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## Bronchoscopy-Related Infections and Pseudoinfections — New York, 1996 and 1998

Bronchoscopy is a useful diagnostic technique that can be performed safely by trained specialists when the bronchoscopes in both inpatient and ambulatory-care settings are reprocessed properly to prevent transmission of infection. The New York State Department of Health received reports of three clusters of culture-positive bronchoscopy specimens obtained in 1996 and 1998 from patients at local health-care facilities. This report summarizes the results of investigations of these clusters, which indicated involvement of *Mycobacterium tuberculosis, M. intracellulare,* or imipenemresistant *Pseudomonas aeruginosa*. Between patient uses, bronchoscopes had been cleaned, visually inspected, leak tested, and processed by STERIS System 1 processors (STERIS, Mentor, Ohio)\*.

#### Cluster 1

During November–December 1996, bronchial specimens from five patients at a health-care facility yielded *M. tuberculosis* with the same restriction fragment length polymorphism (RFLP) pattern suggesting a common source. The index case-patient had tuberculosis with persistent acid-fast bacillus (AFB) smear- and culture-positive specimens. The four subsequent case-patients had no clinical evidence of tuberculosis, although one had a positive tuberculin skin test 6 weeks postbronchoscopy and was treated with isoniazid. Investigators concluded that all specimens from the four patients were contaminated but could not determine whether contamination occurred during the bronchoscopy or in the mycobacteriology laboratory. Specimens from three of the four case-patients were processed in the laboratory on the same day as the index case-patient's specimen.

The bronchoscopies were performed using three Olympus BF-P20D (Olympus America, Inc., Melville, New York) bronchoscopes, each processed in the same STERIS System 1 processor. Cultures from all three bronchoscopes, taken 5 weeks after the last case procedure, were negative. The same cleaning brushes used on all three bronchoscopes also were culture negative. Investigators identified an inconsistency between the disinfection/sterilization procedures recommended in the STERIS manual

<sup>\*</sup>Use of trade names and commercial sources is for identification only and does not imply endorsement by CDC or the U.S. Department of Health and Human Services.

Bronchoscopy-Related Infections — Continued

and those followed by the facility personnel—the biopsy port cap was not replaced before loading for cleaning in the STERIS System 1 processor. The bronchoscope manufacturer did not provide recommendations for processing in the STERIS System 1, but the manual suggests removal of the biopsy port cap before cleaning and replacing it immediately before the next use. At the investigators' request, the STERIS device testing program performed pressure and flow studies with the biopsy port cap removed and observed a 50% flow reduction and a 25% flow pressure reduction. Therefore, STERIS could not assure bronchoscope sterility when the biopsy port cap was not replaced before processing, as specified in the STERIS manual.

#### Cluster 2

During March–April 1998, an increase in positive bronchial specimens for *M. avium-intracellulare* (MAI) occurred among patients in an ambulatory surgery unit (ASU) at a health-care facility. Seven cases without clinical evidence of MAI were identified over a 2-month period compared with two MAI cases during the preceding 8 months. All seven patients had undergone bronchoscopy in the same ASU with the same bronchoscope. Typing by polymerase chain reaction restriction enzyme analysis indicated that all of the isolates from the ASU bronchoscopy-associated patients were *M. intracellulare* (nontypable), and all of the isolates from the environmental and control patients with previously diagnosed atypical mycobacterial disease were *M. avium*. Mycobacterial cultures of the implicated bronchoscope, taken 12 days after diagnosis of the last MAI case, were negative.

The bronchoscope used was an Olympus BF-P20D model and was processed in a STERIS System 1. Olympus connectors were used for processing the bronchoscope in the STERIS System 1 rather than the connector kit and methods specifically developed by STERIS.

#### Cluster 3

During August–October 1998, 18 patients (11 inpatients and seven outpatients) at a health-care facility had bronchial specimens that grew imipenem-resistant *P. aeruginosa* (IRPA). None of the 18 patients had IRPA isolated from sputum cultures obtained before bronchoscopy. At least three patients had persistent infection with IRPA with an associated clinical illness postbronchoscopy. All but one of the isolates from the 18 patients had identical DNA patterns by pulsed-field gel electrophoresis analysis.

In July 1998, the facility began processing bronchoscopes and other endoscopes using a STERIS System 1 processor. The facility used Pentax (Pentax, Orangeburg, New York) and Olympus bronchoscopes but did not document the specific bronchoscope used on each patient. Neither the Pentax nor the Olympus bronchoscopes were connected to the STERIS System 1 in accordance with the STERIS manufacturer's recommendations. The person responsible for cleaning and disinfecting the endoscopes had received training at the STERIS Corporation; however, the specific scopes used at the facility were not demonstrated during the training.

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Bronchoscopy-Related Infections — Continued

**Editorial Note**: The number of bronchoscopy procedures performed in the United States reached an estimated 497,000 in 1996 (1). Although reported infectious complications caused by bronchoscopy are rare (2), the incidence is probably underestimated, with many episodes unrecognized or unreported. Most reported bronchoscopy-related outbreaks or pseudo-outbreaks have been associated with inadequate cleaning and disinfection procedures (3–9).

The findings in this report identified additional problems related to using automated reprocessing machines. Conflicting recommendations for disinfection/sterilization exist between bronchoscope and reprocessor system manufacturers. Some individual bronchoscope models are not compatible with certain automated reprocessing systems. However, users may not be aware of these incompatibilities unless they make a device-specific inquiry to the manufacturers. Personnel using automated reprocessing machines in these clusters did not receive adequate device-specific training, and the wrong set up or connector systems were used. Inadequate documentation in the third cluster about which bronchoscope was used in which patient prevented traceback of the culture-positive respiratory specimens to a particular bronchoscope.

Bronchoscopes are designed with small lumens, multiple ports with obtuse angles, and linings vulnerable to damage and subsequent biofilm formation, presenting obstacles to proper cleaning and disinfection or sterilization. Manual cleaning and sterilization with chemical agents, such as glutaraldehyde, is the reprocessing method most widely recommended by bronchoscopy equipment manufacturers; however, this process is laborious, time consuming, and poses a chemical contact risk to health-care workers. Thus, many health-care facilities use automated reprocessing machines. These machines can become colonized and cause bronchoscopy-related outbreaks or pseudo-outbreaks (5–8).

To address the challenges of reprocessing bronchoscopes, all users should comply with guidelines for cleaning and disinfection/sterilization (2,10). The following additional steps should be taken to reduce bronchoscopy-related infections or pseudoinfections. First, bronchoscope users should obtain and review model-specific reprocessing protocols from both bronchoscope and automated reprocessing system manufacturers. Second, bronchoscope and reprocessor system manufacturers should collaborate to develop and validate device- and model-specific high-level disinfection or sterilization protocols. Third, user education should include on-site training and observation during the set up of each bronchoscope model to clarify device- and modelspecific differences in procedure. Fourth, instruction manuals provided by both bronchoscopy equipment and automated reprocessing system manufacturers should address procedural differences among varying models of bronchoscopes and highlight proper connector system(s) to be used with their machine. Fifth, connector systems should be clearly labeled (e.g., color coded) to ensure proper selection and use. Finally, quality-control procedures should be developed in each health-care facility to include visual inspection of the bronchoscope, regular testing for bronchoscope integrity, maintenance, and surveillance for unusual clusters of organisms.

Under the Safe Medical Devices Act of 1990, facilities are required to report to the Food and Drug Administration (FDA) instances when endoscopes (including bronchoscopes) and endoscope reprocessing systems may have caused or contributed to serious injury or a patient's death. Questions concerning this mandatory reporting

Bronchoscopy-Related Infections — Continued

requirement can be directed to FDA's Center for Devices and Radiological Health, Office of Surveillance and Biometrics, telephone (310) 827-0360. In addition, health-care workers are requested to report bronchoscopy-related colonization episodes, infection, or pseudoinfection to their state health department, to FDA's MedWatch program, telephone (800) 332-1088, fax (800) 332-0178, or World-Wide Web site, http://www.fda.gov/medwatch, and to CDC's Hospital Infections Program, telephone (404) 639-6413 or fax (404) 639-6459.

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### Rubella Outbreak — Westchester County, New York, 1997–1998

Since licensure of rubella vaccines in 1969, the incidence of rubella and congenital rubella syndrome (CRS) in the United States has decreased substantially. Rubella infection during the first trimester of pregnancy can result in miscarriage, stillbirth, or infants with a pattern of birth defects (i.e., CRS) (1). One of the national health objectives for 2000 is to eliminate indigenous rubella and CRS (objective 20.1) (2). During 1997–1998, 524 cases of rubella were reported in the United States (CDC, unpublished data, 1999). This report describes a rubella outbreak in Westchester County, New York, demonstrates the importance of accurately defining and vaccinating at-risk populations to prevent transmission, and underscores how collaboration with community-based organizations can facilitate the development and implementation of control measures.

During the outbreak, a clinical case of rubella was defined as an illness with an acute onset of generalized maculopapular rash, a temperature of >99 F (>37.2 C), and arthralgia/arthritis, lymphadenopathy, or conjunctivitis. Laboratory confirmation of rubella required a positive serologic test for rubella IgM antibody, a substantial increase

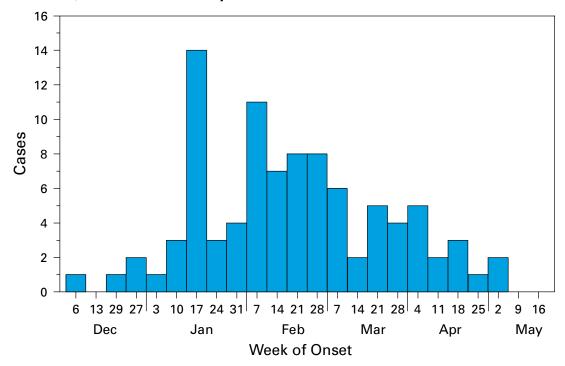
#### Rubella Outbreak — Continued

in acute- and convalescent-phase titers in serum rubella IgG antibody levels by any standard serologic assay, or isolation of rubella virus (3). A confirmed case of rubella required either laboratory confirmation or meeting the clinical case definition and epidemiologic linkage to a laboratory-confirmed case.

From December 1997 through May 1998, 95 confirmed rubella cases were identified in Westchester County (attack rate: 10.7 per 100,000 population); 79 (83%) were laboratory-confirmed and 16 (17%) were linked epidemiologically to a laboratory-confirmed case. During this period, 333 cases were reported in the United States. The outbreak peaked during mid-January and mid-February (Figure 1). The index case-patient in Westchester County was a 23-year-old man from Mexico who first noticed a rash on December 6, 1997. He was exposed previously to a Hispanic co-worker with rubella in Port Chester, New York, who resided in Connecticut, where there was an ongoing rubella outbreak. Port Chester reported 53 (50%) cases; cases were identified in 14 towns, cities, or villages. The outbreak spread through the county along train lines and through work sites.

The median age of case-patients was 23 years (range: 4 months–59 years); 76% were males aged 16–54 years. Of the 22 female patients, 19 were of childbearing age (15–44 years). Of five (26%) pregnant women, three were infected during the first trimester and elected to terminate their pregnancies. The other pregnant women delivered infants with no CRS. Eighty-eight (93%) patients were foreign born; the median time in the United States was 4 years (range: 12 days–26 years). Among foreign-born patients, 34 (39%) were born in Mexico and 31 (35%) in Guatemala. The remaining 23 (27%) patients were born in Colombia, Dominican Republic, El Salvador, Ecuador,

FIGURE 1. Confirmed cases of rubella,\* by week of rash onset — Westchester County, New York, December 1997–May 1998



<sup>\*</sup>n=93. Two patients did not have a rash.

Rubella Outbreak — Continued

Nicaragua, or Portugal. None of the patients born outside the United States had received rubella vaccine. Of the seven U.S.-born patients, four were aged ≥29 years with no history of rubella vaccination, and three were aged <1 year and had parents who were born in Latin American countries.

Local health authorities initiated control measures including case and contact investigations, vaccination of contacts and susceptible persons in the community, and increased awareness to screen pregnant women for susceptibility to rubella and asymptomatic infection. Active surveillance for rash illness was conducted at 28 sites in the county, including emergency departments, health departments, and private providers. Health alerts in Spanish and English were sent to all schools and physicians and distributed in Hispanic communities. Although rubella vaccine was available at no cost at the county health department, special clinics, and work sites, only 248 doses were administered during December 6, 1997–February 9, 1998.

To facilitate rubella-control efforts, health department staff identified community leaders and formed partnerships between Hispanic community-based organizations and Hispanic outreach workers from the Westchester County Health Department. These community-based organizations collaborated with the health department to provide targeted educational materials and one-on-one counseling about the importance of rubella vaccination and bilingual personnel for vaccination sites.

The number of sites offering measles, mumps, and rubella (MMR) vaccine was increased by the health department at work sites (e.g., restaurants, landscaping companies, and cleaning services), special vaccination clinics (e.g., churches, day labor pick-up sites, and a mobile van), and at district public health clinics. The number of vaccinations administered increased, and by the end of May 1998, 4539 doses of MMR vaccine had been administered. The last case of rubella associated with the outbreak was identified on May 2, 1998.

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Editorial Note: The rubella outbreak in Westchester County occurred among young Hispanic adults who were born in countries either without national rubella vaccination programs or where such programs were implemented recently. The demographic characteristics of case-patients were similar to those reported in other recent rubella outbreaks in the United States (4). Most cases occurred among unvaccinated persons aged ≥20 years and among persons who were foreign born, primarily Hispanics (63% of reported cases in 1997) (CDC, unpublished data, 1998). Previous community outbreaks were localized in close-knit, circumscribed, Hispanic neighborhoods (CDC, unpublished data, 1997). The Westchester County outbreak differed in that it did not remain localized, but spread to 14 towns, cities, and villages and occurred among eight different Hispanic nationalities. The wide distribution of cases and the multiple Hispanic nationalities made it difficult to identify and access the at-risk population for targeted control measures. Factors that may have contributed to the low receipt of rubella vaccine included difficulty identifying who the leaders were in the Hispanic communities, limited demographic information about the Hispanic communities, and the Hispanic communities' distrust of persons affiliated with the government because of immigration concerns.

#### Rubella Outbreak — Continued

In outbreaks of rubella in foreign-born populations, both prevention and control measures require a culturally sensitive approach. Collaboration between health departments and community-based organizations may be useful in effectively informing and mobilizing the at-risk population.

In recent years, rubella vaccination programs have been introduced throughout the Americas to decrease the morbidity and mortality from rubella infections during pregnancy. However, because these programs were only recently implemented, persons who have entered the United States as adults probably are not vaccinated and may be susceptible to rubella. Further decreases in rubella incidence in the United States will require increased vaccine coverage in susceptible populations.

During rubella outbreaks, vaccination is the most effective preventive measure. In the United States, two doses of MMR vaccine are recommended at age 12–15 months and 4–6 years (5). For adults who have not received rubella vaccine, a single dose of a rubella-containing vaccine is considered evidence of immunity (6). Reduction in rubella morbidity in Latin America is expected to lower the number of cases imported from this area and indigenous outbreaks in the United States.

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### Notice to Readers

# Thimerosal in Vaccines: A Joint Statement of the American Academy of Pediatrics and the Public Health Service

The Food and Drug Administration (FDA) Modernization Act of 1997 called for FDA to review and assess the risk of all mercury-containing food and drugs. In line with this review, U.S. vaccine manufacturers responded to a December 1998 and April 1999 FDA request to provide more detailed information about the thimerosal content of their preparations that include this compound as a preservative. Thimerosal has been used as an additive to biologics and vaccines since the 1930s because it is very effective in killing bacteria used in several vaccines and in preventing bacterial contamination, particularly in opened multidose containers. Some but not all of the vaccines recommended routinely for children in the United States contain thimerosal.

Thimerosal in Vaccines — Continued

There is a significant safety margin incorporated into all the acceptable mercury exposure limits. Furthermore, there are no data or evidence of any harm caused by the level of exposure that some children may have encountered in following the existing immunization schedule. Infants and children who have received thimerosal-containing vaccines do not need to be tested for mercury exposure.

The recognition that some children could be exposed to a cumulative level of mercury over the first 6 months of life that exceeds one of the federal guidelines on methyl mercury now requires a weighing of two different types of risks when vaccinating infants. On the one hand, there is the known serious risk of diseases and deaths caused by failure to immunize our infants against vaccine-preventable infectious diseases; on the other, there is the unknown and probably much smaller risk, if any, of neurodevelopmental effects posed by exposure to thimerosal. The large risks of not vaccinating children far outweigh the unknown and probably much smaller risk, if any, of cumulative exposure to thimerosal-containing vaccines over the first 6 months of life.

Nevertheless, because any potential risk is of concern, the Public Health Service (PHS), the American Academy of Pediatrics (AAP), and vaccine manufacturers agree that thimerosal-containing vaccines should be removed as soon as possible. Similar conclusions were reached this year in a meeting attended by European regulatory agencies, European vaccine manufacturers, and FDA, which examined the use of thimerosal-containing vaccines produced or sold in European countries.

PHS and AAP are working collaboratively to assure that the replacement of thimerosal-containing vaccines takes place as expeditiously as possible while at the same time ensuring that our high vaccination coverage levels and their associated low disease levels throughout our entire childhood population are maintained.

The key actions being taken are

- A formal request to manufacturers for a clear commitment and a plan to eliminate or reduce as expeditiously as possible the mercury content of their vaccines.
- 2. A review of pertinent data in a public workshop.
- 3. Expedited FDA review of manufacturers' supplements to their product license applications to eliminate or reduce the mercury content of a vaccine.
- 4. Provide information to clinicians and public health professionals to enable them to communicate effectively with parents and consumer groups.
- 5. Monitoring immunization practices, future immunization coverage, and vaccinepreventable disease levels.
- 6. Studies to better understand the risks and benefits of this safety assessment.

PHS and AAP continue to recommend that all children should be immunized against the diseases indicated in the recommended immunization schedule. Given that the risks of not vaccinating children far outweigh the unknown and much smaller risk, if any, of exposure to thimerosal-containing vaccines over the first 6 months of life, clinicians and parents are encouraged to immunize all infants even if the choice of individual vaccine products is limited for any reason.

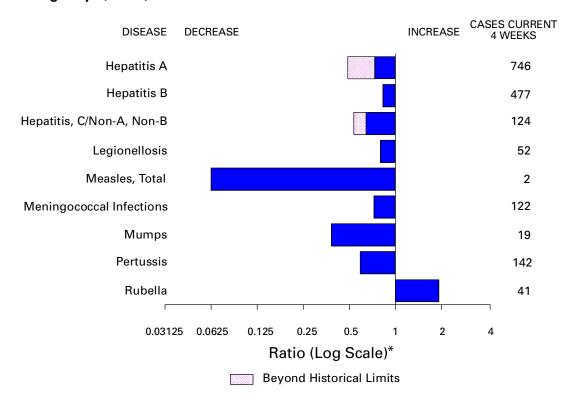
While there is a margin of safety with existing vaccines containing thimerosal, there are steps that can be taken to increase that margin even further. Clinicians and parents can take advantage of the flexibility within the existing schedule for infants born to hepatitis B surface antigen (HBsAg)-negative women to postpone the first

Thimerosal in Vaccines — Continued

dose of hepatitis B vaccine from birth until 2 to 6 months of age when the infant is considerably larger. Preterm infants born to HBsAg-negative mothers should similarly receive hepatitis B vaccine, but ideally not until they reach term gestational age and a weight of at least 5.5 lbs (2.5 kg). Because of the substantial risk of disease, there is no change in the recommendations for infants of HBsAg-positive mothers or of mothers whose status is not known. Also, in populations where HBsAg screening of pregnant women is not routinely performed, vaccination of all infants at birth should be maintained, as is currently recommended. In addition to the key actions mentioned above, the PHS Advisory Committee on Immunization Practices and the AAP Committee on Infectious Diseases will be reviewing these issues and may make additional statements.

Reported by: Public Health Service, US Dept of Health and Human Services. American Academy of Pediatrics, Elk Grove Village, Illinois.

FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending July 3, 1999, with historical data — United States



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

TABLE I. Summary — provisional cases of selected notifiable diseases, United States, cumulative, week ending July 3, 1999 (26th Week)

		Cum. 1999		Cum. 1999
Anthrax Brucellosis* Cholera		- 17 2	HIV infection, pediatric* <sup>§</sup> Plague Poliomyelitis, paralytic	81 2
	ibella syndrome is*	3 11	Psittacosis* Rabies, human Rocky Mountain spotted fever (RMSF)	14 - 148
Encephalitis:	California* eastern equine* St. Louis*	2 2	Streptococcal disease, invasive Group A Streptococcal toxic-shock syndrome* Syphilis, congenital <sup>§</sup>	1,152 22 94
Ehrlichiosis	western equine* human granulocytic (HGE)* human monocytic (HME)*	1 49 6	Tetanus Toxic-shock syndrome Trichinosis	11 63 5
		40 7 24	Typhoid fever Yellow fever	136

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

<sup>\*</sup>Not notifiable in all states.

<sup>\*</sup>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 from reports to the Division of HIV/AIDS Prevention–Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP), last update June 23, 1999.

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

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending July 3, 1999, and July 4, 1998 (26th Week)

								Escherichia coli O157:H7*				
	Al	IDS	Chla	mydia	Cryptosp	oridiosis	NET		PH	LIS		
Reporting Area	Cum. 1999†	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998		
UNITED STATES	23,194	23,725	281,030	286,678	661	972	738	820	361	690		
NEW ENGLAND	1,120	810	9,555	10,113	32	70	106	116	76	103		
Maine N.H.	29 26	18 15	193 458	461 477	10 5	18 3	10 15	10 18	8	19		
Vt. Mass.	6 716	10 372	235 4,521	194 4,118	6 11	9 36	12 42	4 60	2 39	4 60		
R.I.	61	69	1,181	1,229	-	4	6	3	6	1		
Conn.	282	326	2,967	3,634	-	-	21	21	21	19		
MID. ATLANTIC Upstate N.Y.	5,913 725	6,918 856	34,009 N	29,954 N	98 57	296 185	46 40	85 55	11 -	29		
N.Y. City	3,003	3,888	17,606	13,211	22	100	-	7	3	6		
N.J. Pa.	1,158 1,027	1,215 959	4,808 11,595	5,740 11,003	9 10	11 -	6 N	23 N	8 -	19 4		
E.N. CENTRAL	1,502	1,760	40,428	48,935	57	105	122	166	60	136		
Ohio Ind.	241 191	339 323	11,228 5,280	13,281 5,319	18 9	39 20	51 17	36 51	8 13	22 25		
III.	682	693	13,376	12,834	11	31	28	47	12	31		
Mich. Wis.	308 80	305 100	10,544 U	10,844 6,657	19	15	26 N	32 N	14 13	26 32		
W.N. CENTRAL	537	441	14,443	16,891	51	116	145	97	57	98		
Minn.	82 50	64 49	3,264	3,435	14 9	41 20	47 15	30 23	33 6	43 17		
lowa Mo.	261	210	1,225 5,099	2,071 5,990	11	11	15	23 13	13	21		
N. Dak. S. Dak.	4 11	4 9	325 803	498 798	4 3	14 14	3 5	2 6	- 4	6 8		
Nebr.	39	37	1,258	1,421	9	14	50	14	-	-		
Kans.	90	68	2,469	2,678	1	2	10	9	1	3		
S. ATLANTIC Del.	6,366 80	5,825 75	66,029 1,392	54,881 1,241	160	89 -	95 2	56 -	46 -	57 1		
Md.	720	717	4,848	4,131	7	8	6	12	-	7		
D.C. Va.	242 340	480 424	826 7,414	N 5,454	5 10	3 1	29	-	- 17	24		
W. Va. N.C.	31 390	51 389	1,011 11,466	1,171 10,898	- 4	1	4 22	3 12	1 16	2 13		
S.C.	588	381	8,635	9,311	-	-	11	2	3	1		
Ga. Fla.	958 3,017	618 2,690	15,832 14,605	11,919 10,604	86 48	28 48	6 15	21 6	9	9		
E.S. CENTRAL	1,034	933	19,520	19,595	8	15	52	51	19	35		
Ky.	152	126	3,333	3,051	2	5	14 23	15 22	12	23		
Tenn. Ala.	405 257	330 274	6,850 5,211	6,412 5,015	1	6	23 12	11	6	23 11		
Miss.	220	203	4,126	5,117	1	4	3	3	1	1		
W.S. CENTRAL Ark.	2,491 90	2,889 104	40,943 3,058	43,010 1,812	33	15 3	28 5	31 4	11 3	46 4		
La.	463	507	7,726	6,732	21	6	3	-	3	2		
Okla. Tex.	70 1,8 <b>6</b> 8	170 2,108	3,702 26,457	4,858 29,608	2 10	3 3	7 13	6 21	5 -	4 36		
MOUNTAIN	860	816	15,941	15,856	37	65	55	86	27	74		
Mont. Idaho	4 12	15 15	654 617	632 914	7 2	4 14	4 1	6 10	2	2 3		
Wyo.	3	1	333	329	- 4	3	3 22	2	4	16		
Colo. N. Mex.	172 46	146 130	3,726 1,731	3,978 1,878	15	26	3	22 10	12 1	19 6		
Ariz. Utah	427 80	327 65	6,474 946	5,409 1,144	7	10 1	11 9	15 15	4 2	11 10		
Nev.	116	117	1,460	1,572	2	7	2	6	2	7		
PACIFIC	3,371	3,333	40,162	47,443	185	201	89	132	54	112		
Wash. Oreg.	188 88	230 94	5,960 2,894	5,581 2,586	73	22	30 22	27 33	26 14	36 29		
Calif. Alaska	3,036	2,930 12	29,385 925	37,174 950	112	176	37	70 2	13	43		
Hawaii	13 46	67	925	1,152	-	3	-	-	1	4		
Guam	5	-	149	182	-	-	N	N	-			
P.R. V.I.	734 15	995 17	U N	U N	-	-	6 N	N	U U	U U		
Amer. Samoa	-	-	U	U	-	-	N N	N	U	Ü		
C.N.M.I.		-	N	N	-		IN	N	U	U		

N: Not notifiable U: Unavailable

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

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

<sup>\*</sup>Individual cases may be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the

Public Health Laboratory Information System (PHLIS).

†Updated monthly from reports to the Division of HIV/AIDS Prevention–Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention, last update June 23, 1999.

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending July 3, 1999, and July 4, 1998 (26th Week)

	Gono	orrhea	Hepa C/N/		Legion	ellosis	Lyr Dise	
Reporting Area	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998
UNITED STATES	151,968	165,675	1,817	1,538	454	541	2,647	3,495
NEW ENGLAND Maine	2,865 15	2,821	56 1	44	29 4	29 1	475	1,121
N.H.	38	31 46	-	-	3	3	-	18 16
Vt. Mass.	28 1,261	13 985	3 49	2 40	4 9	1 14	260	4 275
R.I. Conn.	304 1,219	179 1,567	3	2	3 6	4 6	77 1 <b>3</b> 8	31 777
MID. ATLANTIC	18,889	17,899	86	117	95	122	1,652	1,791
Upstate N.Y. N.Y. City	3,024 7,494	3,384 5,925	51 -	59 -	26 7	33 26	819 6	800 69
N.J. Pa.	2,760 5,611	3,548 5,042	35	- 58	5 57	5 58	124 703	327 595
E.N. CENTRAL	26,515	32,818	985	283	125	184	49	198
Ohio Ind.	6,668 3,049	8,169 3,066	1	6 4	41 39	65 31	26 20	19 11
III. Mich.	9,481 7,317	10,491 8,269	10 392	27 246	10 32	22 33	2 1	6 8
Wis.	Ū	2,823	582	-	3	33	U	154
W.N. CENTRAL Minn.	5,815 1,208	8,241 1,225	66 2	19 6	23 1	31 3	38 13	29 9
lowa Mo.	306 2.625	666 4,509	- 56	5 6	11 8	5 9	10	10 6
N. Dak. S. Dak.	31 80	44 127	-	-	1	1	1	-
Nebr.	553	539	3	2	2	11	6	2
Kans. S. ATLANTIC	1,012 48,013	1,131 44,210	5 120	- 54	- 54	2 64	8 290	2 266
Del. Md.	840 4,186	673 4,711	29	5	4 7	7 15	9 199	15 199
D.C.	2,490	1,966	10	- 5	-	4 7	1	4
Va. W. Va.	4,944 276	3,079 391	13	4	13 N	N	22 7	21 5
N.C. S.C.	9,750 4,645	9,146 6,043	25 12	12 2	8 7	6 5	34 4	13 2
Ga. Fla.	10,464 10,418	9,717 8,484	1 30	9 17	- 15	2 17	- 14	2 5
E.S. CENTRAL	15,362	18,428	120	80	55	32	44	31
Ky. Tenn.	1,494 5,349	1,753 5,421	8 44	15 62	44 9	17 7	19 13	10 11
Ala. Miss.	4,637 3,882	6,346 4,908	1 67	3	2	3 5	6 6	10 -
W.S. CENTRAL	22,652	25,788	128	278	2	10	7	8
Ark. La.	1,509 6,054	1,988 5,638	3 100	11 10	1	1 1	1	5 -
Okla. Tex.	1,878 13,211	2,635 15,527	6 19	2 255	1 -	6 2	4 2	3
MOUNTAIN Mont.	4,414 21	4,214 23	75 4	252 5	27	32 1	6	3
Idaho	32	83	4	85	-	-	1	1
Wyo. Colo.	11 1,061	15 1,029	25 15	59 13	5	1 6	- -	- -
N. Mex. Ariz.	311 2,305	371 1,951	4 18	52 4	1 4	2 3	1 -	-
Utah Nev.	89 584	112 630	2 3	18 16	11 6	16 3	1 2	- 1
PACIFIC	7,443	11,256	181	411	44	37	86	48
Wash. Oreg.	1,034 411	953 338	8 9	10 10	9 N	5 N	2 5	8
Calif. Alaska	5,718 152	9,571 157	164 -	336 1	34 1	31 -	79 -	37 1
Hawaii	128	237	-	54	-	1	-	-
Guam P.R.	22 145	24 210	-	-	-	2	-	-
V.I. Amer. Samoa	U U	U U	U U	U U	U U	U U	U U	U U
C.N.M.I.	-	19	-	-	-	-	-	-

N: Not notifiable

U: Unavailable

-: no reported cases

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending July 3, 1999, and July 4, 1998 (26th Week)

				-	Salmonellosis*						
	Ma	laria	Rabies,	Animal	NE	TSS	PH	LIS			
Reporting Area	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998			
UNITED STATES	535	595	2,645	3,743	13,207	15,304	9,355	14,287			
NEW ENGLAND	21	22	407 75	688	809 58	1,010	703	949			
Maine N.H.	2	3 3	75 27	127 33	44	72 69	35 39	29 101			
Vt. Mass.	1 8	- 14	60 91	30 223	33 474	45 557	26 407	39 552			
R.I.	2	2	50	36	49	62	48	37			
Conn.	8	-	104	239	151	205	148	191			
MID. ATLANTIC Upstate N.Y.	123 36	172 35	481 307	782 541	1,700 480	2,604 585	1,103 454	2,547 555			
N.Y. City	38	101	U	U	377	856	368	788			
N.J. Pa.	29 20	21 15	101 73	100 141	332 511	525 638	281 -	459 745			
E.N. CENTRAL	55	58	39	56	1,609	2,688	1,199	1,895			
Ohio Ind.	9 8	3 2	11	38 4	396 185	609 295	117 127	522 284			
III.	18	26	-	6	558	819	399	428			
Mich. Wis.	18 2	24 3	25 3	6 2	432 38	528 437	380 176	415 246			
W.N. CENTRAL	23	37	305	397	882	957	729	1,029			
Minn. Iowa	5 6	17 3	52 65	67 82	238 90	248 159	248 60	286 135			
Mo.	10	10	9	20	266	261	321	371			
N. Dak. S. Dak.	-	2	84 44	74 92	15 44	28 40	2 26	45 52			
Nebr.	-	1	2	3	105	79	-	20			
Kans. S. ATLANTIC	2 152	4 128	49 1,043	59 1,268	124 2,925	142 2,641	72 2,007	120 2,150			
Del.	1	1	29	20	43	30	51	48			
Md. D.C.	48 10	44 10	216 -	266	336 39	363 44	296	396			
Va.	30	22	265	336	503	419	371	391			
W. Va. N.C.	1 10	12	62 205	42 325	43 450	67 385	37 414	71 444			
S.C. Ga.	1 12	4 15	78 99	77 103	172 453	167 412	134 543	147 439			
Fla.	39	20	89	99	886	754	161	214			
E.S. CENTRAL	10	16	134	148	696	732	263	627			
Ky. Tenn.	2 5	2 8	22 48	18 84	161 191	170 218	139	89 334			
Ala. Miss.	2 1	4 2	64	44 2	220 124	189 155	107 17	166 38			
W.S. CENTRAL	8	11	54	104	990	1,187	653	1,568			
Ark.	6	1		19	166	123	76	93			
La. Okla.	1	1	54	85	159 145	201 149	66 88	287 58			
Tex.	1	5	-	-	520	714	423	1,130			
MOUNTAIN Mont.	23 3	32	95 35	97 29	1,307 28	940 41	802 1	879 22			
ldaho	1	3	28	-	40	52	35 17	41 27			
Wyo. Colo.	1 8	- 7	1	41 2	15 384	32 236	367	228			
N. Mex. Ariz.	2 5	11 5	2 29	2 21	145 414	91 264	79 250	84 269			
Utah	2	1	-	2	203	145	-	120			
Nev.	1	5	-	-	78	79	53	88			
PACIFIC Wash.	120 10	119 9	87 -	203	2,289 221	2,545 192	1,896 279	2,643 320			
Oreg. Calif.	13 91	11 97	1 80	1 182	180 1,687	141 2,093	205 1,291	184 2,012			
Alaska	-	-	6	20	21	19	6	15			
Hawaii	6	2	-	-	180	100	115	112			
Guam P.R.	-	1 -	36	28	18 184	12 310	-	-			
V.I. Amer. Samoa	U U	U U	Ü	Ü	-	-	-	-			
C.N.M.I.	-	-	-	-	-	13	-	<u> </u>			

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

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending July 3, 1999, and July 4, 1998 (26th Week)

	I	Shige			Sypt	T		
	NE.	TSS		ILIS	(Primary &		Tubero	ulosis
Reporting Area	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999†	Cum. 1998†
UNITED STATES	5,833	8,600	2,007	5,235	3,111	3,418	3,992	4,838
NEW ENGLAND	150	211	126	185	30	37	187	224
Maine N.H.	3 7	7 7	6	9	-	1 1	10 4	5 6
Vt.	4	4	3	-	2	3	-	1
Mass. R.I.	95 14	132 15	82 9	124 12	19 1	23	106 19	117 30
Conn.	27	46	26	40	8	9	48	65
MID. ATLANTIC	384	1,287	185	1,091	126	115	974	1,106
Upstate N.Y. N.Y. City	113 98	245 419	31 81	77 453	17 57	18 25	138 609	151 655
N.J.	103	392	73	382	16	54	227	300
Pa.	70 832	231	-	179	36 606	18	U 428	U 603
E.N. CENTRAL Ohio	256	1,268 283	334 14	642 67	47	505 76	428 U	603 U
Ind.	54	87	11	24	178	91	U	U
III. Mich.	312 162	665 123	218 73	528 4	268 113	211 89	252 137	376 173
Wis.	48	110	18	19	U	38	39	54
W.N. CENTRAL Minn.	514 84	450 79	311 83	196 84	52 5	77 5	241 95	195 66
lowa	7	33	9	27	5	-	26	2
Mo.	361	57 4	201	39 3	34	59	84 2	82 3
N. Dak. S. Dak.	2 8	22	4	3 18	-	1	3	3 14
Nebr.	30	239	-	15	4	4	12	5
Kans. S. ATLANTIC	22 1,106	16 1,678	14 239	10 535	4 1.013	8 1,319	19 815	23 833
Del.	7	1,678	239	2	1,013	1,319	12	033 17
Md. D.C.	59 30	98	15	30	201 42	369 49	U 24	U 58
Va.	40	11 69	10	28	89	87	104	144
W. Va.	5	7	2 54	5	2 243	2	23	24
N.C. S.C.	113 55	142 78	18	83 31	243 125	370 161	209 124	204 161
Ga.	105	453	34	135	156 151	139	319	225
Fla. E.S. CENTRAL	692 626	811 426	104 217	221 252	151 573	127 591	U 284	U 405
Ky.	113	77	-	38	46	59	82	95
Tenn. Ala.	419 55	69 250	197 19	94 118	327 130	285 135	U 146	U 194
Miss.	39	30	1	2	70	112	56	116
W.S. CENTRAL	877	1,695	339	1,883	460	456	752	1,041
Ark. La.	47 76	80 130	21 29	16 159	38 121	60 155	80 U	53 U
Okla.	267	119	77	30	103	25	63	66
Tex.	487	1,366	212	1,678	198	216	609	922
MOUNTAIN Mont.	350 6	536 3	152 -	311 3	111 -	127 -	62 5	134 12
Idaho	6 2	11	3	8	1	-	-	7
Wyo. Colo.	2 52	1 66	1 37	49	- 1	1 8	1 U	2 U
N. Mex.	40	129	13	53	-	18	23	31
Ariz. Utah	197 26	291 16	92	178 13	102 2	87 3	U 18	U 33
Nev.	21	19	6	7	5	10	15	49
PACIFIC	994	1,049	104	140	140	191	249	297
Wash. Oreg.	52 35	57 64	51 34	58 58	39 2	12 1	82 57	124 58
Calif.	885	904	-	-	96	178	U	U
Alaska Hawaii	22	4 20	19	2 22	1 2	-	29 81	26 89
Guam	3	20	-	-	-	-	-	39
P.R. V.I.	23	28	-	-	82 U	113 U	41 U	80 U
v.i. Amer. Samoa	-	-	-	-	Ü	U	Ü	U
C.N.M.I.	-	12	-	-	-	135	-	58

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

†Cumulative reports of provisional tuberculosis cases for 1998 and 1999 are unavailable ("U") for some areas using the Tuberculosis Information System (TIMS)

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending July 3, 1999, and July 4, 1998 (26th Week)

		ienzae,		lepatitis (Vi						les (Rube			
		sive		A		3	Indi	genous	Imp	orted*		tal	
Reporting Area	Cum. 1999 <sup>†</sup>	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	1999	Cum. 1999	1999	Cum. 1999	Cum. 1999	Cum. 1998	
UNITED STATES	610	610	7,729	11,282	3,179	4,473	1	30	-	14	44	40	
NEW ENGLAND	42	41	91	152	53	95	-	5	-	4	9	2	
Maine N.H.	5 9	2 6	4 7	13 7	8	2 10	-	-	-	1	1	-	
Vt.	4	2	3	13	1	4	-	-	-	-	-	-	
Mass. R.I.	17 -	29 2	30 9	51 9	28 16	36 24	-	4	-	2	6	2	
Conn.	7	-	38	59	-	19	-	1	-	1	2	-	
MID. ATLANTIC Upstate N.Y.	85 49	92 29	510 128	856 166	392 103	640 124	-	-	-	2 2	2 2	11 2	
N.Y. City	13	28	82	313	89	219	-	-	-	-	-	-	
N.J. Pa.	23	28 7	57 243	160 217	40 160	107 190	-	-	-	-	-	8 1	
E.N. CENTRAL	83	98	1,499	1,552	304	498	_	1	-	-	1	15	
Ohio Ind.	35 14	34 23	366 98	177 89	45 27	37 55	-	- 1	-	-	- 1	1 3	
III.	27	23 37	220	384	-	133	-	-	-	-	-	-	
Mich. Wis.	7	- 4	789 26	777 125	231 1	225 48	-	-	-	-	-	10 1	
W.N. CENTRAL	49	51	374	876	244	209	-	_	-	-	-		
Minn.	13	37	33	69	19	18	-	-	-	-	-	-	
lowa Mo.	13 16	1 8	76 195	355 365	103 94	33 129	-	-	-	-	-	-	
N. Dak.	-	-	1	3	-	4	U	-	U	-	-	-	
S. Dak. Nebr.	1 3	-	8 33	16 14	1 10	1 9	-	-	-	-	-	-	
Kans.	3	5	28	54	17	15	U	-	U	-	-	-	
S. ATLANTIC Del.	144	112	954 2	863 3	571 -	466	-	1	-	3	4	6 1	
Md.	33	38	159	175	85	88	-	-	-	-	-	i	
D.C. Va.	4 12	12	32 79	30 129	11 51	6 53	-	1	-	2	3	2	
W. Va.	4	4	17	1	13	3	-	-	-	-	-	-	
N.C. S.C.	22 2	15 3	65 19	51 17	117 38	110 9	-	-	-	-	-	-	
Ga. Fla.	38 29	22 18	259 322	247 210	66 190	90 107	-	-	-	- 1	- 1	1 1	
E.S. CENTRAL	46	37	237	225	235	206	_	_	_			1	
Ky.	6	5	37	14	25	23	-	-	-	-	-	-	
Tenn. Ala.	25 13	23 7	125 36	127 45	118 47	142 41	-	-	-	-	-	1	
Miss.	2	2	39	39	45	-	-	-	-	-	-	-	
W.S. CENTRAL Ark.	34 1	30	1,415 26	1,992 43	298 25	1,009 49	-	1	-	2	3	-	
La.	7	13	59	41	72	47	-	-	-	-	-	-	
Okla. Tex.	24 2	15 2	258 1,072	290 1,618	67 134	31 882	-	1	-	2	3	-	
MOUNTAIN	60	77	747	1,725	321	437	-	2	-	-	2	-	
Mont. Idaho	1 1	-	12 27	56 140	16 16	3 17	- U	-	- U	-	-	-	
Wyo.	1	-	4	23	5	2	-	-	-	-	-	-	
Colo. N. Mex.	9 13	14 4	134 29	129 86	45 110	52 168	-	-	-	-	-	-	
Ariz.	29	39	454	1,059	84	107	-	1	-	-	1	-	
Utah Nev.	4 2	3 17	25 62	115 117	17 28	39 49	Ū	1 -	Ū	-	1 -	-	
PACIFIC	67	72	1,902	3,041	761	913	1	20	-	3	23	5	
Wash. Oreg.	2 26	4 30	164 141	570 240	33 50	53 93	-	- 8	-	-	- 8	1	
Calif.	32	31	1,585	2,188	661	752	1	11	-	3	14	4	
Alaska Hawaii	5 2	1 6	3 9	14 29	10 7	7 8	-	1	-	-	- 1	-	
Guam	-	-	2	-	2	2	U	1	U	-	1	-	
P.R. V.I.	1 U	2 U	80 U	25 U	76 U	130 U	Ū	Ū	- U	- U	Ū	- U	
Amer. Samoa	Ü	U	Ü	U	U	U	U	Ü	U	Ü	U	U	
C.N.M.I.	-	-	-	1	-	35	U	-	U	-	-	-	

N: Not notifiable

U: Unavailable

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

<sup>\*</sup>For imported measles, cases include only those resulting from importation from other countries.

<sup>&</sup>lt;sup>†</sup>Of 127 cases among children aged <5 years, serotype was reported for 58 and of those, 13 were type b.

TABLE III. (Cont'd.) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending July 3, 1999, and July 4, 1998 (26th Week)

	Meningococcal Disease			Mumps	1000 (2		Pertussis		Rubella			
Reporting Area	Cum. 1999	Cum. 1998	1999	Cum. 1999	Cum. 1998	1999	Cum. 1999	Cum. 1998	1999	Cum. 1999	Cum. 1998	
UNITED STATES	1,330	1,582	4	180	406	32	2,539	2,448	3	138	291	
NEW ENGLAND	74	69	-	3	1	-	254	450	-	6	37	
Maine N.H.	5 10	4 8	-	1	-	-	- 53	5 34	-	-	-	
Vt.	4	1	-	-	-	-	9	38	-	-	-	
Mass. R.I.	45 2	30 3	-	2	1 -	-	176 8	355 3	-	6	8 -	
Conn.	8	23	-	-	-	-	8	15	-	-	29	
MID. ATLANTIC Upstate N.Y.	117 34	165 43	1	22 5	168 2	9 3	577 498	299 148	-	17 13	130 108	
N.Y. City	27	20	-	3	153	-	10	14	-	-	9	
N.J. Pa.	23 33	39 63	- 1	14	5 8	6	12 57	8 129	-	1 3	12 1	
E.N. CENTRAL	206	244	-	23	49	1	208	229	_	1	-	
Ohio	91	82	-	7	19	-	107	72	-	-	-	
Ind. III.	37 50	43 69	-	3 6	5 8	-	14 38	61 26	-	1 -	-	
Mich. Wis.	27 1	26 24	-	7	17 -	1 -	22 27	32 38	-	-	-	
W.N. CENTRAL	151	133	_	- 7	20	8	92	176	-	71	29	
Minn.	30	24	-	1	10	8	33	100	-	-	-	
lowa Mo.	28 59	19 52	-	3 1	6 3	-	20 15	43 13	-	21 2	2	
N. Dak.	3	-	U	-	1	U	-	-	U	-	-	
S. Dak. Nebr.	8 9	6 8	-	-	-	-	4 1	4 6	-	48	-	
Kans.	14	24	U	2	-	U	19	10	U	-	27	
S. ATLANTIC Del.	231 3	256 1	1 -	36	26	4	142	122 1	3	20	7	
Md.	35	23	-	3	-	1	39	27	-	1	-	
D.C. Va.	1 26	23	-	2 8	- 5	-	13	1 6	-	-	-	
W. Va. N.C.	4 27	9 39	-	8	8	-	1 35	1 44	- 3	- 19	- 5	
S.C.	28	41	-	3	4	-	8	15	- -	-	-	
Ga. Fla.	41 66	58 <b>62</b>	- 1	2 10	1 8	3	16 30	6 21	-	-	2	
E.S. CENTRAL	108	116	-	1	8	-	43	53	_	1	-	
Ky.	29	16	-	-	-	-	3	20	-	-	-	
Tenn. Ala.	38 24	41 40	-	1	1 4	-	25 11	17 14	-	1	-	
Miss.	17	19	-	-	3	-	4	2	-	-	-	
W.S. CENTRAL Ark.	97 22	186 23	-	21	35	1 1	62 7	150 16	-	5	70 -	
La.	34	35	-	3	5	-	3	1	-	-	-	
Okla. Tex.	19 22	27 101	-	1 17	30	-	7 45	15 118	-	5	70	
MOUNTAIN	89	85	-	12	24	3	248	508	-	14	5	
Mont. Idaho	2 8	3 4	- U	1	3	- U	2 93	1 184	- U	-	-	
Wyo.	3	3	-	-	1	-	2	7	-	-	-	
Colo. N. Mex.	24 11	17 15	N	3 N	3 N	3	60 27	120 64	-	-	1	
Ariz. Utah	28 8	30 8	-	- 5	5 3	-	29 33	88 26	-	13	1 2	
Nev.	5	5	Ū	3	9	Ū	2	18	Ū	1	1	
PACIFIC	257	328	2	55	75	6	913	461	-	3	13	
Wash. Oreg.	38 44	41 55	- N	2 N	5 N	3 1	502 18	148 29	-	-	9	
Calif.	166	227	1	46	54	2	383 3	275	-	3	2	
Alaska Hawaii	5 4	1 4	1	1 6	2 14	-	3 7	2 7	-	-	2	
Guam	-	2	U	1	2	U	1	-	U	-	-	
P.R. V.I.	5 U	6 U	Ū	Ū	2 U	Ū	9 U	3 U	Ū	Ū	Ū	
Amer. Samoa	Ü	U	U	U	U	U	U	U	U	U	Ū	
C.N.M.I.	-	-	U	-	2	U	-	1	U	-	-	

N: Not notifiable

U: Unavailable

-: no reported cases

TABLE IV. Deaths in 122 U.S. cities,\* week ending July 3, 1999 (26th Week)

	All Courses By Arra (Vesus)							(Lotti Frook,		All Car	ıses. Rv	/ Age (Y	ears)		•
Reporting Area	All Ages	>65	45-64	25-44	1-24	<1	P&I <sup>†</sup> Total	Reporting Area	All Ages	>65	45-64		1-24	<1	P&l <sup>†</sup> Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn. Cambridge, Mass. Fall River, Mass. Hartford, Conn. Lowell, Mass. Lynn, Mass. New Bedford, Mass. New Haven, Conn. Providence, R.I. Somerville, Mass. Springfield, Mass. Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y.	35 U 6 U 22 64 2,000 58	157 U 14 11 U U 22 24 U 5 U 15 46 1,352 42	3 3 3 0 0 7 0 4 7 0 1 0 3 15 392 12	11 U 4 - U U 2 2 U - U 1 2 162 2	3 U 1	4 U - U - U - 1 U - U - 2 1 39 2	21 1 1 1 1 2 2 4 4 4 0 - 1 2 8 66	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.	91	466 U 94 50 101 U 33 37 30 U 121 U U 528 103 47 64	21 30 U 10 11 4 U 38 U U 171 41 6 17	50 U 15 13 4 U 2 5 1 U 10 U 71 10 5 8	14 U 2 2 3 U 1 - - U 6 U U 29 5 3	12 U 3 - U 1 4 - U 4 U U 9 - 2	50 15 12 7 U 5 2 4 U 5 U 34 10 4 4
Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Erie, Pa.	U 85 U U 41	U 62 U U 30	15 U U 9	U 3 U U	U 3 U U 1	U 2 U U	U 1 U 1	Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala. Nashville, Tenn.	94 200 80 U 123	63 124 55 U 72	19 38 18 U 32	7 23 7 U 11	2 11 - U 8	3 4 U	7 9 - U -
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.	32 1,127 U 22 299 45 34 121 U 33 76 U 27 U	24 755 U 16 183 29 28 87 U 22 57 U 17 U	223 U - 68 8 4 21 U 9 8 U 9	1 101 U 1 34 - 8 U 1 7 U	1 27 U - 10 2 1 2 U 1 - U 1 U 1 U 1	21 U - 4 2 1 3 U - 4 U -	30 8 6 2 11 U 2 5 U · U	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.	751 U 48 U U 114 U 67 130 202 66 124	523 U U 34 U 76 U 47 89 143 47 87	138 U U 11 U 20 U 14 25 33 12 23	50 UU 1 UU 9 UU 10 18 38	23 U U 2 U U 2 U 3 6 5 2 3	17 U U - U U 7 U 2 - 3 2 3	46 U U 3 U U 6 U 4 4 15 7
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.	1,818 54 27 395 70 142 164 112 202 U 79 20	1,233 35 21 249 48 97 122 85 116 U 63	13 2 86 7 29 30 20 50 U 12 5	131 5 2 37 5 9 8 6 24 U 2 2	43 10 3 3 1 1 6 U 1 2	47 1 2 11 7 4 3 - 6 U 1 2	107 1 29 4 6 15 7 7 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	U 204 25 56 25	332 U U 39 U 136 19 43 19 U 76	103 UU 5 UU 49 4 5 3 U 37 231	37 UU 6 U 12 2 7 3 U 7 90	12 U U 2 U 5 - 1 - U 4	6 U U 2 - - - U 4 33	24 U U 1 U 9 1 5 3 U 5
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	149 42 89 53 42 30 96 U 424 55	33 95 32 64 39 31 24 70 U 318 43	30 8 14 9 5 21 U 65 9	4 12 2 8 2 1 1 1 U	2 8 1 2 1 2 U	1 4 - 2 1 - - 2 U	3 9 1 9 2 3 6 U 26 4	Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawaii Long Beach, Calif. Los Angeles, Calif. Pasadena, Calif. Portland, Oreg. Sacramento, Calif. San Diego, Calif. San Francisco, Calif.	19 135 18 67 70 269 U 117 U 134	9 96 16 50 52 183 U 82 U 86 U	6 26 1 12 12 53 U 20 U 28 U	2 8 1 4 3 19 U 5 U 18 U	2 - 1 6 U 4 U 1 U	2 3 1 2 8 U 6 U 1 U	1 10 5 13 18 U 5 U 14 U
Duluth, Minn. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	U 72 25 205 U U 67	U 54 20 151 U U 50 U	33 U U 12	U 4 2 10 U 1 U	U 4 - 6 U U 2 U	U 2 5 U 2 U 2 U	UU313UU5U	San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash. TOTAL	207 U 108 U 72 8,410 <sup>¶</sup>	144 U 66 U 51 5,744	26 U 11	15 U 9 U 6	8 U 1 U 2 210	4 U 6 U - 176	16 U 3 U 2 461

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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☆U.S. Government Printing Office: 1999-733-228/08008 Region IV