antonio, Te: Shreveport, La. Tulsa, Okla.	1,794 133 128 ex. 57 283 97 121 407 65 U x. 285 65 153	1,115 79 88 36 161 71 77 241 37 U 184 49 92	412 29 26 16 77 16 22 101 18 U 60 10 37	151 13 4 1 24 8 9 40 4 U 28 3 17	70 5 8 1 9 2 9 17 5 U 7 3 4	46 7 2 3 12 4 8 1 U 6 3	104 5 5 1 15 6 5 26 2 U 16 9 14
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Garand Rapids, Mic Indianapolis, Ind. Lansing, Mich.	2,032 63 47 U 119 146 211 160 236 55 82 39 ch. 82 239 62 221	1,423 33 U 84 96 139 127 135 62 21 55 169 42 83 83	389 9 U 24 34 45 20 5 9 11 8 14 50 10 23	124 3 U 8 11 15 9 22 3 5 5 4 9 6 9	41 3 U 2 3 4 3 5 1 2 4 1 3 2 4	55 1 U 1 2 8 1 16 - 2 1 8 8 1 2	137 85 U 9 11 7 12 4 2 4 85 11	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. PACIFIC Berkeley, Calif. Glendale, Calif.	1,203 M. 175 olo. 51 100 314 47 142 31 tah 141 161 1,922 12 131 28	804 110 33 42 60 217 35 8 22 88 119 1,345 11 86 20	255 40 7 5 23 79 9 36 6 25 25 354 1 26 5 5	87 14 1 3 11 15 2 13 2 17 9 133 - 17 3	26 4 - 3 3 - 7 1 4 4 4 2 - -	24 3 - 1 3 - 1 5 - 7 4 48 - 2 - 2	84 12 3 4 9 19 6 8 2 14 7 150 2 5 3 7
Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohi W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans Kansas City, Mo. Lincoln, Nebr. Minneapolis. Min	54 77 75 87 0 71 797 49 37 53 58 50 n. 149	38 61 66 62 58 580 42 30 43 30 43 38 109	11 10 8 18 11 127 7 4 15 7 7 26	3 4 1 4 2 42 - 1 4 5 3 8	1 2 - 22 - 2 1 3	1 - 2 - 26 - 2 2 1 3	86271 4684221 12	Long Beach, Cali Los Angeles, Cal Pasadena, Calif. Portland, Oreg. Sacramento, Cali San Diego, Calif. San Diego, Calif. San Jose, Calif. Santa Cruz, Calif Seattle, Wash. Spokane, Wash. Tacoma, Wash.	if. 77 if. 461 20 181 if. 142 176 alif. U 216 . 37 166 71 118	58 320 12 133 88 119 U 157 31 113 52 83	14 89 25 34 33 0 36 4 31 11 24	3 32 13 7 15 17 2 7 4 8	3 12 4 5 3 U 1 9 2 1	1 8 - 6 8 6 U 5 - 6 2 2	, 13 34 3 6 10 15 U 10 5 17 9 11
Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	93 116 87 105	73 70 76 67	9 24 6 22	1 10 2 8	5 7 1 2	5 5 2 6	5 - 6 6	TOTAL	12,962¶	8,789	2,605	952	318	287	830

TABLE IV. Deaths in 122 U.S. cities,* week ending December 1, 2001 (48th Week)

U: Unavailable. -:No reported cases.
* Mortality data in this table are reported voluntarily from 122 cities in the United States, most of which have populations of ≥100,000. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included.
* Pneumonia and influenza.
* Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks.
* Total includes unknown ages.

1098

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