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Update: Isolation of Avian Influenza A(H5N1) Viruses from Humans — Hong Kong, 1997–1998

MORBIDITY AND MORTALITY WEEKLY REPORT

As of January 6, 1998, a total of 16 confirmed and three suspected cases of human infection with avian influenza A(H5N1) viruses have been identified in Hong Kong. Confirmed cases are those from which an influenza A(H5N1) virus was isolated or in which a seroconversion to influenza A(H5N1) virus was detected by a neutralization assay. Suspected cases are those with influenza-like illness (ILI) and preliminary laboratory evidence of influenza A(H5N1) infection. This report summarizes interim findings from the ongoing epidemiologic and laboratory investigation of influenza A(H5N1) cases by health officials in Hong Kong and by CDC.

The first known case of human infection with influenza A(H5N1) occurred in a 3year-old boy who died from respiratory failure in May 1997 (1). Of the 15 remaining confirmed cases, five persons had onset of illness in November and 10 in December; all three persons with suspected cases had onset during December. No cases have been identified with onset after December 28, 1997. Ages of persons with confirmed cases ranged from 1 to 60 years (mean age: 17 years) and, for persons with suspected cases, from 3 to 7 years (mean age: 5 years). Nine (47%) cases occurred among persons aged \leq 5 years. Four persons with confirmed cases have died, and three remain in critical condition.

Testing has been completed of serum samples collected in August as a part of the epidemiologic investigation of the first case of human influenza A(H5N1) infection. Serum samples were obtained from 502 persons who may have had contact with the child or with poultry, including family members, persons who lived in the same neighborhood, children and staff of the child-care center the child attended, health-care workers, poultry workers, and persons working on pig farms. Samples of control serum specimens were obtained from 218 healthy children and 201 healthy adult residents of Hong Kong. These samples were tested for antibody to influenza A(H5N1) virus using a micro-neutralization assay. Of the 502 persons tested who may have had contact with the child or with poultry, elevated neutralization antibody titers to influenza A(H5N1) virus were present in nine (2%). These persons included five (17%) of 29 poultry workers, one (2%) of 54 health-care workers, one (2%) of 63 neighbors, one (1%) of 73 laboratory workers, and one (0.4%) of 261 child-care center contacts. Specimens were negative for the four family members, 18 persons working on pig farms, and the 419 controls. Seropositivity was not associated with reported ILI.

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES

Avian Influenza A Virus — Continued

Antigenic and genetic analyses of viral isolates from seven case-patients indicated two closely related but distinguishable groups of influenza A(H5N1) viruses, suggesting multiple introductions in humans from poultry sources. All seven of the influenza A(H5N1) viruses analyzed from human cases contained all eight RNA gene segments from avian viruses, indicating that genetic reassortment between avian and human influenza viruses has not occurred.

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Editorial Note: The cases reported in Hong Kong represent the first identified instances of human illness associated with infection with influenza A(H5N1) viruses. Goals of the ongoing investigation are to detect new cases, determine sources of infection and mode(s) of transmission, and identify risk factors for influenza A(H5N1) infection. Except for a cluster of two confirmed and two suspected cases in one family, case-patients are not known to have had contact with each other or a common source of exposure and are geographically distributed throughout Hong Kong. All cases of infection have occurred among residents of Hong Kong, and no cases of infection with influenza A(H5N1) viruses have been identified among persons residing outside Hong Kong.

The serologic data obtained as part of the epidemiologic study of the initial case support the preliminary conclusion that persons with high levels of exposure to infected poultry or direct exposure to the virus in the laboratory may be at increased risk for infection with influenza A(H5N1) virus. However, the investigation has not ruled out the possibility of person-to-person transmission from exposure to ill and infectious persons: two seropositive persons who had contact with the first case-patient included a child-care center classmate and a health-care worker, and the classmate had contact with both the ill child and the same potential environmental source of exposure to ill chickens at the school as the ill child. However, the health-care worker reported no history of exposure to the virus in the laboratory or any recent exposure to poultry, and a history of exposure to the child or to poultry was unknown for a seropositive elderly neighbor. On the basis of the overall low rates of infection among contacts and controls and the lack of seropositivity among family members, at this time, the virus probably is not being efficiently transmitted among humans.

Global surveillance for influenza viruses is critical to monitor the circulation of different strains and indicates that human influenza type A(H3N2), type A(H1N1), and type B viruses continue to circulate worldwide. Data from the Hong Kong Department of Health's influenza surveillance system indicate that the number of cases of ILI in Hong Kong is at normal levels for this period; however, during December, the number of human influenza viruses isolated increased. During December, influenza A(H3N2) was the most commonly isolated influenza strain in Hong Kong, although influenza A(H1N1) and B viruses also were identified. The currently available inactivated trivalent influenza vaccine contains influenza A(H1N1), A(H3N2), and B strains repre-

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sentative of those currently circulating among humans and is recommended for persons at increased risk for influenza-related complications (2).

Information about influenza A(H5N1) activity in Hong Kong and the United States and international influenza surveillance data are available through CDC's Influenza Branch, Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases, World-Wide Web site http://www.cdc.gov/ncidod/diseases/flu/fluvirus.htm.

References

- CDC. Isolation of avian influenza A(H5N1) viruses from humans—Hong Kong, May–December 1997. MMWR 1997;46:1204–7.
- CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 1997;46(no. RR-9).

Rubella Among Crew Members of Commercial Cruise Ships — Florida, 1997

During April–July 1997, two different commercial cruse lines notified CDC of rubella outbreaks among crew members. In July 1997, CDC initiated an investigation on one cruise ship to determine the extent of and risk factors for rubella infection among crew members and to assess the potential risk for rubella transmission to passengers—particularly rubella-susceptible pregnant women at risk for giving birth to an infant with congenital rubella syndrome (CRS). This report summarizes rubella outbreaks involving two cruise ships and the results of the CDC investigation on one cruise ship, which demonstrate that crew members can serve as a susceptible population for rubella infection and should be vaccinated with measles-mumps-rubella vaccine (MMR) if they are not immune. Although the outbreaks were limited to crew members, cruise ship travel provides an environment conducive to the potential spread of rubella and other infectious diseases among crew and passengers; therefore, women of childbearing age, particularly pregnant women, should be immune to rubella before traveling on cruise ships to reduce the risks for rubella infection and CRS.

Cruise Ship A

On April 7, cruise line A notified CDC about a rash illness in a crew member aboard one of the ships in its fleet. The cruise ship sailed twice a week from Florida on 3-day cruises to the Bahamas, carrying approximately 900 crew members and 2000 passengers per cruise. During May and June, rash illnesses were reported in six additional crew members; five of the seven cases were confirmed serologically (by immunoglobulin [Ig] M antibodies) as acute rubella infection. A survey of the crew members conducted by the cruise line indicated that a substantial proportion had no documentation of rubella vaccination and that at least 95% were not U.S.-born. Because of evidence of ongoing transmission of rubella among crew members (many of whom were natives of countries without rubella vaccination programs) and the potential for transmission to female crew members and passengers of childbearing age, CDC advised the cruise line to initiate a vaccination campaign with MMR during June. Serologic susceptibility testing was recommended for all crew members ineligible for vaccination, including pregnant women. Cruise line staff and state and local health depart-

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ment personnel vaccinated 865 (96%) of the approximately 900 crew members who had no documented rubella vaccination or immunity. Following the vaccination campaign, one additional rash illness was reported in a crew member and subsequently was serologically confirmed to be consistent with acute rubella infection. This crew member had received MMR <2 weeks before the onset of rash.

Cruise Ship B

On July 25, cruise line B notified CDC about a cluster of rash illnesses among crew members of one of its cruise ships sailing between Florida and the Bahamas. The cruise ship sailed daily from Florida with a crew of 385 and carried approximately 8400 passengers per week. CDC initiated an investigation in July to determine the extent of the outbreak and risk factors for rubella infection among crew members and to assess the potential risk for rubella transmission to passengers, particularly rubella-susceptible pregnant women at risk for serious adverse health outcomes (including CRS).

The investigation included review of the ship's medical logs and interviews and examinations of the 385 crew members. Because approximately 25%-50% of rubella infections are asymptomatic (1), a serosurvey of rubella IgM and IgG antibodies was conducted among 366 consenting crew members. A confirmed case was defined as IgM serology consistent with rubella infection, or signs and symptoms meeting the clinical case definition for rubella and linked epidemiologically to a laboratoryconfirmed case with onset during May 30-August 2. Rubella was confirmed in 16 (4%) crew members; all confirmed cases had IgM serology consistent with rubella infection. Of 16 crew members with IgM-confirmed cases, eight (50%) had no symptoms of infection. An additional 25 (7%) of the 366 crew members surveyed were susceptible to rubella at the time of the serosurvey. The crew interviews indicated that approximately 85% of the crew members were not U.S.-born (representing at least 50 countries), and 75% had negative or unknown rubella vaccination histories. Crew members living aboard the ship were more likely to have confirmed rubella than were crew members living ashore (16 of 288 versus zero of 78; relative risk=9.0 [Woolf's estimate], p=0.03).

To determine demographic characteristics of passengers on cruise ship B and identify pregnant women who, if susceptible to rubella, could be at risk for giving birth to infants with CRS, a questionnaire was administered to passengers sailing on cruises during August 4–8. All passengers (approximately 6000) received a health alert about the rubella outbreak before boarding the ship; 3643 (61%) passengers completed the questionnaire. Among the respondents, approximately 75% of passengers were U.S.born, 12% were born in the Bahamas, and 13% were born in other countries. A total of 1213 (33%) of the 3643 respondents were women of childbearing age (i.e., 15–44 years); 28 (0.8%) of all respondents were pregnant women, of whom 14 (50%) reported being in the first trimester. Although the rubella immune status of these pregnant passengers was not determined, previous serosurveys in the U.S. population suggest that approximately 10% of women of childbearing age may be susceptible to rubella, and up to 85% of susceptible pregnant women who are infected during their first trimester may give birth to an infant with CRS (*2*).

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Editorial Note: Although rubella is typically a mild, self-limited disease in adults, infection in pregnant women can result in serious adverse health outcomes for the fetus, including CRS, a group of birth defects including deafness, cataracts, heart defects, and mental retardation. In the United States, approximately 10% of young adults are susceptible to rubella; in other countries, some without routine vaccination policies for rubella, susceptibility rates for rubella range from 4% to 68% (*3*). During 1994–1996, 12 laboratory-confirmed cases of CRS were reported in the United States (*4*).

Although a definitive quantitation of the risk for transmission of rubella among crew members and passengers on the cruise ships could not be ascertained, risk for infection among those crew members of cruise ship B could be estimated. Results of the serosurvey among crew members indicate that at least 41 (11%) of 366 were acutely infected with or susceptible to rubella at the time of the serosurvey. This serosurvey was conducted after recognition of an ongoing outbreak of rash illnesses among crew members, and it is likely that rubella susceptibility rates at the outset of the outbreak would have been higher.

The risk for transmission of infection and an outcome of CRS in pregnant passengers in their first trimester of pregnancy on cruise ship B was difficult to determine because 1) the rubella immune status of these pregnant passengers was unknown and 2) the consequences of rubella infection in susceptible pregnant women (i.e., CRS) may not be evident for several months after the exposure. If pregnant passengers were exposed, and assuming that approximately 10% of these women were susceptible to rubella and 85% of susceptible pregnant women who are infected during their first trimester will give birth to an infant with CRS, one case of CRS could potentially occur each week among passengers sailing during the outbreak.

Minimizing or eliminating the risk for rubella exposure among susceptible pregnant women is important because of the potential for serious adverse health outcomes for the fetus. To interrupt transmission of rubella among crew members and to prevent transmission of infection and CRS among susceptible pregnant women, CDC recommended administration of MMR to all crew members lacking documented immunity to rubella; serologic testing to determine susceptibility to rubella for all crew members ineligible for vaccination, including pregnant women; active surveillance aboard the ship to detect new rubella infections; prospective notification of the potential risk for rubella exposure to all embarking passengers until 30 days after the last confirmed rubella infection; and retrospective notification to all passengers sailing during the period of potential rubella transmission. These recommendations were effective in interrupting rubella transmission among crew members on cruise ship B: no additional rash illnesses were identified after their implementation.

This report of two clusters of rubella infections on commercial cruise ships demonstrates that crew members—many from countries without routine rubella vaccination programs—are potential groups of susceptible persons at risk for rubella infection. To prevent future rubella outbreaks among such persons, CDC recommends that cruise lines administer MMR to all crew members without documented immunity to rubella. Although reported rubella cases in these two outbreaks were limited to crew mem-

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bers, cruise ship travel provides a semi-closed environment for crew and passenger interactions, conducive to the potential spread of rubella and many other infectious diseases among crew and passengers. To prevent transmission of rubella infection and subsequent CRS, women of childbearing age, particularly pregnant women, should be immune to rubella before cruise ship excursions or international travel.

The outbreaks described in this report illustrate the potential for transmission of infectious disease among persons traveling across international borders, including aboard commercial cruise ships. Previous infectious disease outbreaks reported among crew members and passengers have included diarrheal diseases and other vaccine-preventable diseases such as influenza (*5*). Approximately 4 million persons travel aboard North American cruise ships each year (CDC, unpublished data, 1998). Ensuring routinely recommended adult vaccinations for all crew members will substantially decrease the potential for future outbreaks of vaccine-preventable illnesses aboard cruise ships. All suspected cases of rubella and other notifiable vaccine-preventable diseases should be reported to the nearest state and local health department. State health departments should report all suspected cases of rubella to CDC's Child Vaccine-Preventable Disease Branch, Epidemiology and Surveillance Division, National Immunization Program, telephone (404) 639-8230.

References

- 1. American Academy of Pediatrics. 1997 Red book: report of the Committee on Infectious Diseases. 24th ed. Elk Grove Village, Illinois: American Academy of Pediatrics, 1997.
- 2. Miller E, Cradock-Watson JE, Pollock TM. Consequences of confirmed maternal rubella at successive stages of pregnancy. Lancet 1982;2:781–4.
- Cutts FT, Robertson SE, Diaz-Ortega JL, Samuel R. Control of rubella and congenital rubella syndrome (CRS) in developing countries, part 1: burden of disease from CRS. Bull World Health Organ 1997;75:55–68.
- 4. CDC. Rubella and congenital rubella syndrome—United States, 1994–1997. MMWR 1997; 46:350–4.
- 5. CDC. Update: influenza activity—United States, 1997–98 season. MMWR 1997;46:1094–98.

Enhanced Medical Assessment Strategy for Barawan Somali Refugees — Kenya, 1997

Each year, approximately 100,000 refugees are resettled to the United States. Before resettlement, these refugees undergo medical screening to identify inadmissible conditions (e.g., infectious tuberculosis and human immunodeficiency virus [HIV] infection) among individual refugees. This report describes the implementation and results of an enhanced refugee medical assessment strategy among Barawan Somali refugees in Kenya during July 1997. This strategy employs population-based screening for parasitic infections. The findings indicate that, among these refugees, the prevalences of malaria and intestinal parasites were sufficient to warrant preembarkation therapy to improve the health of both individuals and the total refugee population. This therapy also may prevent local transmission of parasitic infections in the resettlement communities in the United States.

In May 1997, resettlement began for approximately 4000 Barawan Somali refugees encamped since 1992 near Mombasa, Kenya. In 1993, detection of substantial malaria parasitemia (15%) among Somali refugees from this region prompted recommenda-

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tion of antimalarial treatment before resettlement (1). In addition, high prevalences of malaria (30%) and intestinal parasites (60%-80%) had been reported among residents of Kenya living in the coastal region, including Mombasa (S.K. Sharif, M.D., Ministry of Health, Kenya, personal communication, 1997). Because the prevalence of parasitic infections among the Barawan refugees may reflect those of the local community, the International Organization for Migration (IOM) consulted CDC on appropriate preembarkation interventions for Barawan refugees. IOM, a nongovernmental organization, medically screens more than half of the refugees resettling to the United States. CDC interim recommendations included mass pre-embarkation therapy with singledose sulfadoxine-pyrimethamine (SP) for malaria parasitemia and mebendazole (100 mg twice a day for 3 days) for intestinal helminths. During July 1997, CDC conducted a cross-sectional survey of an approximately 10% sample of refugees during the standard medical screening process to 1) determine the prevalences of malaria and intestinal parasites, 2) reevaluate recommended pre-embarkation therapies, 3) assess the effectiveness of the antimalarial regimen, and 4) evaluate the laboratory component of medical screening.

IOM provided information about two groups: refugees examined during February 3–June 23 (travel-approved population, n=3253) and refugees examined during July 7–17 (survey population, n=390). Basic characteristics of the two groups (i.e., age, sex, country of origin, and size of family unit) were similar. Members of the survey population were asked about histories of recent illness and use of medications and other antimalarial preventive measures. A local hospital laboratory screened members of the survey population for malaria by using a qualitative buffy-coat (QBC) test followed by confirmation of all QBC-positives using microscopic examination of Field's-stained blood smears; persons who were positive for malaria were retested 3 and 7 days following completion of antimalarial therapy. Stool specimens were screened at a local hospital for intestinal parasites by direct and formalin ether-concentrated smears. CDC performed quality-control assessments for both the malaria smears and stool samples.

Malaria

Of the 390 survey participants, 26 (7%; 95% confidence interval [CI]=4%–10%) were positive for malaria. Of the 26 who were positive, 25 had *Plasmodium falciparum* parasitemia, and one had *P. ovale* parasitemia. Because of the severity of the parasitemia and symptoms, the local hospital treated seven of the 26 malaria-positive persons with halofantrine or artemether. Nineteen received a weight-adjusted dose of SP. One patient receiving SP was lost to follow-up. Of the remaining 18 patients receiving SP, 13 were malaria-negative on day 3 of follow-up, and all were malaria-negative by day 7.

Of the surveyed population, recent febrile symptoms were reported by 20% and 37% during initial and follow-up questioning, respectively (Table 1). Use of antimalarial therapy (chloroquine, halofantrine, SP, or quinine) was common among those refugees reporting fever (71% and 93%, respectively). Ten percent of the surveyed refugees reported using any malaria chemoprophylaxis, and most (91%) reported using bed nets (Table 1). Of the 229 refugees reporting the condition of their bed nets, 51 (22%) reported holes or tears in the netting (i.e., poor condition). Use of

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bed nets in poor condition compared with use of bed nets in good condition was associated with malaria parasitemia (odds ratio=9.2; 95% CI=3.2–27.5).

A total of 37 randomly selected blood smears from refugees reported as parasitenegative by the local hospital were reviewed by CDC and confirmed as negative. However, of the 26 refugees reported as parasite-positive, two cases of *P. falciparum* parasitemia could not be confirmed by CDC. The smear diagnosed by the local hospital as *P. ovale* was identified by CDC as *P. falciparum*.

Intestinal Parasites

Stool specimens were obtained from 331 persons; of these, specimens from 129 (39%) were positive for one or more pathogenic intestinal parasites, including *Trichuris trichiura* (28%), *Ascaris lumbricoides* (9%), and other pathogens (Table 2). Sex-specific prevalences were similar (41% for females versus 37% for males, chi-square test=0.47, p=0.49). However, age-specific prevalence was higher for persons

		native onses	Total
Survey/Characteristic	No.	(%)	respondents
nitial survey*			
Had fever in July	77	(20)	389
Received any antimalarial therapy [†]	54	(71)	76
Follow-up survey [§]			
Had fever during May–July	131	(37)	358
Received any antimalarial therapy [†]	122	(93)	131
Used chemoprophylaxis	34	(10)	355
Used bed nets	325	(91)	355
Bed nets in poor condition	51	(22)	229

 TABLE 1. Number of affirmative responses to initial and follow-up questions about

 malaria symptoms among Barawan Somali refugees, 1997

*Survey conducted during July 7–17; n=390.

[†]Chloroquine, sulfadoxine-pyrimethamine, quinine, halofantrine, or artemether.

[§]Survey conducted July 12 and July 16; n=358.

TABLE 2. Number and percentage distribution of selected intestinal parasites among
Barawan Somali refugees and comparison of results of stool specimen screening
performed by a local hospital and reviewed by CDC, by parasite, 1997

				Quality	review	
	Survey p	opulation*	Local h	nospital	C	DC [†]
Parasite	No.	(%)	No.	(%)	No.	(%)
Trichuris trichiura	92	(28)	13	(32)	17	(42)
Ascaris lumbricoides	29	(9)	7	(17)	7	(17)
Giardia lamblia	25	(8)	5	(12)	7	(17)
Entamoeba histolytica	7	(2)	2	(5)	4	(10)
Hookworms	3	(1)	1	(2)	0	—
Strongyloides stercoralis	3	(1)	1	(2)	1	(2)
Hymenolepis nana	3	(1)	0	—	0	—

*Stool specimens from survey population tested at a local hospital during July; n=331.

[†]Quality review by CDC during October; n=41.

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aged <15 years (51%) than for persons aged \geq 15 years (32%) (chi-square test=11.95, p<0.01). CDC reviewed randomly selected negative (n=15) and positive (n=26) stool specimens as determined by the local hospital and found that, for 11 (27%) of these 41 specimens, the local hospital either did not detect or misclassified pathogens that were present in sufficient numbers to detect. The most commonly undetected pathogen was *T. trichiura*, and the most commonly misclassified pathogen was *Entamoeba histolytica*.

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Editorial Note: Although the prevalences of parasitic infections among the Barawan refugees were lower than the prevalences of these infections among persons in the surrounding communities, the prevalences of malaria (7%) and intestinal parasites (39%) among Barawan Somali refugees encamped in Kenya were sufficient to warrant pre-embarkation therapies. The strategy of screening for parasitic infections among a subset of refugees before resettlement provided an opportunity to assess the need for public health interventions for the entire Barawan refugee population. This strategy optimized the efficient distribution of these therapies before the refugees were resettled to the United States. This screening strategy also may be used to determine the need for other pre-embarkation therapies among future refugee populations. However, because the magnitudes of exposures and risks may vary among different groups, the use of specific interventions may differ by refugee group.

CDC oversees refugee health screening in accordance with the Refugee Act of 1980.* The law requires that refugees with medical conditions potentially affecting the public's health be identified and treated; the quality of medical screening and related health services be monitored and assessed; and that health officials in resettlement communities be notified of identified medical conditions. Refugee medical assessments previously focused on identifying inadmissible medical conditions. The enhancement of the medical screening process described in this report emphasizes the expansion of screening to include parasitic diseases with the potential for local transmission in the resettlement community (*2,3*) and a broadening of the focus from the individual to a population.

As a result of the findings of the enhanced assessment of Barawan Somali refugees, CDC recommended continuation of mass pre-embarkation therapy (day before departure) for malaria infection with SP for all departing refugees who had no contraindication to therapy (i.e., sulfa allergy). This recommendation was based on three considerations. First, the prevalence of parasitemia (7%) may have been underestimated because of the extensive use of presumptive antimalarial therapy for fever. Second, single-dose SP provides adequate cost-effective therapy for *P. falciparum*. Although the small number of refugees treated with SP (n=19) precluded accurate assessment of the effectiveness of SP, all refugees were malaria-negative by day 7 following SP therapy (n=18, one lost to follow-up). Third, mass pre-embarkation therapy effectively treats symptomatic persons and reduces asymptomatic malaria parasitemia among the entire refugee population, thereby reducing the risk for imported *P. falciparum* malaria.

^{*}Public Law 96-212.

Barawan Somali Refugees — Continued

Because some Barawan Somali refugees were infected with both helminthic and protozoan pathogens, the interim recommendation for mass pre-embarkation therapy with 3-day mebendazole was changed to single-dose albendazole (400 mg per kg of body weight) for all persons except pregnant women.[†] This approach was considered preferable because of the high prevalence of mixed intestinal parasites, the low cost of albendazole, and the ease of single-dose therapy before departure (4–6). The optimal choices of agent(s) and duration of therapy for mass treatment of intestinal parasites among refugee populations remain to be determined.

The program of enhanced screening for and management of infectious diseases among this vulnerable refugee population enabled the implementation of populationbased interventions before members of this group dispersed to multiple locations in ≥20 states. CDC is notifying health officials in the states in which refugees are being resettled of the results of the pre-embarkation medical screening and treatment. CDC also is working with IOM and state refugee health programs to develop a shared database of refugee medical screening results.

References

- Slutsker L, Tipple M, Keane V, McCance C, Campbell CC. Malaria in East African refugees resettling to the United States: development of strategies to reduce the risk of imported malaria. J Infect Dis 1995;171:489–93.
- 2. CDC. Malaria in Montagnard refugees—North Carolina, 1992. MMWR 1993;42:180-3.
- Gyorkos TW, Frappier-Davignon L, MacLean JD, Viens P. Effect of screening and treatment on imported intestinal parasite infections: results from a randomized, controlled trial. Am J Epidemiol 1989;129:753–61.
- 4. Raccurt CP, Lambert MT, Bouloumie J, Ripert C. Evaluation of the treatment of intestinal helminthiases with albendazole in Djohong (North Cameroon). Trop Med Parasitol 1990;41:46–8.
- 5. Pungpak S, Singhasivanon V, Bunnag D. Albendazole as a treatment for *Giardia* infection. Ann Trop Med Parasitol 1996;90:563–5.
- 6. Hall A, Nahar Q. Albendazole as a treatment for infections with *Giardia duodenalis* in children in Bangladesh. Trans R Soc Trop Med Hyg 1993;87:84–6.

Evaluation of HIV Case Surveillance Through the Use of Non-Name Unique Identifiers — Maryland and Texas, 1994–1996

Notifiable disease reporting laws or regulations in states and territories require reporting of acquired immunodeficiency syndrome (AIDS) cases, including patient and physician names, to state or local health authorities. As of January 1, 1998, a total of 31 states were conducting name-based human immunodeficiency virus (HIV) case surveillance by using the same methods as surveillance for AIDS. However, because of concerns about name-based HIV surveillance, Maryland and Texas implemented HIV surveillance using non-name unique identifiers (UI)*. This report summarizes a 3-year collaboration by CDC and these states to evaluate UI surveillance for HIV infec-

[†]Albendazole is currently approved by the Food and Drug Administration for treatment of neurocysticercosis and hydatid disease.

^{*}Reporting in Maryland is exempted for nonstate residents; persons who are tested at anonymous test sites; are blood, semen, or tissue donors; and participants of certain research projects. No exemptions to reporting exist in Texas.

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tion; the findings indicate some limitations to the use of a Social Security numberbased UI for HIV surveillance.

In both Maryland and Texas, UI surveillance for HIV was implemented in early 1994, and both used the same 12-digit numeric UI code (comprising the last four digits of the patient's Social Security number [SSN], six-digit [month/day/year] date of birth [DOB], one-digit code for race/ethnicity, and one-digit code for sex). HIV-infection reports included residence data, diagnosing facility, and date of test, but did not include mode of HIV exposure. In both states, UI HIV surveillance databases were maintained separately from name-based AIDS surveillance databases.

Evaluation criteria included the proportion of reports with full UI codes, timeliness and completeness of HIV reporting, and potential for matching the UI-based case reports to alternate databases. In Texas, selected HIV reports also were evaluated for ability to follow back UI reports to patient records; in Maryland, provider compliance with maintaining patient surveillance logs was assessed. During July 1994–December 1996, Maryland reported 6412 AIDS cases and received 9971 HIV-infection reports, and Texas reported 12,041 AIDS cases and received approximately 23,000 HIVinfection reports.

Maryland

In 1993, the Maryland legislature mandated UI reporting of both positive HIV tests and patients with CD4+ T-lymphocyte counts of <200 cells/ μ L (CD4+)[†]. Health-care providers requesting HIV or CD4+ tests are required to construct the UI code for each patient, include the code on the laboratory slip, and record it in a surveillance log that matches the UI to patient identifiers (e.g., medical record number, patient name, or other patient code) for purposes of case investigation and follow up. Laboratories licensed by Maryland are required to submit the UI-based reports to the state health department through the local health departments.

Of 9971 HIV-infection reports entered during July 1994–December 1996, all UI elements were present for 7119 (71%) (Table 1). Element-specific presence ranged from 78% (SSN) to 98% (DOB and sex). The proportion of reports with full UI increased during July 1994–June 1996, and declined slightly during July–December 1996. The median time from date of HIV test to receipt of report by the state health department was 20 days (range: 1–847 days). During October–November 1997, all 72 providers in nine counties of eastern Maryland (the counties reported 3% of AIDS cases in Maryland in 1996) for whom laboratories had submitted HIV-infection reports were contacted to determine the proportion of providers who maintain the required surveillance log linking UI to patient identifiers; 32 (44%) of these providers maintained logs.

Completeness of HIV-infection reporting was estimated by comparison to cases of AIDS reported in the AIDS surveillance registry. Of AIDS cases with dates of HIV diagnosis from July 1995 through June 1996, data elements to construct UI were available for 633 (85%) cases. Of these, 319 (50%) matched to HIV-infection reports with full UI in the UI database (Table 2).

Data from the Maryland HIV counseling and testing (C&T) system (excluding sites offering only anonymous HIV tests) were used to evaluate the proportion of records

⁺HIV-infected persons with a CD4+ T-lymphocyte count of <200 cells/μL meet the 1993 expanded AIDS surveillance case definition and are reportable by name for AIDS surveillance.

HIV Case Surveillance — Continued

Reports/Data element	State	July– Dec. 1994	Jan.– June 1995	July– Dec. 1995	Jan.– June 1996	July– Dec. 1996	Overall
Total no. reports	MD	2,238	1,691	1,866	1,881	2,295	9,971
	TX*	3,932	3,399	3,597	2,852	2,339	16,119
Data element [†]							
Social Security	MD	69.6	73.1	81.2	83.5	84.5	78.4
number	TX	56.7	68.6	65.0	69.5	75.2	66.0
Date of birth	MD	95.2	96.3	98.7	99.3	98.8	97.6
	TX	88.4	89.8	93.1	96.8	97.6	92.6
Sex	MD	96.8	97.2	98.7	99.2	99.4	98.3
	TX	91.5	97.5	98.4	99.1	97.9	96.6
Race/Ethnicity	MD	85.8	88.5	91.6	94.0	89.9	89.8
	TX	80.8	91.6	94.4	97.1	95.4	91.1
% Reports with full UI	MD	61.3	65.9	74.9	78.5	76.5	71.4
	TX	51.8	61.9	61.6	66.5	71.3	61.6

TABLE 1. Number of reports of HIV infection and percentage of reports that included data elements for unique identifiers (UIs), by reporting period — Maryland (MD) and Texas (TX), July 1994–December 1996

*Excludes approximately 7000 records that had three or more missing UI data elements.

with full UI and completeness of HIV-infection reporting. In early 1995, counselors were instructed to obtain UI code information from clients and record the UI on the HIV C&T record. During 1995–1996, a total of 1093 records with a positive HIV test were entered into the C&T database; of these, all UI elements were present for 94%. HIV C&T reports for persons who had HIV diagnosed from July 1995 through June 1996 were matched to the UI database. Of the 528 reports, 276 (52%) matched.

Texas

In 1994, the Texas Board of Health amended regulations to require named reporting of HIV-infected children aged <13 years and UI reporting of HIV-infected adolescents and adults. Both health-care providers ordering an HIV test and laboratories performing the test report confirmed HIV infections to the Texas Department of Health (TDH) through the local health departments. Neither providers nor laboratories are required to maintain registries linking UI to patient identifiers.

Approximately 23,000 HIV-infection reports were received at TDH during the evaluation period. Since 1995, TDH excluded approximately 7000 paper HIV reports with three or more missing UI data elements. Of 16,119 HIV-infection reports entered into the UI database, all UI elements were present for 9923 (62%) (Table 1). Elementspecific presence ranged from 66% (SSN) to 97% (sex). Overall, 60% of reports were submitted in periodic batches, which had a longer time from date of HIV test to receipt by TDH (median: 173 days; range: 26–974 days) than the 40% of reports submitted individually (median: 59 days; range: 2–906 days).

Completeness of HIV-infection reporting was estimated by comparison to AIDS surveillance data using the same methodology as in Maryland. Data elements to construct UI were available for 1762 (79%) of AIDS cases with dates of HIV diagnosis in the

HIV Case Surveillance — Continued

Characteristic	Maryland (n=9,971)	Texas (n=16,119)
Completeness of reporting		
HIV*	50.4	26.0
CD4+ T-lymphocyte count*	44.4	NA [†]
HIV§	52.3	NA
Availability of UI data elements in alternate databases		
Birth¶	No	No
Death	Yes	Yes
Sexually transmitted		
disease	No	No
Tuberculosis	No	No
Drug assistance**	Yes	Yes
Medical assistance ^{††}	Yes	No
Hospital discharge	No	No
Source of HIV report		
Public	30% ^{§§}	77% ^{§§}
Private	70%¶¶	23%***

TABLE 2. Percentage completeness of HIV-infection reporting, availability of unique identifier (UI) data elements in alternate databases, and sources of report — Maryland and Texas, July 1994–December 1996

*AIDS cases reported through July 1997 compared with the UI database.

[†]Not available.

[§]HIV cases diagnosed from July 1995 through June 1996 in HIV counseling and testing sites compared with the UI database.

[¶]Used for pediatric AIDS surveillance only.

**Federal- and state-funded medication program.

^{††}Federal- and state-funded medical-assistance program.

§§ Includes local health departments and state laboratory.

"Includes community-based organizations and private clinics and laboratories.

***Includes community-based organizations, hospitals, private physicians, clinics, and laboratories.

specified period (Table 2). Of these, 454 (26%) matched to HIV-infection reports with full UI in the UI database.

To evaluate the feasibility of epidemiologic follow up, TDH sampled 765 HIVinfection reports submitted during January 1995–June 1996, in six areas of the state, reflective of variation in geography, demography, HIV morbidity, and reporting sources. Of these, 456 (60%) could be matched to a client record using any combination of UI (including records without full UI), health-care provider name, date of test, residential information, and other locally available information. Matched records that were missing the SSN data element (n=208) were reviewed to determine whether these data could be located. SSN could not be located for 120 (58%) of these records. *Reported by: L Solomon, DrPH, L Eldred, DrPH, J Markowitz, PhD, P Ryan, MS, G Benjamin, MD, Maryland Dept of Health and Mental Hygiene. AS Robbins, PhD, DW Hamaker, SA King, MA, SK Melville, MD, MC Thomas, MS, DM Simpson, MD, State Epidemiologist, Texas Dept of Health. Div of HIV/AIDS Prevention–Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention, CDC.*

HIV Case Surveillance — Continued

Editorial Note: HIV and AIDS surveillance data are needed to provide reliable population-based data to guide public health programs. During 1995–1996, the first declines in the incidence of AIDS-opportunistic infections and AIDS deaths were reported in the United States (6% and 23%, respectively), in part, as a result of increasingly effective HIV therapy (1). On the basis of revised HIV treatment guidelines (2), the impact of treatment advances on AIDS trends is expected to continue and will reduce the usefulness of AIDS data alone to monitor HIV-infection trends and morbidity. CDC and other public health and advocacy organizations have recognized the need for national HIV case surveillance while continuing to discuss the relative merits of HIV surveillance methods based on numeric codes compared to the name-based approach employed for AIDS surveillance (1,3).

CDC uses established criteria to evaluate performance of public health surveillance systems to provide accurate data to target prevention and care programs (4). States conduct active surveillance using existing name-based clinical and public health records to decrease the reporting burden on providers, eliminate duplicate reports, and facilitate epidemiologic follow-up. These methods enable AIDS surveillance to attain high performance standards as reflected by completeness of reporting (>85%) (5) and documentation of risk exposures (\geq 93% of cases) (6). Evaluation of name-based HIV surveillance has shown 74%–97% completeness of reporting (7; CDC, unpublished data, 1997), and documentation of risk exposures (\geq 76% of cases) (6). Secure and confidential surveillance practices are required as a condition for receipt of federal resources for HIV and AIDS surveillance. At the state level, the most comprehensive protections of medical data apply to government-held data, and most specifically to HIV-related data (8). Names are removed before encoded and encrypted AIDS or HIV surveillance data are transmitted to CDC.

The evaluations in Maryland and Texas indicated that the use of UIs limits the performance of an HIV surveillance system and complicates efforts to collect riskbehavior information. Both systems demonstrated timely reporting. Although data from both states indicated increases in reporting of the SSN data element during the evaluation period, overall 22% of reports in Maryland and 34% in Texas were missing the SSN element, which contributed to a high rate of incomplete case reporting. The follow-back investigation in Texas suggests that SSNs are not readily available in client or medical records but, in the controlled environment of the Maryland HIV C&T system, counselors were able to collect SSNs for most clients. The completeness of reporting also may be affected by the ability of providers and laboratories to use UIs as part of routine HIV-testing practices. For example, one large laboratory providing HIV-testing services in Maryland did not report HIV infections during the evaluation period. The difficulty in collecting HIV data when persons are tested out of state also may affect completeness of reporting and the ability to eliminate duplicate reports. Maryland is continuing to evaluate its UI surveillance system, and Texas is exploring alternative HIV surveillance systems with input from community groups.

Effective HIV surveillance systems must include HIV risk information; however, this information often is not available at the time of the initial UI case report, and follow-up with health-care providers is necessary. To supply follow-up information, health-care providers must use lists or other mechanisms to link the UI to patient identifiers. The UI approach complicates efforts to collect this information and increases the number of lists of HIV-infected persons that could be disclosed in a breach of confidentiality.

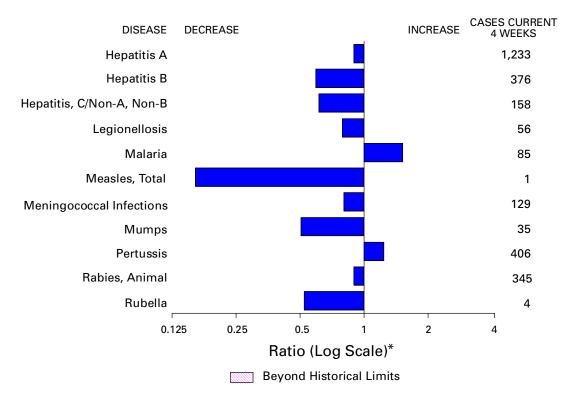


FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending December 27, 1997, with historical data — United States

*Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

TABLE I. Summary — provisional cases of selected notifiable diseases, United States, cumulative, week ending December 27, 1997 (52nd Week)

	Cum. 1997		Cum. 1997
Anthrax Brucellosis Cholera Congenital rubella syndrome Cryptosporidiosis* Diphtheria Encephalitis: California* eastern equine* St. Louis* western equine* Hansen Disease Hantavirus pulmonary syndrome* [†] Hemolytic uremic syndrome, post-diarrheal* HIV infection, pediatric* [§]	0 75 10 4 1,950 5 118 10 12 - 109 18 60 214	Plague Poliomyelitis, paralytic [¶] Psittacosis Rabies, human Rocky Mountain spotted fever (RMSF) Streptococcal disease, invasive Group A Streptococcal toxic-shock syndrome* Syphilis, congenital** Tetanus Toxic-shock syndrome Trichinosis Typhoid fever Yellow fever	4 1 37 2 396 1,404 31 525 42 133 9 344

-:no reported cases *Not notifiable in all states. †Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases (NCID). \$Updated monthly to the Division of HIV/AIDS Prevention–Surveillance, and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP), last update November 25, 1997. ¶One suspected case of polio with onset in 1997 has also been reported to date. **Updated from reports to the Division of STD Prevention, NCHSTP.

					Esche coli O	erichia 157:H7			Нера	atitis
		DS		mydia	NETSS [†]	PHLIS [§]		rrhea	C/N/	-
Reporting Area	Cum. 1997*	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1997	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996
UNITED STATES	53,031	66,213	458,353	437,869	2,292	1,530	284,427	322,818	3,132	3,719
NEW ENGLAND	2,252	2,746	17,362	16,967	200	127	5,687	6,296	56	113
Maine N.H.	51 40	49 93	1,025 799	931 752	17 13	16	66 95	55 162	- 8	-7
Vt.	32	19	421	398	8	3	51	47	2	26
Mass. R.I.	808 142	1,306 171	7,390 1,822	6,791 1,832	109 10	93	2,118 392	2,163 486	39 7	74 6
Conn.	1,179	1,108	5,905	6,263	43	15	2,965	3,383	-	-
MID. ATLANTIC	16,043	18,114	58,700	57,172	141	52	36,918	41,735	368	337
Upstate N.Y. N.Y. City	2,390 8,610	2,422 9,942	N 30,377	N 26,455	97 15	- 8	6,147 14,213	7,606 13,008	287	272 3
N.J.	3,044	3,584	9,524	12,261	29	25	7,176	8,721	-	-
Pa.	1,999	2,166	18,799	18,456	N	19	9,382	12,400	81	62
E.N. CENTRAL Ohio	3,957 798	5,171 1,153	69,621 20,254	83,652 20,653	405 108	280 52	42,597 12,710	57,320 14,946	503 20	490 35
Ind.	488	591	9,487	10,100	81	46	6,175	6,458	11	8
III.	1,715	2,192	10,838	22,459	70	31	5,272	15,776	83	93
Mich. Wis.	716 240	964 271	20,151 8,891	20,277 10,163	146 N	103 48	14,488 3,952	15,267 4,873	389	354
W.N. CENTRAL	1,055	1,600	32,731	31,705	518	402	14,834	15,822	163	111
Minn.	194	304	7,416	5,608	214	203	2,752	2,698	4	10
lowa Mo.	100 505	92 851	4,647 11,922	4,165 12,020	119 55	74 69	1,245 7,792	1,144 8,492	35 108	53 23
N. Dak.	12	12	749	1,022	15	12	52	42	3	-
S. Dak. Nebr.	8 90	14 93	1,438 2,560	1,536 2,878	28 61	32	169 1,087	176 1,213	- 3	- 9
Kans.	90 146	234	3,999	4,476	26	12	1,087	2,057	10	16
S. ATLANTIC	13,084	16,514	89,547	53,331	214	135	87,862	95,569	273	235
Del.	214	285	1,276	1,148	5	4	1,228	1,432	-	1
Md. D.C.	1,811 955	2,246 1,257	7,516 N	U N	25 2	14	12,931 4,256	11,316 4,470	22	4
Va.	1,113	1,146	11,340	11,754	N	41	8,668	9,292	24	17
W. Va. N.C.	121 795	121 898	2,882 17,097	2,301 U	N 74	1 38	928 16,675	850 18,229	17 50	9 46
S.C.	754	842	12,359	Ŭ	13	8	11,235	10,984	38	34
Ga. Fla.	1,604 5,717	2,422 7,297	12,244 24,833	13,333 24,795	41 46	- 29	14,384 17,557	19,810 19,186	U 122	- 124
E.S. CENTRAL	1,908	2,280	31,504	32,507	95	39	31,709	35,982	334	590
Ky.	338	401	6,207	6,687	30	-	3,960	4,229	15	29
Tenn. Ala.	745 512	821 608	12,438 8,587	13,121 8,352	46 15	39	10,975 11,917	11,710 13,334	232 13	400 8
Miss.	313	450	4,272	4,347	4	-	4,857	6,709	74	153
W.S. CENTRAL	5,663	6,793	59,878	59,215	71	26	39,190	40,942	489	515
Ark. La.	216 997	267 1,464	2,128 10,030	1,663 7,330	10 7	5 12	3,571 9,777	5,050 7,976	11 241	8 292
Okla.	275	261	7,374	7,219	12	6	4,725	4,792	7	7
Tex.	4,175	4,801	40,346	43,003	42	3	21,117	23,124	230	208
MOUNTAIN Mont.	1,527 41	2,014 34	22,765 1,084	29,642 1,232	239 24	132	7,925 47	7,459 35	484 23	555 20
Idaho	50	38	1,646	1,505	38	23	156	98	84	99
Wyo. Colo.	14 352	7 519	611 1,896	621 7,282	17 83	12 57	52 2,138	41 1,367	230 40	179 64
N. Mex.	163	206	3,369	3,862	7	-	1,136	891	58	77
Ariz.	374	591	10,550	10,686	N	30	3,596	3,705	25	76
Utah Nev.	134 399	194 425	1,728 1,881	1,554 2,900	59 11	10	269 531	280 1,042	5 19	19 21
PACIFIC	7,542	10,980	76,245	73,678	409	332	17,705	21,693	462	773
Wash.	617	762	9,408	9,236	130	131	1,917	2,020	31	66
Oreg. Calif.	286 6,510	461 9,522	5,009 58,580	5,347 55,926	84 183	95 94	735 14,204	871 17,842	4 266	8 479
Alaska	40	36	1,549	1,345	12	3	371	459	-	3
Hawaii	89	199	1,699	1,824	N	9	478	501	161	217
Guam P.R.	2 1,975	4 2,238	193 U	355 U	N 41	- U	27 526	63 648	- 150	6 180
V.I.	95	18	N	N	N	U	-	-	-	-
Amer. Samoa	- 1	-	- N	- NI	N	U	- 17	- 11	- ว	-
C.N.M.I.	1	-	N	N	N	U	17	11	2	-

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending December 27, 1997, and December 28, 1996 (52nd Week)

N: Not notifiable U: Unavailable -: no reported cases C.N.M.I.: Commonwealth of Northern Mariana Islands

*Updated monthly to the Division of HIV/AIDS Prevention–Surveillance, and Epidemiology, National Center for HIV, STD, and TB Prevention, last update November 25, 1997. [†]National Electronic Telecommunications System for Surveillance. [§]Public Health Laboratory Information System.

	Legion	ellosis	Lyı Dise		Mal	aria	Syp (Primary &	hilis Secondary)	Tubero	culosis	Rabies, Animal
Reporting Area	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997
UNITED STATES	1,033	1,198	10,622	16,455	1,756	1,800	7,787	11,344	16,905	21,168	7,674
NEW ENGLAND	82	80	2,877	4,095	96	84	128	195	441	485	1,230
Maine N.H.	2 7	5 4	11 38	63 47	1 10	10 4	2	2 1	11 15	21 21	221 43
Vt. Mass.	13 27	5 34	8 370	26 321	2 30	8 32	- 68	- 85	6 254	4 262	113 278
R.I.	15	32	409	534	11	12	2	4	36	39	41
Conn. MID. ATLANTIC	18 214	N 263	2,041 6,236	3,104 10,305	42 433	18 467	56 352	103 509	119 3,106	138 3,969	534 1,652
Upstate N.Y.	72	80	2,477	4,900	70	96	39	76	438	535	1,205
N.Y. City N.J.	12 20	19 15	119 1,637	401 2,190	252 78	269 68	83 122	133 177	1,591 686	2,035 816	U 189
Pa.	110	149	2,003	2,814	33	34	108	123	391	583	258
E.N. CENTRAL Ohio	312 121	360 116	96 61	498 32	148 19	170 15	668 211	1,573 584	1,579 244	2,110 301	177 116
Ind.	54	51	29	32	16	15	166	207	150	202	13
III. Mich.	28 92	38 109	6	10 28	55 43	83 41	71 128	420 183	752 311	1,060 433	20 28
Wis.	17	46	U	396	15	16	92	179	122	114	-
W.N. CENTRAL Minn.	70 3	71 15	233 196	365 251	68 36	51 26	181 23	340 47	576 152	547 131	490 61
lowa Mo.	12 31	11 18	10 20	19 52	10 11	3 11	8 113	23 224	73 239	70 224	157 25
N. Dak.	2	-	-	52	3	1	-	- 224	12	8	84
S. Dak. Nebr.	2 15	3 18	1 2	- 5	3 1	- 3	1 7	- 10	19 22	19 22	74 2
Kans.	5	6	4	36	4	7	29	36	59	73	87
S. ATLANTIC Del.	131 13	197 12	768 105	823 173	361 5	340 4	3,155 20	3,780 35	3,180 18	4,014 43	3,022 54
Md.	28	39	490	447	86	87	891	733	308	319	596
D.C. Va.	5 27	9 54	10 62	3 57	20 68	9 60	112 232	123 393	103 305	139 349	5 667
W. Va. N.C.	N 14	N 12	10 34	12 66	1 20	6 30	3 721	9 1,052	54 430	57 554	88 852
S.C.	8	8	3	9	18	13	360	384	260	350	175
Ga. Fla.	2 33	3 60	7 47	1 55	52 91	38 93	521 295	683 368	595 1,107	790 1,413	311 274
E.S. CENTRAL	53	59	83	83	35	42	1,610	2,412	1,190	1,436	270
Ky. Tenn.	12 33	11 26	14 44	26 24	8 11	12 14	136 747	154 850	184 358	259 504	28 149
Ala.	4	5	11	9	10	8	410	530	412	423	88
Miss. W.S. CENTRAL	4 33	17 53	14 100	24 175	6 57	8 158	317 1,195	878 1,815	236 2,396	250 2,898	5 340
Ark.	-	1	25	27	5	2	140	262	179	225	54
La. Okla.	7 3	4 16	6 34	9 42	16 8	12 3	366 117	486 177	270 173	350 200	5 112
Tex.	23	32	35	97	28	141	572	890	1,774	2,123	169
MOUNTAIN Mont.	63 1	58 1	23	9	66 2	65 7	179	165	510 17	711 19	188 50
ldaho	2	-	4	2	1	-	1	4	16	15	-
Wyo. Colo.	1 18	7 12	5 6	3	2 30	7 26	- 14	2 26	2 75	7 104	31 28
N. Mex. Ariz.	3 12	2 21	1 4	1	8 11	3 9	16 134	8 102	53 269	89 282	12 53
Utah	19	8	1	1	3	5	5	3	32	58	6
Nev. PACIFIC	7 75	7 57	2 206	2 102	9 492	8 423	9 319	20 555	46 3,927	137 4,998	8 305
Wash.	10	8	10	18	49	41	16	9	264	285	-
Oreg. Calif.	- 64	43	21 173	19 64	24 405	24 343	9 292	9 533	154 3,291	190 4,227	14 262
Alaska Hawaii	- 1	1 5	2	- 1	5 9	3 12	1 1	- 4	73 145	96 200	29
Guam	-	1	-	-	-	-	3	4	145	107	-
P.R.	-	-	-	-	6	2	238	209	212	222	66
V.I. Amer. Samoa	-	-	-	-	-	1	-	-	-	-	-
C.N.M.I.	-	-	-	-	-	-	9	1	2	-	-

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States,
weeks ending December 27, 1997, and December 28, 1996 (52nd Week)

N: Not notifiable U: Unavailable -: no reported cases

		ienzae,		epatitis (Vi						les (Rubec		
	inva Cum.	sive Cum.	Cum.	A Cum.	L Cum.	3 Cum.	Indi	genous Cum.	lmp	ported [†] Cum.	To Cum.	tal Cum.
Reporting Area	1997*	1996	1997	1996	1997	1996	1997	1997	1997	1997	1997	1996
UNITED STATES	1,041	1,170	27,595	31,032	8,656	10,637	-	78	-	55	133	508
NEW ENGLAND Maine	63 5	55 1	625 62	456 28	147 6	255 8	-	11	-	8 1	19 1	17
N.H.	12	13	34	22	17	21	-	1	-	-	1	-
Vt. Mass.	3 38	2 36	15 241	12 229	7 56	14 111	-	10	-	- 6	16	2 12
R.I. Conn.	3 2	2 1	129 144	26 139	16 45	19 82	-	-	-	- 1	- 1	1 2
MID. ATLANTIC	145	213	1,843	1,985	1,275	1,413	-	19	-	8	27	38
Upstate N.Y. N.Y. City	41 35	50 57	367 694	438 609	322 428	358 491	-	2 9	-	3 2	5 11	12 11
N.J. Pa.	49 20	65 41	246 536	394 544	201 324	279 285	-	3 5	-	- 3	3 8	3 12
E.N. CENTRAL	160	191	2,843	2,619	912	1,103	-	6	-	3	9	21
Ohio Ind.	86 19	95 21	328 322	785 367	93 93	120 143	-	-	-	-	-	6
III.	38	50	702	763	227	335	-	6	-	1	7	3
Mich. Wis.	15 2	12 13	1,336 155	506 198	454 45	416 89	-	-	-	2	2	3 9
W.N. CENTRAL Minn.	65 44	63 48	2,182 196	2,656 176	469 44	572 94	-	13 3	-	5 5	18 8	24 19
lowa	7	4	484	334	47	74	-	1	-	-	1	1
Mo. N. Dak.	10	8	1,100 11	1,414 140	323 5	326 2	Ū	1	Ū	-	1 -	3
S. Dak. Nebr.	2 1	1 1	24 114	43 156	1 16	5 39	-	8	-	-	8	-
Kans.	1	1	253	393	33	32	-	-	-	-	-	1
S. ATLANTIC Del.	172	273 2	2,135 31	1,960 21	1,284 6	1,573 9	-	2	-	13	15	12 1
Md. D.C.	58	76 5	210 36	256 39	186 30	169 32	-	-	-	2 1	2 1	2
Va.	13 4	11	229	218	127	163	-	-	-	1	1	3
W. Va. N.C.	21	11 26	12 209	19 204	16 265	36 337	-	-	-	2	2	2
S.C. Ga.	4 42	5 52	108 656	57 414	97 148	101 61	Ū	-	Ū	1 1	1 1	- 3
Fla.	30	85	644	732	409	665	-	2	-	5	7	1
E.S. CENTRAL Ky.	48 6	45 6	631 75	1,273 53	688 40	914 76	-	-	-	-	-	2
Tenn. Ala.	27 15	25 13	400 87	778 217	446 79	516 78	-	-	-	-	-	2
Miss.	-	1	69	225	123	244	-	-	-	-	-	-
W.S. CENTRAL Ark.	51 1	44	5,501 216	6,807 500	1,185 62	1,616 93	-	3	-	5	8	27
La. Okla.	13 32	6 32	238 1,416	261 2,586	170 51	209 56	-	-	-	- 1	- 1	1
Tex.	5	6	3,631	3,460	902	1,258	-	3	-	4	7	26
MOUNTAIN Mont.	94	57 1	4,333 71	4,573 130	887 12	1,164 21	-	6	-	2	8	157 -
ldaho Wyo.	1 4	1	144 41	247 41	54 40	88 45	Ū	-	Ū	-	-	1 1
Colo.	19 10	16	406 354	512 355	153 258	132 417	-	-	-	-	-	7 17
N. Mex. Ariz.	33	11 20	2,353	1,767	201	237	-	5	-	-	5	8
Utah Nev.	3 24	8	547 417	1,073 448	92 77	129 95	-	- 1	-	1 1	1 2	118 5
PACIFIC	243	229	7,502	8,703	1,809	2,027	-	18	-	11	29	210
Wash. Oreg.	6 35	10 33	673 378	1,001 875	80 108	158 129	-	1	-	1	2	38 14
Calif. Alaska	188 7	178 6	6,283 34	6,653 54	1,589 21	1,710 16	-	15	-	8 -	23	46 63
Hawaii	7	2	134	120	11	14	-	2	-	2	4	49
Guam P.R.	-	2	- 257	7 292	3 1,376	1 1,195	U -	-	U -	-	-	- 3
V.I. Amer. Samoa	-	-	-	41	-	44	U U	-	U U	-	-	-
C.N.M.I.	6	10	1	1	34	5	Ŭ	1	Ŭ	-	1	-

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination,
United States, weeks ending December 27, 1997,
and December 28, 1996 (52nd Week)

N: Not notifiable U: Unavailable -: no reported cases

 * Of 241 cases among children aged <5 years, serotype was reported for 126 and of those, 48 were type b.

[†]For imported measles, cases include only those resulting from importation from other countries.

		jococcal ease		Mumps			Pertussis			Rubella	
Reporting Area	Cum. 1997	Cum. 1996	1997	Cum. 1997	Cum. 1996	1997	Cum. 1997	Cum. 1996	1997	Cum. 1997	Cum. 1996
UNITED STATES	3,078	3,437	5	606	751	55	5,461	7,796	3	160	238
NEW ENGLAND	198	171	-	12	5	6	974	1,866	1	2	27
Maine N.H.	18 18	15 13	-	- 1	- 1	- 3	11 133	55 197	-	-	-
Vt.	4	4	-	-	1	2	256	280	-	-	2
Mass. R.I.	98 20	71 18	-	4 6	1 1	1	522 17	1,245 40	-	1	21
Conn.	40	50	-	1	1	-	35	49	1	1	4
MID. ATLANTIC Upstate N.Y.	318 73	381 102	-	57 12	96 28	1 1	384 152	952 533	-	31 4	13 5
N.Y. City	45	56	-	3	20	-	62	61	-	27	5
N.J. Pa.	72 128	79 144	-	7 35	4 44	-	11 159	31 327	-	-	2 1
E.N. CENTRAL	456	475	-	80	135	14	517	837	-	5	3
Ohio	163	159	-	35	52	5	164	289	-	-	-
Ind. III.	58 148	64 142	-	14 13	8 24	9	85 124	128 192	-	- 2	- 1
Mich. Wis.	53 34	51 59	-	15 3	48 3	-	61 83	59 169	-	- 3	2
WIS. W.N. CENTRAL	229	59 264	-	3 18	3 24	- 16	83 579	573	1	3 2	-
Minn.	34	39	-	6	7	15	369	433	-	-	-
lowa Mo.	48 104	56 98	-	10	3 10	- 1	108 68	32 74	- 1	2	-
N. Dak.	2	5	U	-	2	U	2	1	U	-	-
S. Dak. Nebr.	5 14	10 29	-	2	-	-	5 14	4 15	-	-	-
Kans.	22	27	-	-	2	-	13	14	-	-	-
S. ATLANTIC Del.	555 5	659 3	2	85	131	6	436 1	793 26	1	83	101
Md.	42	58	-	10	37	4	124	278	-	-	-
D.C. Va.	9 58	5 67	-	- 19	- 19	-	3 56	4 108	-	1 1	1 2
W. Va. N.C.	18 97	18 79	-	- 12	- 27	-	6	7 186	-	- 59	- 86
S.C.	59	65	-	11	7	-	118 30	49	-	59 19	1
Ga. Fla.	105 162	147 217	U 2	10 23	9 32	U 2	13 85	35 100	U 1	- 3	- 11
E.S. CENTRAL	236	246	-	23	23	-	141	202	-	-	2
Ky.	48	31	-	3	-	-	61	142	-	-	-
Tenn. Ala.	85 84	65 95	-	6 9	1 6	-	39 33	24 26	-	-	2
Miss.	19	55	-	9	16	-	8	10	-	-	N
W.S. CENTRAL Ark.	278 32	365 35	-	75 1	67 1	-	295 60	201 14	-	4	9
La.	47	66	-	16	21	-	20	15	-	-	1
Okla. Tex.	44 155	46 218	-	- 58	1 44	-	48 167	21 151	-	- 4	- 8
MOUNTAIN	185	183	1	57	25	7	1,288	660	-	6	9
Mont. Idaho	9 13	9 25	- 1	- 5	-	- 2	19 621	37 115	-	- 1	2
Wyo.	4	4	ΰ	1	1	U	7	8	U	-	-
Colo. N. Mex.	49 29	44 27	N	3 N	5 N	1 3	342 193	336 64	-	-	3
Ariz.	46	37	-	33	1	-	41	33	-	5	3
Utah Nev.	16 19	18 19	-	8 7	3 15	- 1	26 39	26 41	-	-	- 1
PACIFIC	623	693	2	195	245	5	847	1,712	-	27	74
Wash. Oreg.	92 126	116 123	N	21 N	26 N	5	406 10	830 64	-	5	15 1
Calif.	395	437	2	147	185	-	403	780	-	14	55
Alaska Hawaii	3 7	9 8	-	4 23	3 31	-	14 14	3 35	-	- 8	- 3
Guam	1	5	U	1	10	U	-	-	U	-	-
P.R. V.I.	10	13	- U	7	2 2	- U	2	3	- U	-	-
Amer. Samoa	-	-	U	-	2 -	Ű	-	-	U	-	-
C.N.M.I.	-	-	U	4	-	U	-	-	U	-	-

TABLE III. (Cont'd.) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending December 27, 1997, and December 28, 1996 (52nd Week)

N: Not notifiable U: Unavailable -: no reported cases

	ļ	All Cau	ises, By	/ Age (Y	ears)		P&I [†]			All Cau	ises, Βγ	/ 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, Mass. New Bedford, Mass. New Haven, Conn. Providence, R.I. Somerville, Mass. Springfield, Mass. Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J.	54 U 3 44 24 65 2,284 45 27 74 28 19	407 129 22 12 23 34 18 5 14 41 U 3 37 18 51 1,587 20 58 22 20 58 22 20 58 22 20	31 4 5 10 3 2 3 4 U - 6 8 396 5 7 12 4 2	23 8 5 - - 3 U - 4 3 202 1 - 3 1 1	15 7 1 1 1 2 U - 1 1 69 3 - 1	11 2 - - - - - - - - - - - - - - - - - -	47 21 3 1 3 4 U 9 136 4 1 7 7 1	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla. Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, Fla. Tampa, Fla. Washington, D.C. Wilmington, Del. E.S. CENTRAL Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala.	91 37 178 80 49	554 U 100 55 733 22 57 113 92 372 0 47 66 20 20 115 52 35	161 U 32 14 20 U 12 4 14 296 - 127 U 133 21 8 40 23 8 40 23 8	61 U 13 2 9 U U 4 2 1 9 U U 4 2 1 9 U U 4 2 1 9 U U 7 3 7 U 7 3 5 11 1	21 U 3 3 1 U U 3 - 1 6 4 - 19 U 1 - 1 9 4 4	19 U 11 3 1 U U - 1 - 3 - 13 U 1 1 3 3 2 1	52 U 11 5 4 U U 1 3 8 8 12 - 32 U 5 2 3 10 - 8
Erie, Pa. Jersey City, N.J. New York City, N.Y. Newark, 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.	47 38 1,323 59 13 200 35 43 113 18 25 130 29 18 U	39 25 926 27 10 91 23 84 16 22 103 21 16 U	7 236 17 1 47 5 22 1 3 15 2 1 15 2	2 4 114 8 2 40 3 3 5 1 - 8 5 1 U	2 230 3 21 1 2 2 1 2 1 U	- 17 4 - 1 3 1 - - 2 - - - U	2 1 64 3 10 5 2 11 2 3 13 5 2 U	Nashville, Tenn. W.S. CENTRAL Austin, Tex. Baton Rouge, La. Corpus Christi, Tex. Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La. San Antonio, Tex. Shreveport, La. Tulsa, Okla.	148 68 56 230 42 198 126 U 69	37 688 33 20 95 51 40 134 32 113 84 U 47	16 230 12 11 3 25 12 14 66 8 41 26 U 12	9 98 5 4 1 21 21 22 22 10 U 7	30 1 3 1 3 1 1 4 10 4 U 2	2 25 3 2 4 - 5 - 9 1 U 1	4 69 9 3 4 2 4 23 3 15 0 6
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Celveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Gary, Ind. Grand Rapids, Mich Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio	132 17 80 39 45 46 62 U	1,084 29 263 644 711 127 71 85 0 0 0 1 51 97 8 565 256 326 324 30 324 30 0 20 20 20 20 20 20 20 20 20 20 20 20	4 93 127 35 19 30 0 1 4 21 9 9 10 8 7 10 0	108 2 1 3 6 8 13 3 9 U U 2 2 7 7 2 1 3 6 U	45 2 - 183 - 5 1 5 UU - 24 - 21 - 1 1 U 2	38 1 10 15 2 - 3 UU - - 3 - 6 1 - 3 2 U	101 4 32 9 5 20 5 8 U U 4 2 3 3 2 2 2 U	MOUNTAIN Albuquerque, N.M. Boise, Idaho Colo. Springs, Colo Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, Utah Tucson, Ariz. PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawaii Long Beach, Calif. Dasadena, Calif. Portland, Oreg. Sacramento, Calif.	141 180 33 1599 18 116 128 1,207 19 104 U 75 69 U 26 U 202	651 666 288 399 87 108 266 108 14 876 15 74 57 57 57 57 57 57 0 23 0 0 23 0 0 23 77 77	170 8 5 8 26 54 27 17 20 180 2 180 11 13 U 2 U 314	78 6 13 13 13 2 10 8 93 2 3 U 4 4 U 15 11	27 - - - - - - - - - - - - - - - - - - -	27 2 7 2 6 1 4 3 28 4 U 3 · U U 2 3	80 6 4 5 11 9 20 8 11 9 0 2 13 0 4 0 2 13 0 4 0 2 13 0 4 0 2 19
W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	568 U 27 8 110 31 94 85 89 79 45	402 U 19 56 25 73 63 65 61 34	U 6 18 6 9 18 14 10	37 U 2 10 - 7 3 4 7 2	12 U 6 3 - 3	14 U 2 2 1 6 1 2	36 U 2 - 5 5 13 3 - 4 4	San Diego, Calif. San Francisco, Calif San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash. TOTAL	197 41 112 29 87	77 98 150 30 64 23 66 6,621	14 27 23 5 23 2 8 1,733	11 13 17 3 14 2 5 737	2 1 2 8 - 3 268	3 5 1 3 2 5 205	9 18 22 4 1 2 8 666

TABLE IV. Deaths in 122 U.S. cities,* week ending December 27, 1997 (52nd 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 or more. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included. *Pneumonia and influenza. *Because of changes in reporting methods in 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.

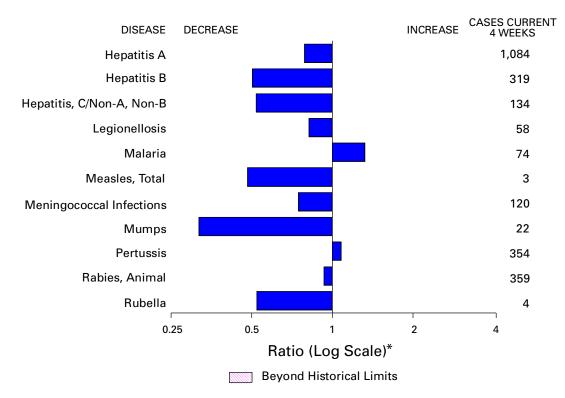


FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending January 3, 1998, with historical data — United States

*Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

TABLE I. Summary — provisional cases of selected notifiable diseases, United States, cumulative, week ending January 3, 1998 (53rd Week)

		Cum. 1997		Cum. 1997
St. Lou wester Hansen Disease Hantavirus pulmonary	nia* 1 equine* is* n equine* 1 syndrome* [†] drome, post-diarrheal*	0 76 10 4 1,963 5 120 10 10 12 - 109 18 61 231	Plague Poliomyelitis, paralytic [¶] Psittacosis Rabies, human Rocky Mountain spotted fever (RMSF) Streptococcal disease, invasive Group A Streptococcal toxic-shock syndrome* Syphilis, congenital** Tetanus Toxic-shock syndrome Trichinosis Typhoid fever Yellow fever	4 1 37 2 400 1,431 33 548 43 134 9 346

-:no reported cases *Not notifiable in all states. †Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases (NCID). §Updated monthly to the Division of HIV/AIDS Prevention–Surveillance, and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP), last update December 23, 1997. ¶One suspected case of polio with onset in 1997 has also been reported to date. **Updated from reports to the Division of STD Prevention, NCHSTP.

				Esche coli O	erichia 157:H7			Hepatitis			
	AI	DS	Chla	mydia	NETSS [†]	PHLIS [§]	Gono	rrhea	C/NA,NB		
Reporting Area	Cum. 1997*	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1997	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	
UNITED STATES	56,492	66,213	467,792	437,869	2,316	1,552	289,870	322,818	3,164	3,719	
NEW ENGLAND	2,353	2,746	17,436	16,967	204	127	5,671	6,296	58	113	
Maine N.H.	51 55	49 93	1,067 812	931 752	17 14	16	66 96	55 162	- 8	-7	
Vt.	35	19	434	398	8	3	53	47	3	26	
Mass. R.I.	832 158	1,306 171	7,330 1,857	6,791 1,832	111 10	93	2,077 397	2,163 486	39 8	74 6	
Conn.	1,222	1,108	5,936	6,263	44	15	2,982	3,383	-	-	
MID. ATLANTIC	16,880	18,114	61,423	57,172	144	52	38,337	41,735	380	337	
Upstate N.Y. N.Y. City	2,645 8,941	2,422 9,942	N 31 <i>.</i> 669	N 26,455	99 16	- 8	6,249 14,841	7,606 13,008	293	272 3	
N.J.	3,206	3,584	9,593	12,261	29	25	7,251	8,721	-	-	
Pa.	2,088	2,166	20,161	18,456	N	19	9,996	12,400	87	62	
E.N. CENTRAL Ohio	4,221 802	5,171 1,153	71,549 20,866	83,652 20,653	406 108	281 52	43,813 13,016	57,320 14,946	503 20	490 35	
Ind.	523	591	9,487	10,100	81	46	6,175	6,458	11	8	
III. Mich.	1,841 812	2,192 964	10,941 21,287	22,459 20,277	71 146	31 103	5,322 15,302	15,776 15,267	83 389	93 354	
Wis.	243	271	8,968	10,163	140 N	49	3,998	4,873	- 309	- 504	
W.N. CENTRAL	1,109	1,600	33,107	31,705	519	405	14,936	15,822	165	111	
Minn. Iowa	214 110	304 92	7,502 4,647	5,608 4,165	214 119	206 74	2,782 1,245	2,698 1,144	4 35	10 53	
Mo.	512	851	12,154	12,020	55	69	7,851	8,492	109	23	
N. Dak.	13	12	766	1,022	15	12	55	42	3	-	
S. Dak. Nebr.	11 90	14 93	1,472 2,567	1,536 2,878	29 61	32	177 1,089	176 1,213	3	9	
Kans.	159	234	3,999	4,476	26	12	1,737	2,057	11	16	
S. ATLANTIC	13,628	16,514	92,133	53,331	218	136	89,775	95,569	279	235	
Del. Md.	229 1,875	285 2,246	1,276 7,750	1,148 U	5 26	4 14	1,250 13,129	1,432 11,316	22	1 4	
D.C.	998	1,257	N	Ň	2	-	4,277	4,470	-	-	
Va. W. Va.	1,116 125	1,146 121	11,617 2,883	11,754 2,301	N N	41 1	8,730 928	9,292 850	24 18	17 9	
N.C.	850	898	17,680	Ū	74	38	17,149	18,229	51	46	
S.C. Ga.	800 1,722	842 2,422	12,433 12,957	U 13,333	13 41	9	11,288 14,969	10,984 19,810	38 U	34	
Fla.	5,913	7,297	25,537	24,795	49	29	18,055	19,186	126	124	
E.S. CENTRAL	2,061	2,280	31,736	32,507	96	39	31,883	35,982	336	590	
Ky. Tenn.	360 784	401 821	6,302 12,456	6,687 13,121	30 47	- 39	4,004 10,990	4,229 11,710	16 233	29 400	
Ala.	570	608	8,706	8,352	15	-	12,032	13,334	13	8	
Miss.	347	450	4,272	4,347	4	-	4,857	6,709	74	153	
W.S. CENTRAL Ark.	6,283 242	6,793 267	59,917 2,128	59,215 1,663	79 10	27 5	39,219 3,571	40,942 5,050	489 11	515 8	
La.	1,031	1,464	10,030	7,330	14	12	9,777	7,976	241	292	
Okla. Tex.	294 4,716	261 4,801	7,413 40,346	7,219 43,003	13 42	7 3	4,754 21,117	4,792 23,124	7 230	7 208	
MOUNTAIN	1,799	2,014	23,884	29,642	239	132	8,391	7,459	492	555	
Mont.	41	34	1,121	1,232	24	-	51	35	24	20	
daho Wyo.	52 16	38 7	1,680 635	1,505 621	38 17	23 12	157 54	98 41	86 230	99 179	
Colo.	380	519	1,896	7,282	83	57	2,138	1,367	42	64	
N. Mex. Ariz.	168 405	206 591	3,381 11,529	3,862 10,686	7 N	30	1,137 4,045	891 3,705	60 26	77 76	
Utah	152	194	1,761	1,554	59	-	278	280	5	19	
Nev.	585	425	1,881	2,900	11	10	531	1,042	19	21	
PACIFIC Wash.	8,158 678	10,980 762	76,607 9,551	73,678 9,236	411 130	347 145	17,845 1,962	21,693 2,020	462 31	773 66	
Oreg.	305	461	5,009	5,347	84	95	735	871	4	8	
Calif. Alaska	7,029 52	9,522 36	58,705 1,587	55,926 1,345	185 12	95 3	14,282 381	17,842 459	266	479 3	
Hawaii	94	199	1,755	1,824	N	9	485	459 501	- 161	217	
Guam	2	4	193	355	Ν	-	27	63	-	6	
P.R. V.I.	2,040 99	2,238	U N	U N	41 N	U U	532	648	150	180	
v.i. Amer. Samoa	-	18	-	-	N	U	-	-	-	-	
C.N.M.I.	1	-	N	N	N	U	17	11	2	-	

TABLE II. Provisional cases of selected notifiable diseases, United States,
weeks ending January 3, 1998, and December 28, 1996 (53rd Week)

N: Not notifiable U: Unavailable -: no reported cases C.N.M.I.: Commonwealth of Northern Mariana Islands

*Updated monthly to the Division of HIV/AIDS Prevention–Surveillance, and Epidemiology, National Center for HIV, STD, and TB Prevention, last update December 23, 1997. [†]National Electronic Telecommunications System for Surveillance. [§]Public Health Laboratory Information System.

	Legionellosis		Lyı Dise	me ease	Mal	aria		hilis Secondary)	Tubero	Rabies, Animal	
Reporting Area	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997
UNITED STATES	1,054	1,198	10,979	16,455	1,772	1,800	7,917	11,344	17,158	21,168	7,853
NEW ENGLAND	85	80	3,038	4,095	96	84	132	195	445	485	1,255
Maine N.H.	2 7	5 4	11 39	63 47	1 10	10 4	2	2 1	13 17	21 21	226 48
Vt. Mass.	13 27	5 34	8 387	26 321	2 30	8 32	- 70	- 85	6 254	4 262	113 282
R.I.	16	32	409	534	11	12	2	4	36	39	41
Conn. MID. ATLANTIC	20 227	N 263	2,184 6,421	3,104 10,305	42 442	18 467	58 364	103 509	119 3,180	138 3,969	545 1,719
Upstate N.Y.	74	80	2,557	4,900	72	96	41	76	449	535	1,227
N.Y. City N.J.	13 20	19 15	122 1,639	401 2,190	259 78	269 68	87 122	133 177	1,622 701	2,035 816	U 190
Pa.	120	149	2,103	2,814	33	34	114	123	408	583	302
E.N. CENTRAL Ohio	312 121	360 116	96 61	498 32	148 19	170 15	695 213	1,573 584	1,616 270	2,110 301	177 116
Ind.	54	51	29	32	16	15	166	207	150	202	13
III. Mich.	28 92	38 109	6	10 28	55 43	83 41	71 153	420 183	755 318	1,060 433	20 28
Wis.	17	46	U	396	15	16	92	179	123	114	-
W.N. CENTRAL Minn.	72 3	71 15	233 196	365 251	70 36	51 26	181 23	340 47	594 158	547 131	492 61
lowa Mo.	12 32	11 18	10 20	19 52	10 13	3 11	8 113	23 224	73 238	70 224	158 25
N. Dak.	2	-	-	2	3	1	-	- 224	12	8	84
S. Dak. Nebr.	2 15	3 18	1 2	- 5	3 1	- 3	1 7	- 10	19 22	19 22	74 2
Kans.	6	6	4	36	4	7	29	36	72	73	88
S. ATLANTIC Del.	133 13	197 12	778 105	823 173	363 5	340 4	3,226 22	3,780 35	3,227 18	4,014 43	3,102 67
Md.	28	39	499	447	87	87	920	733	314	319	603
D.C. Va.	5 27	9 54	10 62	3 57	20 68	9 60	117 236	123 393	106 305	139 349	5 669
W. Va. N.C.	N 14	N 12	10 34	12 66	1 21	6 30	3 736	9 1,052	54 464	57 554	89 893
S.C.	8	8	3	9	18	13	360	384	260	350	175
Ga. Fla.	2 35	3 60	7 48	1 55	52 91	38 93	537 295	683 368	595 1,111	790 1,413	325 276
E.S. CENTRAL	53	59	85	83	35	42	1,610	2,412	1,194	1,436	270
Ky. Tenn.	12 33	11 26	15 45	26 24	8 11	12 14	136 747	154 850	184 358	259 504	28 149
Ala. Miss.	4	5 17	11 14	9 24	10 6	8 8	410 317	530 878	416 236	423 250	88 5
W.S. CENTRAL	4 34	53	14	24 175	59	0 158	1,196	1,815	2,449	2,898	341
Ark.	-	1	25	27	5	2	141	262	179	225	55
La. Okla.	8 3	4 16	6 34	9 42	18 8	12 3	366 117	486 177	277 176	350 200	5 112
Tex.	23	32	35	97	28	141	572	890	1,817	2,123	169
MOUNTAIN Mont.	63 1	58 1	23	9	66 2	65 7	192	165	515 17	711 19	190 52
ldaho Wyo.	2 1	- 7	4 5	2 3	1 2	-7	1	4 2	17 2	15 7	- 31
Colo.	18	12	6	-	30	26	14	26	76	104	28
N. Mex. Ariz.	3 12	2 21	1 4	1	8 11	3 9	16 147	8 102	53 272	89 282	12 53
Utah Nev.	19 7	8 7	1 2	1 2	3 9	5 8	5 9	3 20	32 46	58 137	6 8
PACIFIC	, 75	, 57	205	102	493	423	321	555	3,938	4,998	307
Wash. Oreg.	10	8	10 21	18 19	49 24	41 24	17	9	264 154	285 190	14
Calif.	64	43	172	64	406	343	293	533	3,297	4,227	264
Alaska Hawaii	- 1	1 5	2	- 1	5 9	3 12	1 1	- 4	73 150	96 200	29
Guam	-	1	-	-	-	-	3	3	13	107	-
P.R. V.I.	-	- 1	-	-	6	2 1	240	209	212	222	66
Amer. Samoa	-	-	-	-	-	-	-	-	-	-	-
C.N.M.I.	-	-	-	-	-	-	9	1	2	-	-

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending January 3, 1998, and December 28, 1996 (53rd Week)

N: Not notifiable U: Unavailable -: no reported cases

	H. influ	ienzae,	н	epatitis (Vi	ral), by typ	De	Measles (Rubeola)					
	-	sive		A	E	-	Indi	genous	lmp	orted [†]	-	tal
Reporting Area	Cum. 1997*	Cum. 1996	Cum. 1997	Cum. 1996	Cum. 1997	Cum. 1996	1997	Cum. 1997	1997	Cum. 1997	Cum. 1997	Cum. 1996
UNITED STATES	1,056	1,170	27,799	31,032	8,749	10,637	-	78	2	57	135	508
NEW ENGLAND	65	55	632	456	153	255	-	11	-	8	19	17
Maine N.H.	5 12	1 13	66 35	28 22	6 18	8 21	-	- 1	-	1 -	1 1	-
Vt. Mass.	3 40	2 36	15 241	12 229	10 56	14 111	-	10	-	- 6	- 16	2 12
R.I. Conn.	3	2	130 145	26 139	18 45	19 82	-	-	-	- 1	1	1
MID. ATLANTIC	147	213	1,917	1,985	45 1,315	1,413	-	19	-	8	27	38
Upstate N.Y. N.Y. City	41 35	50 57	373 705	438 609	331 434	358 491	-	2 9	-	3	5 11	12 11
N.J.	51	65	287	394	222	279	-	3	-	-	3	3
Pa. E.N. CENTRAL	20 161	41 191	552 2,858	544 2,619	328 916	285 1,103	-	5 6	-	3 3	8 9	12 21
Ohio	86	95	332	785	94	120	-	-	-	-	-	6
lnd. III.	19 38	21 50	322 706	367 763	93 227	143 335	-	6	-	- 1	-7	3
Mich. Wis.	15 3	12 13	1,342 156	506 198	456 46	416 89	-	-	-	2	2	3 9
W.N. CENTRAL	65	63	2,201	2,656	472	572	-	13	-	5	18	24
Minn. Iowa	44 7	48 4	196 492	176 334	44 47	94 74	-	3 1	-	5	8 1	19 1
Mo. N. Dak.	10	8	1,109 11	1,414 140	326 5	326 2	-	1	-	-	1	3
S. Dak.	2	1	27	43	1	5	-	8	-	-	8	-
Nebr. Kans.	1 1	1 1	109 257	156 393	16 33	39 32	-	-	-	-	-	- 1
S. ATLANTIC	174	273	2,154	1,960	1,302	1,573	-	2	2	15	17	12
Del. Md.	- 58	2 76	31 215	21 256	6 195	9 169	-	-	-	2	2	1 2
D.C. Va.	- 13	5 11	36 233	39 218	30 128	32 163	-	-	2	3 1	3 1	- 3
W. Va. N.C.	4 21	11 26	12 211	19 204	16 265	36 337	-	-	-	2	2	2
S.C.	4	5	110	57	98	101	-	-	-	1	1	-
Ga. Fla.	42 32	52 85	657 649	414 732	148 416	61 665	-	2	-	1 5	1 7	3 1
E.S. CENTRAL	48	45	637	1,273	692	914	-	-	-	-	-	2
Ky. Tenn.	6 27	6 25	76 402	53 778	40 449	76 516	-	-	-	-	-	2
Ala. Miss.	15	13 1	90 69	217 225	80 123	78 244	Ū	-	Ū	-	-	-
W.S. CENTRAL	54	44	5,515	6,807	1,188	1,616	-	3	-	5	8	27
Ark. La.	1 14	- 6	221 239	500 261	63 171	93 209	-	-	-	-	-	- 1
Okla. Tex.	34 5	32 6	1,424 3,631	2,586 3,460	52 902	56 1,258	- U	- 3	Ū	1 4	1 7	26
MOUNTAIN	99	57	4,369	3,400 4,573	898	1,258	-	6	-	2	8	157
Mont.	- 1	1 1	72 150	130	12 54	21 88	-	-	-	-	-	- 1
Wyo.	4	-	41	247 41	40	45	-	-	-	-	-	1
Colo. N. Mex.	21 10	16 11	407 359	512 355	154 263	132 417	-	-	-	-	-	7 17
Ariz. Utah	36 3	20 8	2,376 547	1,767 1,073	206 92	237 129	-	5	-	- 1	5 1	8 118
Nev.	24	-	417	448	77	95	U	1	U	1	2	5
PACIFIC Wash.	243 6	229 10	7,516 673	8,703 1,001	1,813 80	2,027 158	-	18 1	-	11 1	29 2	210 38
Oreg.	35	33	378	875	109	129	-	_ 15	-	-	-	14 46
Calif. Alaska	188 7	178 6	6,296 34	6,653 54	1,592 21	1,710 16	-	-	-	8	23	63
Hawaii Guam	7	2	135	120 7	11 3	14 1	-	2	-	2	4	49
P.R.	-	2	257	292	3 1,376	1,195	U	-	U 	-	-	3
V.I. Amer. Samoa	-	-	-	41	-	44	U U	-	U U	-	-	-
C.N.M.I.	6	10	1	1	34	5	Ŭ	1	Ű	-	1	-

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination,
United States, weeks ending January 3, 1998,
and December 28, 1996 (53rd Week)

N: Not notifiable U: Unavailable -: no reported cases

 * Of 242 cases among children aged <5 years, serotype was reported for 126 and of those, 47 were type b.

[†]For imported measles, cases include only those resulting from importation from other countries.

		ococcal ease		Mumps			Pertussis		Rubella			
Reporting Area	Cum. 1997	Cum. 1996	1997	Cum. 1997	Cum. 1996	1997	Cum. 1997	Cum. 1996	1997	Cum. 1997	Cum. 1996	
UNITED STATES	3,117	3,437	4	612	751	59	5,519	7,796	-	161	238	
NEW ENGLAND	205	171	-	12	5	10	990	1,866	-	2	27	
Maine N.H.	18	15	-	- 1	- 1	-	11 143	55 197	-	-	-	
Vt.	18 4	13 4	-	-	1	10	262	280	-	-	2	
Mass.	102	71	-	4	1 1	-	522	1,245	-	1	21	
R.I. Conn.	21 42	18 50	-	6 1	1	-	17 35	40 49	-	- 1	- 4	
MID. ATLANTIC	325	381	-	59	96	-	409	952	-	32	13	
Upstate N.Y.	75 46	102 56	-	13 3	28 20	-	161 62	533 61	-	5 27	5 5	
N.Y. City N.J.	73	79	-	3 7	20	-	11	31	-	- 27	2	
Pa.	131	144	-	36	44	-	175	327	-	-	1	
E.N. CENTRAL	461	475	-	80 25	135	3	544	837	-	5	3	
Ohio Ind.	164 58	159 64	-	35 14	52 8	1	165 85	289 128	-	-	-	
III.	148	142	-	13	24	2	126	192	-	2	1	
Mich. Wis.	53 38	51 59	-	15 3	48 3	-	62 106	59 169	-	- 3	2	
W.N. CENTRAL	235	264	-	18	24	17	599	573	-	2	-	
Minn.	34	39	-	6	7	15	384	433	-	-	-	
lowa Mo.	48 109	56 98	-	10	3 10	2	113 68	32 74	-	- 2	-	
N. Dak.	2	5	-	-	2	-	2	1	-	-	-	
S. Dak. Nebr.	6 14	10 29	-	2	-	-	5 14	4 15	-	-	-	
Kans.	22	27	-	-	2	-	13	14	-	-	-	
S. ATLANTIC	558	659	-	85	131	-	438	793	-	83	101	
Del. Md.	5 42	3 58	-	- 10	37	-	1 125	26 278	-	-	-	
D.C.	9	5	-	-	-	-	3	4	-	1	1	
Va. W. Va.	58 18	67 18	-	19	19	-	56 6	108 7	-	1	2	
N.C.	97	79	-	12	27	-	118	, 186	-	59	86	
S.C. Ga.	61 106	65 147	-	11 10	7 9	-	30 14	49 35	-	19	1	
Fla.	162	217	-	23	32	-	85	100	-	3	11	
E.S. CENTRAL	237	246	1	28	23	3	144	202	-	-	2	
Ky. Tenn.	48 85	31 65	- 1	3 7	- 1	- 1	61 40	142 24	-	-	-	
Ala.	85	95	-	9	6	2	35	26	-	-	2	
Miss.	19	55	U	9	16	U	8	10	U	-	N	
W.S. CENTRAL Ark.	282 34	365 35	-	75 1	67 1	1	296 60	201 14	-	4	9	
La.	48	66	-	16	21	-	20	15	-	-	1	
Okla. Tex.	45 155	46 218	Ū	- 58	1 44	1 U	49 167	21 151	Ū	- 4	- 8	
MOUNTAIN	190	183	2	58 59	25	25	1,252	660	0	4 6	9	
Mont.	9	9	-	-	- 25	-	19	37	-	-	-	
Idaho Wyo.	15 4	25 4	1	6 1	- 1	14	570 7	115 8	-	1	2	
Colo.	51	44	-	3	5	2	348	336	-	-	3	
N. Mex. Ariz.	30 46	27 37	N 1	N 34	N 1	5 4	198 45	64 33	-	- 5	- 3	
Utah	16	18	-	8	3	-	45 26	33 26	-	5	-	
Nev.	19	19	U	7	15	U	39	41	U	-	1	
PACIFIC	624 92	693 116	1	196 21	245	-	847 406	1,712 830	-	27 5	74 15	
Wash. Oreg.	126	123	N	21 N	26 N	-	406	64	-	-	1	
Calif.	396	437	-	147	185	-	403	780	-	14	55	
Alaska Hawaii	3 7	9 8	- 1	4 24	3 31	-	14 14	3 35	-	- 8	- 3	
Guam	1	5	U	1	10	U	-	-	U	-	-	
P.R.	10	13	-	7	2	-	2	3	-	-	-	
V.I. Amer. Samoa	-	-	U U	-	2	U U	-	-	U U	-	-	
C.N.M.I.	-	-	Ū	4	-	Ū	-	-	Ū	-	-	

TABLE III. (Cont'd.) Provisional cases of selected notifiable diseases preventable
by vaccination, United States, weeks ending January 3, 1998,
and December 28, 1996 (53rd Week)

N: Not notifiable U: Unavailable -: no reported cases

All Causes, By Age (Years)				P&I [†]	I [†]		All Causes, By Age (Years)								
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	P&l [†] Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn. Cambridge, Mass. Fall River, Mass. Hartford, Conn. Lowell, Mass. New Bedford, Mass. New Bedford, Mass. New Haven, Conn. Providence, R.I. Somerville, Mass. Springfield, Mass. Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J.	619 169 255 57 31 200 34 68 9 44 41 81 2,582 25 79 35	452 112 U 14 20 366 26 12 18 53 53 53 53 53 53 53 53 53 53 53 53 53	26 U 1 3 11 5 4 4 6 9 2 4 6 15 449 8 3 10 11	44 20 U 1 7 - 3 2 3 2 - 3 2 - 3 2 2 216 2 5 6	11 1 U 1 2 2 2 1 1 60 2 2 1 4	16 10 - - 1 - 1 2 - 1 - 1 38 3 - 2 1	47 14 U 1 2 4 - 1 5 15 133 3 1 8 8	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla. Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, Fla. Tampa, Fla. Washington, D.C. Wilmington, Del. E.S. CENTRAL Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn.	U 120 27 600 118	525 92 67 96 52 U 53 316 40 75 13 410 711 16 51 38 726	141 U 33 21 17 21 U 13 6 8 U 22 - 120 24 10 14 125 21	99 U 19 5 12 17 U 10 4 5 U 13 14 40 14 5 5 5 8	30 0 6 2 5 5 5 0 6 1 - U 5 - 18 6 1 2 - 1 7	22 U 4 4 4 1 U 4 - U 5 - 10 1 1 - 3 3 1	49 15 9 9 1 U 6 3 3 U 3 - 52 3 2 8 5 1 1
Elizabeth, N.J. Erie, Pa. Jersey City, N.J. New York City, N.Y. Newark, N.J. Paterson, N.J. Philadelphia, Pa. Pittsburgh, Pa.§ Reading, Pa. Rochester, N.Y. Schenectady, N.Y. Scranton, Pa. Syracuse, N.Y. Trenton, N.J. Utica, N.Y. Yonkers, N.Y. E.N. CENTRAL	34 29 54	20 21 37 995 28 10 282 33 30 94 26 21 46 23 10 U 1,394	5 6 8 246 3 70 6 3 23 7 3 10 7 3 U	6 1 5 121 13 4 35 2 5 6 2 - 1 2 - U 110	1 31 5 1 8 3 - 1 - 1 - U 37	-3 -4 15 11 2 2 	2 3 61 3 19 2 4 11 1 2 4 U 111	Montgomery, Ala. Nashville, Tenn. W.S. CENTRAL Austin, Tex. Baton Rouge, La. Corpus Christi, Tex. Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La. San Antonio, Tex. Shreveport, La. Tulsa, Okla.	1 129 1,260 65 26 27 163 61 79 293 71 121 184 46 124 664	1 95 828 44 15 22 105 38 51 192 49 60 127 30 95 479	24 252 9 7 36 10 11 63 16 30 40 10 17 110	8 113 7 4 11 10 7 29 3 16 13 2 10 44	1 42 1 6 2 4 6 1 14 3 3 2 12	1 24 5 - 5 6 3 2 1 1 1 1 1 9	12 67 2 1 3 7 20 3 - 13 27 58
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, Mich Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Moa. Lincoln, Nebr.	60 48 418 63 119 133 88 230 46 49 5	44 41 276 49 86 94 68 94 68 149 42 38 104 25 81 343 573 45 60 60 573 108 26 47 26	$\begin{array}{c} 12\\ 5\\ 9\\ 9\\ 21\\ 49\\ 2\\ 8\\ 2\\ 5\\ 32\\ 10\\ 19\\ 10\\ 10\\ 121\\ 10\\ 8\\ 8\\ 10\\ \end{array}$	3 1 3 5 5 8 4 19 2 1 1 4 6 1 4 - 3 4 7 1 8 8 6 - 4 3 2	1 - 9 - 2 1 2 8 - 2 - 2 5 1 1 - 1 - 2 - 20 5 - 1 2 1 2 1	- 1 1 8 - 5 	. 1 1 2 8 1 0 1 1 2 3 2 5 2 7 3 4 5 7 3 5 1 2 4 2 8 4 5 7 3 5 1 2 4 2 8 4	Albuquerque, N.M. Boise, Idaho Colo. Springs, Colo Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, Utah Tucson, Ariz. PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawaii Long Beach, Calif. Pasadena, Calif. Pasadena, Calif. Pasadena, Calif. Pasadena, Calif. San Diego, Calif. San Diego, Calif. San Jose, Calif. San Jose, Calif. San Jose, Calif. Santa Cruz, Calif. Seattle, Wash.	80 122 14 89 25 95 128 1,151 25 0 4 71 83 214 29 83 214 29 83 U 214	59 25 0 63 73 10 61 20 73 95 826 19 0 4 4 866 160 152 0 84 102 113 19 59	10 6 U 9 38 1 11 4 7 24 205 4 U - 17 10 41 10 26 29 3 3 3 8 3 8	63U48141077-U-346410U8475124	1 1 U - 3 1 3 - 2 1 23 - U - 3 1 4 - 3 U 5 2 1 2 2 -	U 4 - 10 - 31 20 2 U - 2 3 - U 5 2 5 - 1 - 1	3 1 U 10 5 2 6 1 15 10 2 U 1 2 15 1 7 2 U 6 1 3 4 6 10
Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.		26 81 51 102 53 66	14 6 15 10	2 5 4 9 1 4	2 6 3	1 2 5 2	4 2 4 10 2	Tacoma, Wash.	′Ü 10,396 [¶]	U	U	Ü 781	U 253	U 200	Ŭ 676

TABLE IV. Deaths in 122 U.S. cities,* week ending January 3, 1998 (53rd 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 or more. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included. *Pneumonia and influenza. *Because of changes in reporting methods in 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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MMWR

HIV Case Surveillance — Continued

CDC has recommended that all states and territories conduct HIV case surveillance as an extension of their AIDS surveillance systems (1). In addition, CDC is developing technical guidance to enhance security practices, standardize confidentiality laws and regulations, and promote uniform standards for HIV case surveillance systems. These guidelines will assist states and territories in implementing HIV case surveillance using data-collection and data-storage methods that provide high quality HIV surveillance data while assuring the confidentiality of surveillance information.

References

- 1. CDC. Update: trends in AIDS incidence, 1996. MMWR 1997;46:861-7.
- Carpenter CC, Fischl MA, Hammer SM, et al. Antiretroviral therapy for HIV infection in 1997: updated recommendations of the International AIDS Society-USA panel. JAMA 1997;277:1962– 9.
- Council of State and Territorial Epidemiologists. CSTE: position statement ID-4. National HIV surveillance: addition to the National Public Health Surveillance System. Atlanta, Georgia: Council of State and Territorial Epidemiologists, 1997.
- Klaucke DN. Evaluating public health surveillance. In: Teutsch SM, Churchill RE, eds. Principals and practices of public health surveillance. New York, New York: Oxford University Press, 1994:158–74.
- 5. Rosenblum L, Buehler JW, Morgan ME, et al. The completeness of AIDS case reporting, 1988: a multisite collaborative surveillance project. Am J Public Health 1992;82:1495–9.
- 6. CDC. HIV/AIDS surveillance report. Atlanta, Georgia: US Department of Health and Human Services, Public Health Service, 1997. (Vol 8, no. 1).
- 7. Meyer PA, Jones JL, Garrison CZ. Completeness of reporting of diagnosed HIV-infected hospital inpatients. J AIDS 1994;7:1067–73.
- 8. Gostin LO, Lazzarini Z, Neslund VS, Osterholm MT. The public health information infrastructure: a national review of the law on health information privacy. JAMA 1996;275:1921–7.

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The Morbidity and Mortality Weekly Report (MMWR) Series is prepared by the Centers for Disease Control and Prevention (CDC) and is available free of charge in electronic format and on a paid subscription basis for paper copy. To receive an electronic copy on Friday of each week, send an e-mail message to *listserv@listserv.cdc.gov*. The body content should read SUBscribe mmwr-toc. Electronic copy also is available from CDC's World-Wide Web server at http://www.cdc.gov/ or from CDC's file transfer protocol server at ftp.cdc.gov. To subscribe for paper copy, contact Superintendent of Documents, U.S. Government Printing Office, Washington, DC 20402; telephone (202) 512-1800.

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