

MORBIDITY AND MORTALITY

WEEKLY REPORT

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Influenza B Virus Outbreak on a Cruise Ship — Northern Europe, 2000

During June 23–July 5, 2000, an outbreak of respiratory illnesses occurred on the MS Rotterdam (Holland America Line & Windstar Cruises) during a 12-day Baltic cruise from the United Kingdom to Germany via Russia. The ship carried 1311 passengers, primarily from the United States, and 506 crew members from many countries. Although results of rapid viral testing for influenza A and B viruses were negative, immunofluorescence staining and viral culture results implicated influenza B virus infection as the cause of the outbreak. This report summarizes the findings of the outbreak investigation conducted by the ship's medical department and describes the measures taken to control the outbreak. Travelers at high risk for complications of influenza who were not vaccinated with influenza vaccine during the preceding fall or winter should consider receiving influenza vaccine before travel with large tourist groups at any time of year or to certain regions of the world.

On June 26, nine crew members presented to the ship's infirmary with cough, sore throat, and fever \geq 100.0 F (\geq 37.8 C). All had developed symptoms during the preceding 24 hours. Oropharyngeal specimens from two crew members were tested by a commercial rapid influenza diagnostic test designed to detect both influenza A and B viruses but not to distinguish between them. Although test results were negative, three crew members with high fevers were started on rimantadine therapy for clinically suspected influenza A infection.

To characterize and control the suspected outbreak among crew members, ship's medical staff implemented a respiratory illness protocol that included surveillance for cases of respiratory illness. A case of acute respiratory illness (ARI) was defined as cough or sore throat. Influenza-like illness (ILI), a subset of ARI cases, was defined as ARI with fever ≥ 100.0 F (≥ 37.8 C) or self-reported feverishness. Active surveillance was initiated among crew members. Supervisors on each work shift observed and asked crew members about symptoms of influenza and required any crew member with symptoms to report to the ship's infirmary for evaluation. Crew members with confirmed ILI were relieved of duty and placed in cabin isolation either alone or with other ill crew members. Passive surveillance was initiated among passengers and identified any passenger who presented to the ship's infirmary with respiratory illness. A commercial rapid influenza diagnostic test, designed to detect both influenza A and B viruses but not to distinguish between them, was used selectively to assist in diagnosis. Medical and demographic information, including country of residence, cabin number, and crew duties (if applicable), was collected from ill patients.

U.S. DEPARTMENT OF HEALTH & HUMAN SERVICES

Influenza B Virus — Continued

By June 29, 38 crew members and 26 passengers had been seen in the infirmary for ARI; of these, 32 (84%) crew members and 11 (42%) passengers had ILI. Eight crew members were tested by rapid influenza diagnostic testing; all had negative results. Because the etiology of crew respiratory illnesses remained uncertain, four symptomatic crew members disembarked in Stockholm, Sweden, for medical evaluation that included testing of nasopharyngeal specimens by immunofluorescence staining and viral culture. Two of four nasopharyngeal specimens tested positive for influenza B virus by immunofluorescence staining; one of the two specimens also was positive by culture. Neither of the two crew members diagnostic test. On the basis of immunofluorescence results, crew members on rimantadine therapy, which is effective only against influenza A infection, were advised to discontinue their medication. Oseltamivir, an antiviral agent that is effective against both influenza A and B infection, was sent to the ship for treatment of ill crew members and passengers.

A total of 64 (13%) crew members and 54 (4%) passengers were identified with ARI during the cruise. Of 63 crew members and 54 passengers with ARI for whom clinical information was known, 45 (71%) and 25 (46%), respectively, also had ILI (Figure 1). The median age of ill crew members was 32 years (range: 21–56 years) and of passengers, 68 years (range: 7–85 years). By cross-referencing crew duties, cabin locations of ill crew members and passengers, and dates of illness, medical staff identified the potential index case-patient as a 78-year-old U.S. passenger who boarded the ship ill with unconfirmed ILI after visiting London. She remained in her cabin except for occasional meals and did not seek medical attention until the fifth day of the cruise (June 28). Two of the 13 crew members with ILI, who were seen in the infirmary on June 25 and 26, were her cabin and dining room stewards. Both had worked, socialized, or shared cabins with other crew members who became ill. Surveillance among passengers and crew members was continued during the subsequent cruise and showed a decrease in the number of ARI and ILI cases.

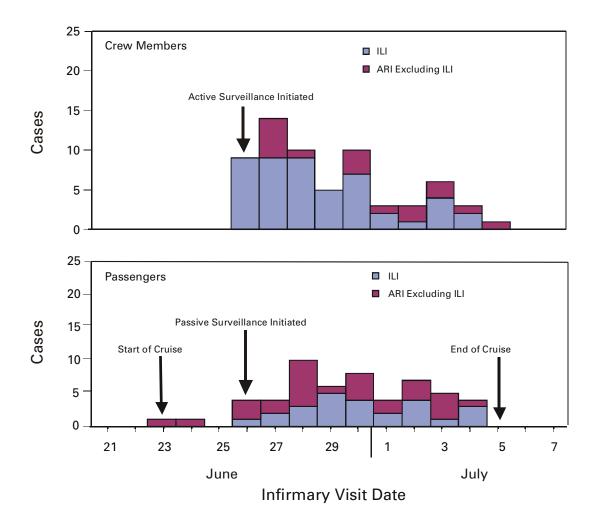
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Editorial Note: The findings of this investigation implicated influenza B virus as the cause of a respiratory illness outbreak onboard a cruise ship. Although the results of rapid viral testing for influenza A and B viruses were negative, influenza B infection was confirmed by viral culture and immunofluorescence antibody testing in two crew members. Although these tests were not performed on passengers, epidemiologic evidence suggested that respiratory illness cases among crew members and passengers were related and that an ill passenger might have transmitted infection to crew members.

Rapid viral diagnostic testing for influenza can be useful for patient management and influenza outbreak control. However, these tests are not as accurate in detecting influenza infection as viral culture (1). If an influenza outbreak is suspected, nasopharyngeal specimens should be collected simultaneously for rapid viral tests and viral isolation. Viral isolation is essential for identifying new or unusual strains of influenza and for selecting influenza vaccine strains.

Influenza B Virus — Continued

FIGURE 1. Acute respiratory illness (ARI) and influenza-like illness (ILI) among crew members and passengers, by infirmary visit date — MS Rotterdam, June 23–July 5, 2000



Influenza A outbreaks have been reported on cruise ships sailing in the Northern Hemisphere during the summer, but influenza B outbreaks have not been documented (2–7). Early suspicion of a potential influenza outbreak among crew members and rapid implementation of a respiratory illness control protocol probably limited the size of the outbreak. Key elements of the protocol included 1) implementation of active and passive surveillance using standard case definitions; 2) use of targeted rapid influenza diagnostic testing and viral cultures to confirm cases of influenza virus infection; 3) isolation of all crew members meeting the ILI case definition or those with confirmed influenza; 4) use of antiviral agents for treatment and, if indicated, for prophylaxis; and 5) monitoring of intervention results (8).

Influenza B Virus — Continued

Because influenza viruses usually are spread by droplets and aerosols produced by an infected person who is coughing or sneezing, isolation can limit the spread of infection in semienclosed environments such as cruise ships (2). Although the number of days crew members with ILI were isolated from noninfected crew members and passengers was not reported, isolation measures ideally should have covered the first 5 days of illness, a period based on the duration of influenza virus shedding in adults (8).

Summertime influenza outbreaks among passengers and crew members on cruise ships suggest that traveling in large groups can pose a risk for exposure to influenza viruses, even when the group is traveling in regions where influenza is not in seasonal circulation. Both passengers and crew members can serve as potential reservoirs of influenza infection. Travelers at high risk for complications of influenza (e.g., persons aged \geq 50 years, immunocompromised persons, and persons with chronic disorders of the pulmonary or cardiovascular systems) who were not vaccinated with influenza vaccine during the preceding fall or winter should consider receiving influenza vaccine before travel 1) with large organized tourist groups at any time of year; 2) to the tropics; or 3) to the Southern Hemisphere from April through September (the time of increased influenza activity in that hemisphere) (9). Cruise lines should attempt to achieve at least an 80% vaccination rate among crew members on each ship each year (8).

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Blood and Hair Mercury Levels in Young Children and Women of Childbearing Age — United States, 1999

Mercury (Hg), a heavy metal, is widespread and persistent in the environment. Exposure to hazardous Hg levels can cause permanent neurologic and kidney impairment (1–3). Elemental or inorganic Hg released into the air or water becomes methylated in the environment where it accumulates in animal tissues and increases in concentration

Blood and Hair Mercury Levels — Continued

through the food chain. The U.S. population primarily is exposed to methylmercury by eating fish. Methylmercury exposures to women of childbearing age are of great concern because a fetus is highly susceptible to adverse effects. This report presents preliminary estimates of blood and hair Hg levels from the 1999 National Health and Nutrition Examination Survey (NHANES 1999) and compares them with a recent toxicologic review by the National Research Council (NRC). The findings suggest that Hg levels in young children and women of childbearing age generally are below those considered hazardous. These preliminary estimates show that approximately 10% of women have Hg levels within one tenth of potentially hazardous levels indicating a narrow margin of safety for some women and supporting efforts to reduce methylmercury exposure.

CDC's NHANES is a continuous survey of the health and nutritional status of the U.S. civilian, noninstitutionalized population with each year of data constituting a representative population sample. 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 and hair samples, consisting of approximately 100 strands, were cut from the occipital position of the head of children aged 1-5 years and women aged 16–49 years. Whole blood specimens were analyzed for total Hg and inorganic Hg for children aged 1-5 years and women aged 16-49 years by automated cold vapor atomic absorption spectrophotometry in CDC's trace elements laboratory. The detection limit was 0.2 parts per billion (ppb) for total Hg and 0.4 ppb for inorganic Hg (4). Hairs of 0.6 inches (1.5 cm) closest to the scalp (approximately 1 month's growth) were analyzed for total Hg concentration using cold vapor atomic fluorescence spectroscopy (5). The limit of detection for total Hg in hair varied by analytic batch; the maximum limit of detection (0.1 parts per million [ppm]) was used in these analyses. Blood Hg levels less than the limit of detection were assigned a value equal to the detection limit divided by the square root of two for calculation of geometric mean values.

The geometric mean total blood Hg concentration for all women aged 16–49 years and children aged 1–5 years was 1.2 ppb and 0.3 ppb, respectively; the 90th percentile of blood Hg for women and children was 6.2 ppb and 1.4 ppb, respectively (Table 1). Almost all inorganic Hg levels were undetectable; therefore, these measures indicate blood

	G	eometr	ic	Selected percentiles (95% CI*)								
	No.	mean	(95% CI)	10th	25th	50th	75th	90th				
Blood Hg [†]												
Children	248	0.3	(0.2–0.4)	<lod<sup>§</lod<sup>	<lod< td=""><td>0.2 (0.2-0.3)</td><td>0.5 (0.4–0.8)</td><td>1.4 (0.7–4.8)</td></lod<>	0.2 (0.2-0.3)	0.5 (0.4–0.8)	1.4 (0.7–4.8)				
Women	679	1.2	(0.9–1.6)	0.2 (0.1–0.3)	0.5 (0.4–0.7)	1.2 (0.8–1.6)	2.7 (1.8–4.5)	6.2 (4.7–7.9)				
Hair Hg [¶]												
Children	338	**	×	<lod< td=""><td><lod< td=""><td><lod< td=""><td>0.2 (0.1–0.4)</td><td>0.4 (0.3–1.8)</td></lod<></td></lod<></td></lod<>	<lod< td=""><td><lod< td=""><td>0.2 (0.1–0.4)</td><td>0.4 (0.3–1.8)</td></lod<></td></lod<>	<lod< td=""><td>0.2 (0.1–0.4)</td><td>0.4 (0.3–1.8)</td></lod<>	0.2 (0.1–0.4)	0.4 (0.3–1.8)				
Women	702			<lod< td=""><td><lod< td=""><td>0.2 (0.2–0.3)</td><td>0.5 (0.4–0.8)</td><td>1.4 (0.9–1.7)</td></lod<></td></lod<>	<lod< td=""><td>0.2 (0.2–0.3)</td><td>0.5 (0.4–0.8)</td><td>1.4 (0.9–1.7)</td></lod<>	0.2 (0.2–0.3)	0.5 (0.4–0.8)	1.4 (0.9–1.7)				

TABLE 1. Selected percentiles and geometric means of blood and hair mercury (Hg) concentrations for children aged 1–5 years and women aged 16–49 years — National Health and Nutrition Examination Survey, United States, 1999

* Confidence interval.

[†] Parts per billion.

[§] Limit of detection.

[¶] Parts per million.

** Not calculated. Proportion <LOD too high to be valid.

Blood and Hair Mercury Levels — Continued

methylmercury levels. The 90th percentile of hair Hg for women and children was 1.4 ppm and 0.4 ppm, respectively. Geometric mean values were not calculated for hair Hg values.

Reported by: Center for Food Safety and Applied Nutrition, Food and Drug Administration. US Environmental Protection Agency. National Energy Technology Laboratory, Dept of Energy. National Marine Fisheries Laboratory, National Oceanic and Atmospheric Administration. National Center for Health Statistics; National Center for Environmental Health, CDC.

Editorial Note: The NHANES1999 blood and hair Hg data are the first nationally representative human tissue measures of the U.S. population's exposure to Hg. Previous estimates of methylmercury exposure in the general population were based on exposure models using fish tissue Hg concentrations and dietary recall survey data (1). The NRC review provided guidance to the Environmental Protection Agency (EPA) for developing an exposure reference dose for methylmercury (i.e., an estimated daily exposure that probably is free of risk for adverse effects over the course of a person's life) (3). The NRC report recommended statistical modeling of results from an epidemiologic study conducted in the Faroe Islands near Iceland, where methylmercury exposures are high because of the large amount of seafood eaten by the local population. Results of this study were used to calculate a benchmark dose (BMD), an estimate of a methylmercury exposure in utero associated with an increase in the prevalence of abnormal scores on cognitive function tests in children. The lower 95% confidence limit of the BMD (BMDL*) was recommended to calculate the EPA reference dose. The NRC committee recommended a BMDL of 58 ppb Hg in cord blood (corresponding to 12 ppm Hg in maternal hair) (3). In the NHANES 1999 sample, there were no measurements of blood values \geq 58 ppb or hair values \geq 12 ppm. A margin-of-exposure analysis (i.e., an evaluation of the ratio of BMDL to estimated population exposure levels) showed ratios of <10 when comparing BMDL with NHANES 1999 estimates of the 90th percentile for blood and hair Hg levels in women of childbearing age. Margin-of-exposure measures of this magnitude indicate a narrow margin of safety (3) and suggest that efforts aimed at decreasing human exposure to methylmercury should continue.

The findings in this study are subject to at least three limitations. First, the ratio of Hg in cord and maternal blood is uncertain. The NRC committee summarized some studies that suggest that cord blood values may be 20%–30% higher than corresponding maternal blood levels. However, other studies suggest that the ratio is closer to 1:1 (*3*); therefore, the NHANES values may not be directly comparable to BMDL recommended by NRC. Second, NHANES cannot provide estimates of Hg exposure in certain highly exposed groups (e.g., subsistence fishermen and others who eat large amounts of fish). Published data from studies of highly exposed U.S. populations indicated that some persons attain Hg tissue levels above BMDL (*1*). Third, the sample size of NHANES 1999 was small and the 1999 survey was conducted in only 12 locations. More data are needed to confirm these findings.

^{*}A BMD of 85 ppb Hg in cord blood or 17 ppm Hg in maternal hair was estimated to result in an increase in the proportion of abnormal scores on the Boston Naming Test for children exposed in utero from an estimated background prevalence of 5% to a prevalence of 10% (6). BMDL recommended by NRC is the lower 95% confidence bound of the BMD.

Blood and Hair Mercury Levels — Continued

The long-term strategy for reducing exposure to Hg is to lower concentrations of Hg in fish by limiting Hg releases into the atmosphere from burning mercury-containing fuel and waste and from other industrial processes. On the basis of data from EPA's National Toxics Inventory, air emissions of Hg decreased approximately 21% during 1990–1996, largely because of regulations for waste incineration (7). EPA expects this trend to continue as regulations are implemented for waste incineration and chlorine production facilities and are developed for electric power utilities (8,9). Fish is high in protein and nutrients and low in saturated fatty acids and cholesterol and should be considered an important part of the diet. The short-term strategy to reduce Hg exposure is to eat fish with low Hg levels and to avoid or to moderate intake of fish with high Hg levels. Statebased fish advisories and bans identify fish species contaminated by Hg and their locations and provide safety advice (http://www.epa.gov/ost/fish[†]). The Food and Drug Administration advises that pregnant women and those who may become pregnant should not eat shark, swordfish, king mackerel, and tile fish known to contain elevated levels of methylmercury. Information is available at http://www.fda.gov/bbs/ topics/ANSWERS/2001/advisory.html⁺.

U.S. population estimates of Hg tissue levels by race/ethnicity, region, and fish consumption will become available after 2 additional years of NHANES data collection. NHANES will provide the opportunity to measure tissue Hg levels and to monitor the effectiveness of continuing efforts to reduce methylmercury exposure in the U.S. population.

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[†] 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.

Progress Toward Poliomyelitis Eradication — Afghanistan, 1999–2000

In 1988, the World Health Assembly of the World Health Organization (WHO) resolved to eradicate poliomyelitis globally by 2000. During the same year, the Eastern Mediterranean Region* (EMR) of WHO passed a resolution to join the global initiative. Since then, substantial progress has been made worldwide and in EMR member countries (*1,2*). Afghanistan, with ongoing civil conflict, initiated polio eradication activities in 1994. Since then, a countrywide surveillance system for acute flaccid paralysis (AFP) was established and National Immunization Days (NIDs)[†] were implemented (*3*). This report summarizes the achievements toward polio eradication in Afghanistan during 1999–2000.

Routine Vaccination

In 1996, an estimated 30% of infants aged <1 year had received three doses of oral poliovirus vaccine (OPV) (3). In 1998, a review of the Expanded Program on Immunization (EPI) documented wide variations in vaccination coverage by geographic area; levels were particularly low in the north as a result of civil conflict. In 1999, EPI acceleration campaigns provided vaccinations to 82,000 unvaccinated children aged <2 years. In 2000, a comprehensive 5-year plan was drafted to set targets and strategies for the coming years.

Supplemental OPV Vaccination

During 1994–1996, supplemental vaccination activities against polio began with multivaccine subnational campaigns that delivered diphtheria and tetanus toxoids and pertussis vaccine, OPV, and measles vaccine to children aged <5 years. NIDs using OPV were initiated during April–May 1997, and since have been conducted annually. High coverage was achieved during four NID rounds in 1999 and another four in 2000 (Table 1). Of 330 districts in Afghanistan, 325 were reached during the fall 1999 NIDs. During the spring 2000 NIDs, all districts were reached except two north of the capital (Kabul) where most of the population had left the area because of ongoing civil conflict. Supplemental vaccination activities in Afghanistan have been coordinated with neighboring countries, particularly Iran and Pakistan. Because surveillance data indicate that Afghanistan and Pakistan are one epidemiologic block, supplemental campaigns have been conducted simultaneously in both countries when possible. Since the fall of 1999, careful district level NID planning and well-supervised house-to-house vaccination have led to incremental improvements in the quality and coverage of each NID.

AFP Surveillance

In 1997, 37 AFP sentinel reporting sites were established. Since then, surveillance has expanded to 234 sites with emphasis on areas with high population density. In 2000, Afghanistan exceeded the WHO established target for a nonpolio AFP rate indicative of sensitive surveillance (i.e., \geq 1.0 per 100,000 population aged <15 years) with a rate of 1.2 (Table 1). During 1999–2000, the number of AFP cases increased from 230 to 253, and the number of wild polioviruses isolated from AFP cases decreased from 63 to 28 (Figure 1). The

^{*}Djibouti, Egypt, Libya, Morocco, Somalia, Sudan, and Tunisia in northern and eastern Africa; Bahrain, Kuwait, Oman, Qatar, Saudi Arabia, United Arab Emirates, and Yemen in the Arabian peninsula; Iraq, Jordan, Lebanon, Syria, and the Palestinian National Authority in the Middle East; Afghanistan, Iran, and Pakistan in Asia; and Cyprus.

[†]Mass campaigns over a short period (days to weeks) in which two doses of OPV are administered to all children in the target group (usually aged 0–4 years), regardless of vaccination history, with an interval of 4–6 weeks between doses.

Poliomyelitis Eradication — Continued

Surveillance indicators	NID round	1999	2000
AFP cases		230	253
Nonpolio AFP rate [†]		0.66	1.22
Confirmed poliomyelitis cases		150	103
Confirmed wild poliovirus cases		63	28
Percentage of persons with AFP			
with adequate stool samples [§]		53%	50%
No. children vaccinated	1	4,026,094	5,155,049
	2	4,293,368	5,250,648
	3	4,610,861	5,704,009
	4	4,220,681	5,761,400

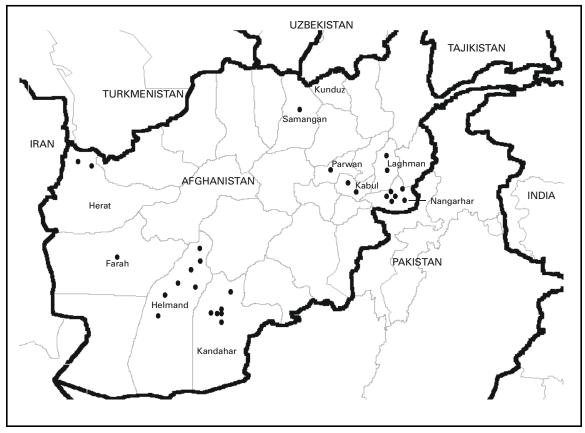
TABLE 1. Acute flaccid paralysis (AFP) surveillance and National Immunization
Day (NID)* coverage — Afghanistan, 1999 and 2000

* Mass campaigns over a short period (days to weeks) in which two doses of oral poliovirus vaccine are administered to all children in the target group (usually aged 0–4 years), regard-less of vaccination history, with an interval of 4–6 weeks between doses.

[†] Number of nonpolio AFP case-patients per 100,000 population aged <15 years.

[§] Two stool specimens, collected 24 to 48 hours apart within 14 days of onset of paralysis, that arrive in the laboratory in good condition.

FIGURE 1. Location of poliomyelitis cases* confirmed through wild poliovirus isolation — Afghanistan, 2000[†]



* n=28.

⁺ As of February 26, 2001.

Poliomyelitis Eradication — Continued

National Institute of Health (NIH), Islamabad, Pakistan, has provided laboratory support for the Afghanistan program. All stool specimens are flown from Afghanistan to Islamabad on United Nations' flights and transported to the NIH laboratory.

A remaining challenge is the timely collection of adequate stool specimens[§] from AFP case-patients. In 2000, 50% of AFP cases reported nationally had adequate stool specimens, which was substantially short of the WHO target of 80%. This low level is partly the result of AFP being identified late in patients' illness, which precludes the collection of stool specimens soon after paralysis onset. Intensified efforts are being made to improve surveillance quality by the immediate investigation of all AFP cases and weekly active surveillance visits to major hospitals and shrines.

Reported by: Afghanistan Country Office, World Health Organization, Islamabad, Pakistan. Eastern Mediterranean Regional Office, World Health Organization, Cairo, Egypt. Vaccines and Biologicals Dept, World Health Organization, Geneva, Switzerland. Respiratory and Enteric Viruses Br, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases; Vaccine Preventable Disease Eradication Div, National Immunization Program, CDC.

Editorial Note: Although polio remains endemic in Afghanistan, progress during 1999–2000 demonstrates that key strategies can be implemented successfully in countries experiencing internal strife. During 1999–2000, the nonpolio AFP rate almost doubled and the number of districts reached by NIDs increased steadily. Careful planning and supervision of house-to-house vaccination and support from an increasing number of local partners resulted in the largest number of children ever being reached. Monitoring by nongovernment organizations, United Nations' agencies, and local authorities has increased the quality of NIDs. During the spring 2000, the days of tranquility were respected by all warring factions and their local commanders, greatly facilitating the implementation of NIDs.

Civil conflict, massive population shifts (returning refugees and traditional nomadic movements), a drought, rebuilding the public health infrastructure, geographic barriers, extreme climate, and the need to access areas that can be reached only by several days' travel on muleback are some of the obstacles facing eradication efforts in Afghanistan. Until 2000, negotiated cease-fires and days of tranquility agreements during NIDs had been only partly successful. Cessation of polio vaccination activities in mid-1997 in northern Afghanistan as a result of ongoing conflict may have facilitated the large polio outbreak that occurred in Kunduz province in 1999 (*4*).

Innovative measures and local peace initiatives will continue to be needed to create opportunities for reaching and vaccinating isolated populations. Afghanistan is preparing the implementation of five NID rounds in 2001. Plans are being developed to conduct focal mass campaigns in large, high-risk areas during the summer of 2001. Improved and timely stool specimen collection from AFP case-patients will be necessary to obtain data for targeting these campaigns and eliminating the last reservoirs of poliovirus circulation. Meeting these challenges will require the continued support of polio eradication partners[¶].

[§] Two stool specimens collected 24 to 48 hours apart within 14 days of onset of paralysis that arrive in the laboratory in good condition.

[¶] Polio eradication in Afghanistan is supported by the national government. External support is provided by global polio eradication partners, including Rotary International, United Nations Children's Fund (UNICEF), WHO, the governments of the United States, Great Britain, Denmark, Norway, Netherlands, Sweden, Luxemburg, Germany, and the European Community.

Poliomyelitis Eradication — Continued

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Public Health Dispatch

Outbreak of Poliomyelitis — Dominican Republic and Haiti, 2000–2001

During July 12, 2000–February 8, 2001, 12 laboratory-confirmed poliomyelitis cases attributed to vaccine-derived poliovirus type 1 were identified in the Dominican Republic (1). Of these, 11 (92%) case-patients were aged \leq 6 years (range: 9 months–14 years), and the date of paralysis onset of the last case was January 2, 2001. All case-patients were inadequately vaccinated or unvaccinated. In Haiti, one confirmed polio case attributed to vaccine-derived type 1 poliovirus was reported in an unvaccinated child aged 2 years with paralysis onset on August 30, 2000. As of February 21, 33 acute flaccid paralysis (AFP) cases from the Dominican Republic and three AFP cases from Haiti were pending final classification.

Extensive control efforts are under way. The Dominican Republic held nationwide mass vaccination campaigns with oral poliovirus vaccine (OPV) in December 2000 and February 2001, with a third round planned for April 2001. All children aged <5 years are being targeted, with approximately 1.2 million OPV doses given in the first campaign. AFP surveillance has been strengthened with intensification of active case-finding and weekly reporting. Haiti has initiated regional OPV campaigns to be conducted approximately every 2 months.

Travelers to the Dominican Republic and Haiti who are not vaccinated adequately are at risk for polio. All travelers should be vaccinated against polio according to national vaccination policies $(2)^*$.

Reported by: Ministry of Health, Pan American Health Organization, Santo Domingo, Dominican Republic. Ministry of Health, Pan American Health Organization, Port-au-Prince, Haiti. Caribbean Epidemiology Center Laboratory, Pan American Health Organization, Trinidad and Tobago. Div of Vaccines and Immunization, Pan American Health Organization, Washington, DC. Respiratory and Enteric Viruses Br, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases; Vaccine Preventable Disease Eradication Div, National Immunization Program, CDC.

^{*}Recommendations for children in the United States include a 4-dose vaccination series with inactivated poliovirus vaccine (IPV) at ages 2, 4, 6–18 months, and 4–6 years. Unvaccinated adults should receive three doses of IPV, the first two doses at intervals of 4–8 weeks and the third dose 6–12 months after the second. If three doses cannot be administered within the recommended intervals before protection is needed, alternative schedules are proposed. For incompletely vaccinated persons, additional IPV doses are recommended to complete a series. Booster doses of IPV may be considered for persons who previously have completed a primary series of polio vaccination and who may be traveling to areas where polio is endemic.

Public Health Dispatch — Continued

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Notice to Readers

International Course in Applied Epidemiology

CDC and Emory University's Rollins School of Public Health will co-sponsor a course, "International Course in Applied Epidemiology" during September 24–October 19, 2001, in Atlanta, Georgia. This basic course in epidemiology is directed at public health professionals from countries other than the United States.

The course's content includes presentations and discussions of epidemiologic principles, basic statistical analysis, public health surveillance, field investigations, surveys and sampling, and discussions of the epidemiologic aspects of current major public health problems in international health. Included are small group discussions of epidemiologic case exercises based on field investigations. Participants are encouraged to give a short presentation reviewing some epidemiologic data from their own country. Computer training using Epi Info 2000 (Windows® version), a software program developed at CDC and the World Health Organization for epidemiologists, is included. Prerequisites are familiarity with the vocabulary and principles of basic epidemiology or completion of CDC's "Principles of Epidemiology" home-study course (SS3030) or equivalent. Preference will be given to applicants whose work involves priority public health problems in international health. Early registration deadline is June 1, 2001; late registration deadline is September 1, 2001. There is a tuition charge.

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; World-Wide Web site, http://www.sph.emory.edu/EPICOURSES*; or e-mail pvaleri@sph.emory.edu.

^{*}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.

Notices to Readers — Continued

Notice to Readers

Introduction to Public Health Surveillance Course

CDC and Emory University's Rollins School of Public Health will co-sponsor a course, "Introduction to Public Health Surveillance" during June 18–22, 2001, in Atlanta, Georgia. The course is designed for state and local public health professionals.

The course will provide practicing public health professionals with the theoretical and practical tools necessary to design, implement, and evaluate effective surveillance programs. Topics include overview and history of surveillance systems; planning considerations; sources and collection of data; analysis, interpretation, and communication of data; surveillance systems technology; ethics and legalities; state and local concerns; and future considerations. There is a tuition charge.

Deadline for application is May 4. Additional information and applications are available from Emory University, International Health Dept.(PIA), 1518 Clifton Road N.E., Room 746, Atlanta, GA 30322; telephone (404) 727-3485; fax (404) 727-4590; or World-Wide Web site, http://www.sph.emory.edu/EPICOURSES*; or e-mail pvaleri@sph.emory.edu.

Erratum: Vol. 50, No. 7

In the article, "Prevalence of Disabilities and Associated Health Conditions Among Adults—United States, 1999," in the first full paragraph on page 121 in the sentence that begins "Of the total percentage of disabilities, 63% occurred among working adults," the age range should read "aged *18*–64" years.

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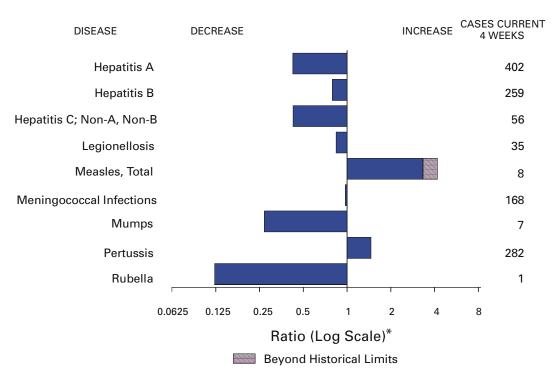


FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending February 24, 2001, with historical data

* 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.

		Cum. 2001		Cum. 2001
Anthrax			Poliomyelitis, paralytic	-
Brucellosis*		-	Psittacosis*	2
Cholera		-	O fever*	1 1
Cyclosporiasis	*	4	Rabies, human	-
Diphtheria			Rocky Mountain spotted fever (RMSF)	9
Ehrlichiosis:	human granulocytic (HGE)*	3	Rubella, congenital syndrome	-
2	human monocytic (HME)*	1	Streptococcal disease, invasive, group A	365
Encephalitis:	California serogroup viral*	-	Streptococcal toxic-shock syndrome*	13
2	eastern equine*	-	Syphilis, congenital [¶]	1
	St. Louis*	-	Tetanus	1
	western equine*	-	Toxic-shock syndrome	14
Hansen diseas		2	Trichinosis	2
	Imonary syndrome**	1 1	Tularemia*	1
	mic syndrome, postdiarrheal*	5	Typhoid fever	15
HIV infection,		10	Yellow fever	-
Plague	r	-		

TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending February 24, 2001 (8th Week)

-: 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).

³Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP). Last update January 30, 2001.

¹Updated from reports to the Division of STD Prevention, NCHSTP.

	AID	ne l	Chlam	vdiat	Cruntos	ooridiosis	NET		<i>coli</i> O157:H7 PH	
Demonstrum Arrow	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
Reporting Area	2001⁵ 2,792	2000 4,895	2001 77,539	2000 95,997	2001 132	2000 156	2001 115	2000 193	2001 56	2000 162
NEW ENGLAND	91	497	2,976	3,510	5	7	13	15	7	18
Maine N.H.	3 5	3 6	151	209 162	-	1	- 4	1 3	2	1 4
/t. Mass.	5 51	360	96 1,405	88 1,489	2	4 2	- 9	1 5	- 5	2 4
R.I. Conn.	11 16	17 111	471 853	370 1,192	- 1 2	-	-	- 5	-	- 7
MID. ATLANTIC	555	1,283	3,613	7,556	9	15	9	23	6	38
Jpstate N.Y. N.Y. City	4 360	60 770	N 1,870	N 3,826	3 6	8 4	9	21 1	6	31
۷.J.	157	300	308	2,020	-	-	-	1	-	2
	34	153	1,435	1,710	-	3	N	N 21	-	5
E.N. CENTRAL Dhio	224 46	545 85	9,867 214	17,380 4,696	43 17	36 6	23 11	31 5	11 6	8 3
nd. II.	26 121	28 352	1,898 2,576	1,975 5,179	9	3 5	4 4	1 14	- 3	1
Mich.	23	67	4,051	3,027	17	3	2	6	-	2
Vis. V.N. CENTRAL	8 44	13 96	1,128 3,847	2,503 5,562	- 4	19 4	2 14	5 39	2 9	2 32
Minn.	12	31	805	1,286	-	-	3	5	4	12
owa Mo.	9 7	7 23	442 1,185	403 2,066	2	-	2 6	8 19	- 2	4 8
N. Dak.	-	- 1	109	159	-	1 1	- 1	2	- 1	2
S. Dak. Nebr.	6	4	279 201	267 475	2	2	-	3	-	- 4
Kans. 6. ATLANTIC	10 734	30 1,220	826 16,080	906 18,317	- 21	- 17	2 19	2 17	2 4	2 16
Del.	15	15	437	450	-	-	-	-	-	-
Иd. Э.С.	41 62	136 24	1,697 446	1,603 448	2 2	1	-	5	Ū	1 U
/a.	48	75	2,176	1,971	2	-	2	3	3	5
V.Va. N.C.	6 57	5 71	321 2,383	308 2,757	- 4	3	- 13	1 5	- 1	1 2
S.C. Ga.	61 104	107 98	1,260 3,105	2,726 3,776	-	-7	1 1	- 1	-	- 3
-la.	340	689	4,255	4,278	11	6	2	2	-	4
E.S. CENTRAL Ky.	148 18	168 36	6,780 1,324	6,409 1,177	3	5	5	10 4	3 2	7 2
lenn.	80	35	2,232	1,816	-	-	2	3	1	5
Ala. Miss.	25 25	50 47	1,533 1,691	1,920 1,496	2 1	5	3	1 2	-	-
V.S. CENTRAL	409	524	14,364	15,087	4	11	2	11	8	19
Ark. _a.	19 130	20 83	1,387 2,707	605 2,748	2 1	1 -	-	2	- 5	3
Okla. Tex.	20 240	17 404	1,599 8,671	1,393 10,341	1	1 9	2	3 6	2 1	3 8
NOUNTAIN	145	178	3,631	5,438	14	11	12	23	5	7
Mont. daho	1 -	3 3	148 292	185 296	2	- 1	2	5 3	-	-
Nyo. Colo.	- 38	1 52	117 247	129 1,356	- 6	1 3	- 6	2 8	- 2	2 2
N. Mex.	38 7	25	604	709	3	1	-	-	-	-
Ariz. Jtah	52 11	22 28	1,718 67	1,816 344	1 2	2 3	4	3 1	2 1	2 1
Nev.	36	44	438	603	-	-	-	1	-	-
PACIFIC Wash.	442 26	384 46	16,381 2,123	16,738 1,921	29 N	50 U	18 3	24 1	3	17 7
Dreg.	17	11	675	454	6	1	3	4	1	4
Calif. Alaska	398 1	303	12,862 299	13,480 339	23	49	12	15	-	3
Hawaii	-	24	422	544	-	-	-	4	2	3
Guam P.R.	2 48	6 116	436	- U	-	-	N -	N -	U U	U U
/.I. Amer. Samoa	1	-	U U	Ŭ U	U U	U U	U U	U U	Ŭ	Ŭ U
C.N.M.I.	-		Ŭ	Ŭ	Ŭ	Ŭ	Ŭ	Ŭ	Ŭ	Ŭ

 TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending February 24, 2001, and February 26, 2000 (8th Week)

N: Not notifiable. 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. * Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention. Last update January 30, 2001.

	Gonor	rhea	Hepatit Non-A, N	is C; Ion-B	Legione	llosis	Listeriosis	Lyı Dise	
Reporting Area	Cum. 2001⁵	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2001	Cum. 2000
UNITED STATES	37,055	49,703	186	532	73	96	39	252	547
NEW ENGLAND	855	1,057	2	2	1	7	5	84	95
Лаine N.H.	- 15	10 16	-	-	-	2	-	42	- 12
/t. Aass.	14 430	4 423	2	- 2	1	- 4	- 3	-7	- 14
R.I.	123	87	-	-	-	-	-	-	-
Conn.	273	517	-	-	-	1	2	35	69 25 C
MID. ATLANTIC Jpstate N.Y.	2,351 594	3,865 443	10 7	78 1	2 1	11 3	1 1	90 65	356 65
N.Y. City N.J.	925 207	1,599 1,119	-	- 72	-	-	-	-	14 47
Pa.	625	704	3	5	1	8	-	25	230
.N. CENTRAL	5,084	10,333	26	50	26	36	5	9	14
Dhio nd.	168 871	2,751 911	1 -	-	15 3	15 4	2	9	2
ll. Mich.	1,238 2,380	3,577 1,964	- 25	5 45	- 8	3 7	- 3	-	1
Wis.	427	1,130	-	-	-	7	-	U	11
W.N. CENTRAL	1,715	2,268	37	73	7	4	2	5	8
Vlinn. owa	271 130	488 111	-	-	- 2	1 1	-	3	2
Ио. N. Dak.	844 4	1,102 6	36	70	3	2	1	2	2
S. Dak.	32	36	-	-	-	-	-	-	-
lebr. Kans.	43 391	149 376	- 1	1 2	1 1	-	- 1	-	- 4
S. ATLANTIC	10,411	14,300	6	9	13	20	6	49	61
Del. Ad.	251 1,016	239 1,093	2	2	- 5	1 7	- 1	- 44	8 45
D.C.	480	376	-	-	-	-	-	1	-
/a. V. Va.	1,231 57	1,445 87	-	- 1	2 N	3 N	1 1	2	1 3
N.C. S.C.	1,968 1,338	1,901 3,529	2	5	2	1 2	-	2	4
Ga.	1,626	2,502	- 2	- 1	- 4	- 6	1 2	-	-
⁻ la. E.S. CENTRAL	2,444	3,128	2		4	2	2 4	2	-
ζу.	4,514 566	4,753 505	-	79 5	2	-	1	2	-
Fenn. Ala.	1,582 1,275	1,477 1,583	7	17 3	- 1	1 1	2 1	-	-
Miss.	1,091	1,188	16	54	-	-	-	-	-
V.S. CENTRAL	7,435	8,071	53	173	1	4	-	-	2
Ark. .a.	935 1,894	332 2,130	1 5	97	- 1	- 2	-	-	- 2
Okla. Tex.	791 3,815	639 4,970	- 47	- 76	-	-2	-	-	-
MOUNTAIN	1,103	1,498	13	39	4	5	3	-	-
Nont.	5 18	17	-	-	-	-	-	-	-
daho Vyo.	12	12	3	25	-	-	-	-	-
Cólo. N. Mex.	318 117	537 135	4 5	6 4	3	2	1 1	-	-
Ariz.	472	541	-	4	1	-	1	-	-
Jtah Nev.	9 152	51 205	-	-	-	2	-	-	-
ACIFIC	3,587	3,558	16	29	16	7	13	13	11
Vash. Dreg.	485 135	369 56	2 3	2 8	3 N	2 N	- 1	- 2	- 1
Calif.	2,862	3,031	11	19	13	5	12	11	10
Alaska Hawaii	33 72	34 68	-	-	-	-	-	Ň	Ň
Guam	-	-	-	-	-	-	-	-	-
P.R. /.I.	126 U	76 U	Ū	1 U	2 U	Ū	-	N U	N U
Amer. Samoa	U	U	U	Ŭ	U	U	-	U	U
C.N.M.I.	U	U	U	U	U	U	-	U	U

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending February 24, 2001, and February 26, 2000 (8th Week)

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

					<u> </u>	Salmor	nellosis*	
		laria		s, Animal		TSS		LIS
Reporting Area	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000
UNITED STATES	106	127	492	622	2,394	3,410	1,646	3,193
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Const	10 - - 3 - 7	3 - - 3 -	66 11 2 13 16 8	64 14 1 4 23 4	198 9 16 10 121 11	205 17 12 5 133 3	141 7 10 9 64 18	225 10 12 5 138 14
Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	6 4 2 -	- 23 7 10 3 3	16 84 66 1 17	18 106 80 U 13 13	31 160 58 77 - 25	35 482 59 141 175 107	33 221 64 96 27 34	46 557 130 166 98 163
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	22 4 7 11	16 2 - 9 5 -	3 - 1 - 2 -	5 1 - - 4	391 153 31 88 82 37	526 139 36 183 74 94	301 74 19 100 77 31	263 90 55 1 78 39
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	3 1 1 - - -	7 2 - 1 - 1 3	44 11 11 2 8 6 - 6	62 18 6 2 8 16 - 12	159 31 23 53 1 13 9 29	155 27 12 54 2 6 20 34	135 53 1 59 2 7 7 13	184 62 13 49 13 12 14 21
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	26 1 11 2 8 - 1 - 3	30 17 7 4 - 2	213 - 43 - 15 56 7 24 25	202 7 42 55 15 57 13 - 13	660 12 90 13 76 3 152 56 106 152	514 8 97 - 48 17 115 55 55 119	366 8 78 U 48 11 45 50 126	512 11 89 0 63 11 80 47 159 52
E.S. CENTRAL Ky. Tenn. Ala. Miss.	5 - 3 2 -	4 1 - 3	4 2 2	25 5 17 3	182 36 41 83 22	178 29 42 63 44	63 21 39 3	126 19 59 40 8
W.S. CENTRAL Ark. La. Okla. Tex.	2 - 1 - 1	1 - 1 -	10 - 10 -	105 - 8 97	74 30 9 14 21	310 28 38 22 222	139 13 40 15 71	369 22 61 26 260
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	8 1 - 3 1 1 1	8 - - 4 - 2 2	27 4 9 - 1 13 -	27 9 - 13 - 1 4 -	200 7 6 7 53 28 67 21 11	296 11 21 5 69 28 89 45 28	128 4 1 34 10 59 20	250 13 3 59 30 96 49
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	24 1 4 18 1	35 2 5 27 1	41 - 24 17	26 - - 21 5 -	370 18 35 313 4	744 23 45 625 11 40	152 31 85 36	707 88 56 520 10 33
Guam P.R. V.I. Amer. Samoa C.N.M.I.	- - U U U	2 U U U	11 U U U	7 U U U	5 U U U	36 U U U	U U U U	

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States,
weeks ending February 24, 2001, and February 26, 2000 (8th Week)

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).

		Shige		•	Sv	philis			
	NET			HLIS	(Primary 8	Secondary)		rculosis	
Reporting Area	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	
UNITED STATES	1,175	2,043	561	1,181	584	910	688	1,294	
NEW ENGLAND Maine N.H. Vt. Mass.	20 - - 16	55 2 1 1 42	15 - - 9	43 - 1 - 29	6 - - 4	9 - - 7	40 - 1 - 25	37 1 1 - 21	
R.I. Conn.	- 4	3	6	6 7	2	1	3 11	2 12	
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	93 56 29 - 8	120 19 49 37 15	65 2 39 8 16	123 21 47 26 29	32 32 30 6 3	37 1 20 7 9	159 20 56 57 26	174 12 108 47 7	
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	216 70 37 53 52 4	358 18 22 146 140 32	104 20 5 48 29 2	123 8 9 2 101 3	60 5 17 11 25 2	182 12 64 71 23 12	98 17 10 57 - 14	113 19 3 83 3 5	
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	180 66 25 48 8 2 9 22	102 21 15 52 1 8 5	127 85 34 1 - 7	85 40 16 20 - 6 3	5 4 - - - - -	20 3 5 10 - 1 1	39 25 - 1 5 -	48 23 3 17 - 2 1 2	
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	176 2 17 8 12 2 51 12 7 65	158 - - 10 1 12 3 6 110	53 - 3 U 5 5 19 7 13 13 1	70 1 5 U 12 1 5 1 24 21	228 1 31 4 15 - 63 31 21 62	277 1 53 11 20 1 78 23 38 52	112 11 9 13 5 10 8 48 8	202 16 5 5 17 18 44 97	
E.S. CENTRAL Ky. Tenn. Ala. Miss.	105 47 13 26 19	96 19 43 5 29	31 13 15 - 3	70 13 51 4 2	87 7 43 21 16	126 7 88 18 13	50 3 36 11	84 5 21 39 19	
W.S. CENTRAL Ark. La. Okla. Tex.	66 31 8 1 26	356 33 50 5 268	97 10 25 - 62	373 3 20 4 346	98 10 18 12 58	146 9 36 37 64	19 15 - 4	258 8 6 7 237	
MOUNTAIN Mont. Idaho Wyo. Colo.	107 - 4 - 20	179 21 1 31	52 - - 12	66 - 15 1 13	26 - - 1	28 - - 1	21 - - 9	58 - - -	
N. Mex. Ariz. Utah Nev.	20 23 52 3 5	18 65 5 38	7 28 5	13 13 19 5	1 19 4 1	2 23 2	1 10 1	8 7 15 4 24	
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	212 25 17 170	619 126 72 412 2 7	17 - 15 - 2	228 178 44 - 1 5	42 13 25 - 25	85 8 1 76 -	150 25 119 6	320 24 1 281 3 11	
Guam P.R. V.I. Amer. Samoa C.N.M.I. N: Not notifiable.	- - - - - - - - - - - - - - - - - - -	- 7 U U U Vailable.	U U U U -: No repo		32 U U U U	29 U U U	- U U U	- 17 U U U	

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending February 24, 2001, and February 26, 2000 (8th Week)

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).

			1		-			Measles (Rubeola)							
		<i>ienzae,</i> isive		epatitis (V	iral), By Typ B	be	Indige	10115	Impo		ola) Tota				
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.		Cum.		Cum.	Cum.	Cum.			
Reporting Area	2001 ⁺ 176	2000 190	2001 980	2000 1,948	2001 552	2000 827	2001	<u>2001</u> 8	2001 2	2001 5	2001 13	2000 10			
NEW ENGLAND	8	18	50	46	9	15	_	3	-	1	4	-			
Maine	-	-	1	1	1	1	-	-	-	-	-	-			
N.H. Vt.	-	2 2	3 1	6 1	2 1	3 2	-	- 1	-	-	- 1	-			
Mass. R.I.	8	14	11 3	17	1 4	1	-	2	-	1	3	-			
Conn.	-	-	31	21	-	8	-	-	-	-	-	-			
MID. ATLANTIC	17	28	41	110	44	133	-	-	-	-	-	3			
Upstate N.Y. N.Y. City	6 6	12 9	16 22	32 60	9 27	7 75	-	-	-	-	-	- 3			
N.J.	4	5	-	5	-	7	-	-	-	-	-	-			
Pa.	1	2	3	13	8	44	-	-	-	-	-	-			
E.N. CENTRAL Ohio	22 16	27 9	135 40	298 61	84 17	87 17	-	-	2	2	2	3 2			
Ind. III.	5	2 13	4 24	5 128	2 2	1 2	-	-	- 2	- 2	- 2	-			
Mich.	1	3	67	92	ຜ້	66	-	-	-	-	-	1			
Wis.	-	-	-	12	-	1	-	-	-	-	-	-			
W.N. CENTRAL Minn.	2	4	71 1	178 18	34 1	53	-	1	-	-	1	-			
lowa	-	-	6	17	3	9	-	-	-	-	-	-			
Mo. N. Dak.	2	3 1	17 -	116	24	37	-	-	-	-	-	-			
S. Dak. Nebr.	-	-	- 17	- 4	1 4	- 4	-	-	-	-	-	-			
Kans.	-	-	30	23	1	3	-	1	-	-	1	-			
S. ATLANTIC	67	42	135	161	87	123	-	2	-	1	3	-			
Del. Md.	- 15	20	- 47	- 25	16	- 25	-	2	-	- 1	- 3	-			
D.C.	5	10	3 20	28	2	21	-	-	-	-	-	-			
Va. W. Va.	3	1	-	19	11 1	-	-	-	-	-	-	-			
N.C. S.C.	14 1	3 1	10 9	45 3	29	55 1	-	-	-	-	-	-			
Ga.	10	6	1	14	1	2	-	-	-	-	-	-			
Fla.	19	1	45	27	27	19 05	-	-	-	-	-	-			
E.S. CENTRAL Ky.	9	10 7	40 6	84 4	54 3	65 8	-	-	-	-	-	-			
Ténn. Ala.	5 4	3	20 14	23 15	23 20	30 5	-	-	-	-	-	-			
Miss.	-	-	-	42	8	22	Ū	-	Ū	-	-	-			
W.S. CENTRAL	2	15	127	381	30	86	-	-	-	-	-	-			
Ark. La.	-	- 6	16 10	27 18	14 4	10 29	-	-	-	-	-	-			
Okla. Tex.	2	9	26 75	55 281	11 1	8 39	-	-	-	-	-	-			
MOUNTAIN	- 40	- 25	75 152	123	67	39 63	-	-		- 1	- 1	-			
Mont.	-	-	2	1	-	2	-	-	-	-	-	-			
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TABLE III. Provisional cases of selected notifiable diseases preventable
by vaccination, United States, weeks ending February 24, 2001,
and February 26, 2000 (8th Week)

N: Not notifiable. U: Unavailable. - : No reported cases. *For imported measles, cases include only those resulting from importation from other countries. † Of 32 cases among children aged <5 years, serotype was reported for 10 and of those, 1 was type b.

Importing Area 2001 2001 2000			jococcal ease		Mumps	-		Pertussis			Rubella	
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N: Not notifiable: No reported cases.		-				-	U	U	U	U	U	U

TABLE III. (Cont'd) Provisional cases of selected notifiable diseases preventable
by vaccination, United States, weeks ending February 24, 2001,
and February 26, 2000 (8th Week)

		All Cau	ises, By	Age (Ye	ears)		P&I [†]			All Cau	ises, 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. Springfield, Mass Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Erie, Pa.§	. 14 33 38 34 14 ss. 32 . 38 U 6 . 41	414 119 40 12 28 29 12 23 21 U 5 29 27 43 1,650 15 15 32 91 15 33	33 5 2 4 8 4 2 8 9 U - 9 4	32 11 3 - 1 4 1 - 1 - 1 - 2 4 167 6 - 7 - 7 - 3 2	10 3 - - 2 U 2 2 2 1 35 - 4 4 -	9 3 - - 2 U 1 1 2 34 - 2 34 -	58235-142-36001346 1454-15-1	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, I Tampa, Fla. Washington, D. Wilmington, De E.S. CENTRAL Birmingham, Al Chattanooga, Te Knoxville, Tenn. Lexington, Ky. Memphis, Tenn Mobile, Ala. Montgomery, A Nashville, Tenn.	89 71 61 61a. 58 210 C. 101 1. 25 880 a. 156 900 . 80 71 53 . 205 102	874 111 118 86 91 63 47 37 36 48 139 73 36 48 139 73 578 104 578 104 54 47 333 136 66 83 8 100	243 44 26 30 24 14 13 17 6 4 49 16 - 182 30 18 18 19 45 21 44 34	109 15 23 11 10 11 6 4 2 37 79 10 6 11 7 20 8 9 8	28 6 2 2 3 1 3 2 2 2 2 3 - 16 3 1 - 1 2 3 2 4	19 32 2 3 - 2 1 1 - 1 3 2 2 4 8 1 2 2 4 1 2 3 2 2 4 1 3 3 2 2 4 1 3 3 2 2 3 5 - 2 2 3 5 - 2 2 3 5 - 2 2 2 3 5 2 2 2 3 5 2 2 2 3 5 2 2 2 2 3 5 2 2 2 2	98 4 20 18 12 1 6 3 4 17 1 - 70 15 6 6 2 12 5 8 16
Jersey City, N.J. Newark, N.J. Paterson, N.J. Philadelphia, Pa. Pittsburgh, Pa.§ Reading, Pa. Rochester, N.Y. Schenectady, N.Y. Scranton, Pa.§ Syracuse, N.Y. Trenton, N.J. Utica, N.Y. Yonkers, N.Y.	73 24 294 87 25 136	41 843 38 11 191 60 20 107 18 21 54 24 28 U	15 11 69 24 3 19 5 2 8	2 92 15 1 23 2 1 8 - 1 4 - U	17 1 6 - 1 - 1 - U	- 14 3 - 5 1 1 1 - 1 2 1 - U	- 86 5 1 20 5 4 9 1 2 7 3 3 U	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.	Tex. 66 265 75 118 360 64 . U	1,063 61 53 48 177 49 82 205 43 U 176 94 75	318 17 22 13 40 16 18 82 14 U 50 20 26	129 12 7 4 27 7 44 2 U 16 2 1	56 1 3 1 0 2 3 24 3 U 4 3 2	41 2 11 1 5 2 U 5 3 2	110 7 3 1 20 5 5 31 2 U 16 12 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, Mid. 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, Kans St. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	182 40 162 55 64 111 0 39 673 83 28 28 28 28 28 28 0 0 1 0 0 1 0 0 1 0 0 0 0 0 0 0 0 0 0	1,257 47 30 88 1592 115 43 118 46 57 133 118 46 57 48 57 48 57 48 57 48 57 48 57 48 57 48 57 48 57 54 30 48 57 54 50 56 56 56 56 56 56 56 56 56 56 56 56 56	19 24 355 70 8 13 2 2 30 6 28 6 11 22 6 12 19 3 8 U 8 20 17 2 15	116 3 - U 4 8 14 6 25 3 1 - 3 14 1 1 3 4 6 9 1 8 3 1 1 U 3 3 6 1 1 6 4	35 1 U 1 4 10 1 1 - 2 1 1 - 1 - 1 - 9 1 - U 2 1 - 2 1 2 1 2	41 2 2 U 6 2 5 2 3 2 1 - 1 2 - 4 3 2 1 2 1 2 3 U - 5 7 2 3 2	11964U04201624·5114155146· 6864U6135685	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. Fresno, Calif. Glendale, Calif. Honolulu, Hawa Long Beach, Cal Los Angeles, Ca Pasadena, Calif. Portland, Oreg. Sacramento, Ca San Diego, Calif San Francisco, C San Jose, Calif. Santa Cruz, Cali Seattle, Wash. Tocma.	40 solo. 52 121 206 30 159 42 tah 120 1,529 25 141 11 ii 74 if. 100 lif. 349 24 U lif. 172 172 talif. U 161 f. 27 113	$\begin{array}{c} 724\\ 76\\ 32\\ 34\\ 76\\ 147\\ 79\\ 118\\ 1,130\\ 18\\ 112\\ 79\\ 118\\ 1,130\\ 18\\ 102\\ 73\\ 246\\ 16\\ 0\\ 133\\ 133\\ 133\\ 133\\ 133\\ 199\\ 75\\ 46\\ 74\\ 8,172 \end{array}$	192 21 5 7 27 35 4 24 7 29 33 27 6 4 23 2 10 21 8 3 U 26 3 U 26 3 U 26 3 U 26 3 U 26 3 U 27 5 4 4 2 7 27 5 4 4 7 27 5 4 4 7 27 5 4 27 5 27 5	82 10 1 9 11 14 2 16 4 7 8 69 3 5 2 2 3 3 3 U 10 2 U 7 2 4 - 3 821	22 2 1 - 3 5 - 6 - 1 4 3 5 - 6 - 1 4 3 5 - 6 - 1 4 3 5 - 6 - 1 4 3 5 - 2 2 9 2 U 2 6 U 2 6 U 4 - 4 4 2 9 2 U 2 6 0 - 4 0 2 6 0 2 8 5 2 9 2 9 2 9 2 9 2 9 2 9 2 9 2 9 2 9 2	25 - 1 2 4 5 2 4 - - - - - - - - - - - - -	82 8 4 3 9 10 - 17 5 15 11 16 2 10 - 5 16 27 4 U 12 5 U 15 2 15 5 13 9 0 - 17 5 15 11 16 2 10 - 5 16 27 4 U 22 5 15 2 15 15 15 15 11 16 2 10 - 5 16 27 4 U 22 5 15 2 15 2 15 15 13 10 2 10 - 5 16 27 4 U 22 5 15 2 15 2 15 15 13 10 2 10 - 5 16 27 4 U 22 5 15 2 15 2 15 13 2 15 15 13 10 2 10 - 5 13 10 2 10 - 5 13 13 10 10 10 15 15 13 13 10 10 15 15 13 13 10 10 15 15 13 13 10 10 15 15 13 13 10 10 15 15 13 10 10 10 15 15 13 10 15 13 10 15 15 13 10 10 15 15 13 10 10 15 15 13 10 15 15 13 10 10 15 15 13 10 15 15 13 10 15 15 15 13 10 15 15 15 13 10 10 15 15 15 13 10 10 15 15 15 15 15 15 15 15 15 15

TABLE IV. Deaths in 122 U.S. cities,* week endingFebruary 24, 2001 (8th Week)

U: Unavailable. -: No reported cases. *Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of ≥100,000. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included. *Pneumonia and influenza.

[®]Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks. [®]Total includes unknown ages.

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