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The Great American Smokeout — November 19, 1998

In 1995, an estimated 47 million U.S. adults smoked cigarettes; in 1997, at least 4.5 million U.S. adolescents were cigarette smokers (*1,2*). Since 1977, the American Cancer Society (ACS) has sponsored the Great American Smokeout to promote community-based activities designed to encourage smokers to refrain from smoking cigarettes for at least 24 hours. In 1997, nearly 11.3 million smokers (approximately 24% of smokers) reported participating in the Smokeout, and 19% of participants reported smoking less or not at all 1–5 days after the Smokeout (*3*). This year, the Great American Smokeout on Thursday, November 19, will focus on preventing the use of all tobacco products and encouraging children and adolescents never to start using tobacco.

As part of the Great American Smokeout, ACS volunteers will conduct smokingprevention and smoking-cessation activities for persons of all ages at shopping malls, worksites, hospitals, military installations, and other locations. Activities will include the ACS *Commit to Quit* program, which helps smokers select a method of quitting that meets their personal needs.

Additional information is available from ACS, telephone (800) 227-2345; CDC, telephone (800) 232-1311 or (770) 488-5705; or the ACS Great American Smokeout World-Wide Web site http://www.cancer.org.

Reported by: American Cancer Society, Atlanta, Georgia. Office on Smoking and Health, National Center for Chronic Disease Prevention and Health Promotion, CDC.

References

1. CDC. Cigarette smoking among adults-United States, 1995. MMWR 1997;46:1217-20.

- Substance Abuse and Mental Health Services Administration. Preliminary results from the 1997 National Household Survey on Drug Abuse. Rockville, Maryland: US Department of Health and Human Services, Substance Abuse and Mental Health Services Administration, 1998.
- 3. Lieberman Research Inc. A study of public reactions to the 1997 Great American Smokeout and American Cancer Society: a partnership with Nicoderm CQ and Nicorette gum. Atlanta, Georgia: American Cancer Society, 1997.

State-Specific Prevalence Among Adults of Current Cigarette Smoking and Smokeless Tobacco Use and Per Capita Tax-Paid Sales of Cigarettes — United States, 1997

In the United States each year, tobacco use causes approximately 400,000 deaths and is the single most preventable cause of death and disease (1,2). Consequently, state and local public health agencies closely monitor tobacco use and its correlates (3). In 1996, the prevalence of current cigarette smoking among adults was the first health behavior and the first noninfectious condition added by the Council of State and Territorial Epidemiologists (CSTE) to the list of nationally notifiable conditions reported to CDC (4). In 1998, per capita sales of cigarettes (along with prevalence among youth of current cigarette smoking and current smokeless tobacco use) was added by CSTE to the list of notifiable conditions reported by states to CDC. This report summarizes state-specific findings for current cigarette and current smokeless tobacco use by adults from the Behavioral Risk Factor Surveillance System (BRFSS) and number of packs of tax-paid cigarettes sold per capita in each state from data compiled annually by The Tobacco Institute. The findings indicate that current adult cigarette smoking prevalence by state ranged from 13.7% to 30.8%, annual per capita tax-paid cigarette sales ranged from 49.1 packs to 186.8 packs, and adult smokeless tobacco use prevalence ranged from 1.4% to 8.8%.

State- and sex-specific prevalences of current cigarette smoking and current smokeless tobacco use among adults are available from the 1997 BRFSS. The BRFSS is a state-specific, random-digit-dialed telephone survey of health behaviors of the civilian, noninstitutionalized U.S. population aged \geq 18 years (5) conducted by state health departments with assistance from CDC. In 1996 and 1997, respondents were asked, "Have you smoked at least 100 cigarettes in your entire life?" and "Do you now smoke cigarettes every day, some days, or not at all?" Current cigarette smokers were defined as persons who reported having smoked at least 100 cigarettes during their lifetime and who currently smoke every day or some days. To determine current smokeless tobacco use, respondents were asked, "Have you ever used or tried any smokeless tobacco products such as chewing tobacco or snuff?" and "Do you currently use any smokeless tobacco products such as chewing tobacco or snuff?" Current smokeless tobacco users were defined as persons who reported having ever used or tried any smokeless tobacco product and who currently use a smokeless tobacco product. To estimate prevalence, responses for each state were weighted to the current age, race, and sex distribution of the state's population (i.e., crude prevalence). To allow comparison of findings across states that had different age distributions, ageadjusted prevalences for each state were estimated by using direct standardization to 10-year age groups of the U.S. population in 1997 derived from U.S. census estimates (6). The number of packs of tax-paid cigarettes sold per capita in each state is compiled yearly by The Tobacco Institute by using information on federal, state, and local excise taxes and total population estimates (7).

In 1997, the median state prevalence of current cigarette smoking by adults was 23.2%; prevalence was 25.5% for men and 21.3% for women (Table 1). The crude median prevalence of current cigarette smoking was similar in 1997 and in 1996 (25.5% for men, 22.0% for women, and 23.6% for both groups combined) (4). In 1997, for

Cigarettes Smoking and Smokeless Tobacco Use - Continued

		Men	N	/omen	1	Total	Per capita tax-paid sales of cigarettes
State	%	(95% Cl [§])	$\begin{array}{ c c c c c c c c c c c c c c c c c c c$		(95% CI)	(in packs)	
Alabama	28.6	(±3.3)	21.3	(±2.5)	24.7	(±2.0)	104.9
Alaska	27.4	(±4.9)	25.8	(±4.3)	26.7	(±3.3)	81.7
Arizona	22.1	(±3.9)	20.2	(±3.6)	21.1	(±2.5)	64.6
Arkansas	32.1	(±4.4)	25.2	(±3.0)	28.5	(±2.6)	108.7
California	22.4	(±2.3)	14.5	(±1.6)	18.4	(±1.4)	53.8
Colorado	24.0	(±3.2)	21.2	(±3.0)	22.6	(±2.2)	81.3
Connecticut	21.4	(±3.2)	22.2	(±2.7)	21.8	(±2.1)	75.9
Delaware	29.3	(±3.5)	24.2	(±2.5)	26.6	(±2.1)	124.1
District of Columbia	22.7	(±4.0)	15.5	(±2.8)	18.8	(±2.4)	54.3
Florida	26.0	(±2.6)	21.4	(±1.9)	23.6	(±1.6)	93.0
Georgia	25.2	(±3.2)	19.9	(±2.7)	22.4	(±2.1)	100.6
Hawaii	21.4	(±2.9)	15.8	(±2.5)	18.6	(±1.9)	49.1
ldaho	21.8	(±2.2)	18.0	(±1.9)	19.9	(±1.4)	75.0
Illinois	25.0	(±2.7)	21.6	(±2.2)	23.2	(±1.7)	79.6
Indiana	29.2	(±3.2)	23.7	(±2.7)	26.3	(±2.1)	135.3
lowa	25.5	(±2.4)	20.9	(±2.0)	23.1	(±1.6)	93.9
Kansas	26.8	(±3.4)	18.9	(±2.3)	22.7	(±2.0)	89.2
Kentucky	33.1	(±2.9)	28.7	(±2.1)	30.8	(±1.8)	186.8
Louisiana	29.3	(±4.1)	20.4	(±2.7)	24.6	(±2.4)	105.3
Maine	25.2	(±3.3)	20.4	(±2.8)	22.7	(±2.2)	101.1
Maryland	21.8	(+2.4)	19.4	(±2.0)	20.6	(±1.6)	72.7
Massachusetts	21.8	(±3.7)	19.2	(±2.6)	20.4	(±2.2)	66.7
Michigan	29.6	(±3.0)	22.8	(±2.2)	26.1	(±1.9)	75.6
Minnesota	24.1	(±2.0)	19.8	(±1.6)	21.8	(±1.3)	84.1
Mississippi	28.3	(±4.2)	18.6	(±2.8)	23.2	(±2.5)	106.3
Missouri	31.7	(±4.1)	26.0	(±2.9)	28.7	(±2.5)	120.6
Montana	20.8	(±3.0)	20.2	(±2.6)	20.5	(±2.0)	88.9
Nebraska	24.4	(±3.1)	20.2	(±2.6)	22.2	(±2.0)	88.5
Nevada	25.7	(±5.0)	29.8	(±4.6)	27.7	(±3.4)	95.6
New Hampshire	26.0	(±4.1)	23.7	(±3.0)	24.8	(±2.5)	174.4
New Jersey	23.3	(±3.0)	19.8	(±2.3)	21.5	(±1.9)	77.0
New Mexico	21.6	(±3.2)	22.6	(±2.7)	22.1	(±2.1)	61.8
New York	25.0	(±2.6)	21.5	(±2.0)	23.1	(±1.6)	64.5
North Carolina	29.7	(±2.7)	22.3	(±2.0)	25.8	(±1.7)	125.6
North Dakota	24.3	(±3.2)	20.3	(±2.7)	22.2	(±2.1)	77.5
Ohio	26.3	(±3.2)	24.0	(±2.5)	25.1	(±2.0)	108.6
Oklahoma	25.2	(±3.7)	24.1	(±3.0)	24.6	(±2.4)	111.8
Oregon	22.1	(±2.7)	19.4	(±2.1)	20.7	(±1.7)	89.5
Pennsylvania	26.2	(±2.6)	22.5	(±2.0)	24.3	(±1.6)	92.9
Rhode Island	25.6	(±3.6)	23.0	(±3.2)	24.2	(±2.4)	90.0
South Carolina	29.5	(±3.5)	17.8	(±2.3)	23.4	(±2.1)	124.5
South Dakota	28.1	(±3.3)	20.8	(±2.6)	24.3	(±2.1)	88.8
Tennessee	27.9	(±3.1)	26.0	(±2.2)	26.9	(±1.9)	118.9
Texas	28.0	(±3.1)	17.5	(±2.2)	22.6	(±1.9)	72.6
Utah	16.1	(±2.4)	11.5	(±2.0)	13.7	(±1.6)	57.0
Vermont	25.1	(±2.9)	21.5	(±2.4)	23.2	(±1.9)	97.7
Virginia	26.2	(±3.4)	23.1	(±2.6)	24.6	(±2.1)	108.0
Washington	25.1	(±2.8)	22.7	(±2.2)	23.9	(±1.8)	55.6
West Virginia	27.1	(±3.1)	27.7	(±2.6)	27.4	(±2.0)	114.5
Wisconsin	25.6	(±3.4)	21.0	(±2.8)	23.2	(±2.2)	91.9
Wyoming	24.0	(±3.8)	24.1	(±2.8)	24.0	(± 2.4)	108.8

TABLE 1. Prevalence of current cigarette smoking* among adults, by state and	sex,
and per capita tax-paid sales of cigarettes, by state [†] — United States, 1997	

* Percentage of persons aged ≥18 years who reported having smoked ≥100 cigarettes during their lifetime and who currently smoke every day or some days. Estimates are weighted to the age, race, and sex distribution of the state population (crude prevalence). Source: Behavioral Risk Factor Surveillance System. [†] Source: The Tobacco Institute. Data are for July 1, 1996, through June 30, 1997 (7).

Cigarettes Smoking and Smokeless Tobacco Use -- Continued

every state except Florida, the crude prevalence of current cigarette smoking was within 1% of the age-adjusted prevalence for that state.

Current adult cigarette smoking prevalence differed approximately twofold across the states (Table 1). In 1997, the current cigarette smoking prevalence was highest in Kentucky (30.8%), Missouri (28.7%), Arkansas (28.5%), Nevada (27.7%), and West Virginia (27.4%), and lowest in Utah (13.7%), California (18.4%), Hawaii (18.6%), the District of Columbia (18.8%), and Idaho (19.9%). The current cigarette smoking prevalence for men was highest in Kentucky (33.1%), and for women in Nevada (29.8%). For both men and women, current smoking prevalence was lowest in Utah.

Per capita tax-paid sales of cigarettes for July 1, 1996, through June 30, 1997, varied approximately fourfold across the states (Table 1). The state median tax-paid cigarette sales was 90 packs per person per year. Sales were highest in Kentucky (186.8 packs) and lowest in Hawaii (49.1 packs).

Questions about current adult smokeless tobacco use were included in the 1997 BRFSS in 17 states (Table 2). The difference in prevalence was more than sixfold (from 1.4% in Arizona to 8.8% in West Virginia). Among men, the prevalence of current smokeless tobacco use was highest in West Virginia (18.4%) and Wyoming (14.7%); five states (Alabama, Alaska, Kansas, Kentucky, and Montana) reported prevalences of 9%–12%, and 10 states reported prevalences of \leq 8%. For women, the prevalence of current smokeless tobacco use was \leq 1.7% in all 17 states.

Reported by the following BRFSS coordinators: J Cook, Alabama, MBA; P Owen, Alaska; B Bender, MBA, Arizona; J Senner, PhD, Arkansas; B Davis, PhD, California; M Leff, MSPH, Colorado; M Adams, MPH, Connecticut; F Breukelman, Delaware; C Mitchell, District of Colum-

		Men	W	/omen	Т	otal
State	%	(95% CI ⁺)	%	(95% CI)	%	(95% CI)
Alabama	9.9	(±2.2)	1.4	(±0.7)	5.4	(±1.1)
Alaska	9.2	(±3.2)	1.6	(±1.0)	5.5	(±1.7)
Arizona	2.6	(±1.3)	0.3	(±0.3)	1.4	(±0.7)
Georgia	6.4	(±1.8)	1.7	(±0.9)	4.0	(±1.0)
Indiana	6.8	(±1.7)	0.0	(±0.0)	3.2	(±0.8)
Kansas	10.3	(±2.5)	0.2	(±0.3)	5.1	(±1.2)
Kentucky	12.2	(±3.0)	0.6	(±0.5)	6.1	(±1.5)
Louisiana	7.6	(±2.1)	0.3	(±0.4)	3.7	(±1.1)
Montana	10.5	(±2.5)	0.2	(±0.3)	5.3	(±1.3)
Ohio	5.1	(±1.6)	0.0	(±0.1)	2.4	(±0.8)
Oklahoma	7.7	(±2.2)	0.3	(±0.3)	3.8	(±1.1)
Pennsylvania	7.4	(±1.7)	0.4	(±0.3)	3.8	(±0.9)
South Carolina	4.8	(±1.7)	1.0	(±0.6)	2.8	(±0.9)
Virginia	6.1	(±1.4)	0.1	(±0.1)	3.0	(±0.7)
Washington	5.6	(±1.4)	0.2	(±0.2)	2.9	(±0.7)
West Virginia	18.4	(±2.6)	0.2	(±0.2)	8.8	(±1.3)
Wyoming	14.7	(±2.3)	0.7	(±0.4)	7.6	(±1.2)

 TABLE 2. Prevalence of current smokeless tobacco use* among adults, by state and sex — United States, 1997

*Percentage of persons aged ≥18 years who reported having ever used or tried smokeless tobacco products such as chewing tobacco or snuff and who currently use a smokeless tobacco product. Estimates are weighted to the age, race, and sex distribution of the state population (crude prevalence). Source: Behavioral Risk Factor Surveillance System.

[†]Confidence interval.

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Editorial Note: This report includes information about two CSTE-recommended indicators of tobacco use for all states (current cigarette smoking by adults and per capita tax-paid sales of cigarettes) and current smokeless tobacco use among adults for 17 states. Information on cigarette and smokeless tobacco use by youth in 1997 is available elsewhere (8). National surveys provide information about tobacco use and are useful for monitoring overall trends, but their effectiveness is limited for monitoring state-level year-to-year changes in tobacco consumption. National surveys also mask the twofold variation in current adult cigarette smoking prevalence among the states.

In the BRFSS, the crude and age-adjusted prevalences of current adult cigarette smoking were similar, indicating that differences in prevalence among states are related primarily to factors other than differences in adult age distributions. Although the median prevalence for current cigarette smoking among adults was nearly the same in 1996 and 1997, the twofold difference in prevalence among states, the wide variation in per capita tax-paid cigarette sales, and the wide variation in smokeless tobacco prevalence among adults suggest that further reductions in tobacco use are achievable.

The findings in this report are subject to at least three limitations. First, the BRFSS standardizes procedures among states, but the quality and completeness of the surveys can vary by state and year. Second, the changes in questions about current cigarette use in 1996 limit comparisons with previous years (9). Finally, estimates of per capita tax-paid cigarette sales provide populationwide rather than individual-based estimates of behaviors; because these estimates are based on tax revenues they may not accurately estimate actual consumption (10).

By monitoring tobacco-related health effects, policy changes, and public attitudes at state and local levels, tobacco-related activities can be evaluated and public health programs can be tailored to local populations. CDC and state health departments are working together to improve state-specific measures of tobacco-related health outcomes, policy interventions, and related activities to improve the prevention and control of tobacco use. In 1999, CDC will provide all states with funding for tobacco-use prevention and control programs. CDC also is collaborating with states that have other sources of funding for activities related to tobacco-use prevention to develop effective public health intervention, surveillance, and evaluation activities.

References

1. CDC. Smoking-attributable mortality and years of potential life lost—United States, 1984. MMWR 1997;46:444–51.

Cigarettes Smoking and Smokeless Tobacco Use — Continued

- 2. McGinnis JM, Foege WH. Actual causes of death in the United States. JAMA 1993;270:2207–12.
- 3. CDC. State and national tobacco control highlights. World-Wide Web site http://www.cdc.gov/nccdphp/osh/statehi/statehi.htm. Accessed November 3, 1998.
- CDC. Addition of prevalence of cigarette smoking as a nationally notifiable condition—June 1996. MMWR 1996;45:537.
- CDC. State and sex-specific prevalence of selected characteristics, Behavioral Risk Factor Surveillance System, 1994 and 1995. MMWR 1997;46(no. SS-3).
- Bureau of the Census, Economics and Statistics Administration, US Department of Commerce. Resident Population of the United States: estimates, by age and sex. World-Wide Web site http://www.census.gov/population/estimates/nation/intfile2-1.txt. Accessed November 3, 1998.
- 7. The Tobacco Institute. The tax burden on tobacco: historical compilation, vol 32. Washington, DC: The Tobacco Institute, 1997.
- 8. CDC. Youth Risk Behavior Surveillance—United States, 1997. MMWR 1998;47(no. SS-3).
- 9. CDC. Cigarette smoking among adults—United States, 1992, and changes in the definition of current cigarette smoking. MMWR 1994;43:342–6.
- 10. Clarke TM. A tax study: cigarette consumption in Washington state. Olympia, Washington: Washington State Department of Health, January 1997.

Nosocomial *Burkholderia cepacia* Infection and Colonization Associated with Intrinsically Contaminated Mouthwash — Arizona, 1998

During August 1996–June 1998, 74 patients at two hospitals in Arizona had cultures positive for *Burkholderia cepacia*. Most isolates were from the respiratory tracts of patients in intensive-care units (ICUs). Because of the large number of *B. cepacia* isolates, personnel at both hospitals requested the Arizona Department of Health Services assist in an investigation. This report summarizes the results of the investigation.

A case of infection or colonization was defined as a positive culture for *B. cepacia* from the respiratory tract of any ICU patient at these hospitals during August 31, 1996– June 12, 1998 (epidemic period). Hospital microbiology records were reviewed to identify all isolates of *B. cepacia* during the pre-epidemic (January 1, 1994–August 30, 1996) and epidemic periods. Case-patient medical records, respiratory therapy procedures, and ICU nursing procedures were reviewed.

A total of 69 patients had positive cultures and had illness that met the case definition, compared with one ICU patient during the pre-epidemic period. Case-patients ranged in age from 17 to 87 years (median: 73 years), and 36 (52%) were male. Casepatients were admitted to the ICU with various diagnoses. None had medical conditions associated with infection with *B. cepacia* (e.g., cystic fibrosis or chronic granulomatous disease). Hospital clinicians identified 33 (48%) case-patients as having infections and 36 case-patients as having *B. cepacia* respiratory tract colonization.

All case-patients had been intubated and mechanically ventilated during their ICU stay. All mechanically ventilated patients had received routine oral care that included swabbing with an alcohol-free mouthwash (Kentron Alcohol Free Mouthwash and Gargle[™], product #711-04, manufactured for Kentron Health Care, Inc., Phoenix Cosmetics, Holbrook, New York). The active ingredient in this product is cetyl pyridium chloride; the formulation does not contain alcohol. This product was produced only during 1994–1995 and was distributed throughout the United States. The extent of use of this product in ICU patients at other hospitals is unknown.

Burkholderia cepacia Infection and Colonization - Continued

Cultures of unopened 4-oz. bottles of the mouthwash grew *B. cepacia, Alcaligenes xylosoxidans*, and *Pseudomonas fluorescens putida* group. *B. cepacia* isolates from case-patients and mouthwash were similar by pulsed-field gel electrophoresis. Other potential reservoirs (e.g., lotion, povidone-iodine solution, water supplies, and a name-brand mouthwash) were culture-negative for *B. cepacia*.

On June 12, the two hospitals discontinued use of the product, and no further respiratory isolates of *B. cepacia* have occurred in their ICU patients. On June 16, the Kentron company initiated a voluntary recall of this product.

Reported by: L Matrician, G Ange, S Burns, L Fanning; C Kioski, G Cage, G Harter, D Reese, D McFall, K Komatsu, R Englund, State Epidemiologist, Arizona Dept of Health Svcs. Investigation Br, Phoenix Resident Post, Food and Drug Administration. Hospital Infections Program, National Center for Infectious Diseases; and an EIS Officer, CDC.

Editorial Note: *B. cepacia* (formerly *Pseudomonas cepacia*) is a motile aerobic gramnegative bacillus commonly found in liquid reservoirs and moist environments. *B. cepacia* is a well-known nosocomial pathogen that is intrinsically resistant to aminoglycosides and first- and second-generation cephalosporins; it is responsible for 0.6% of all ventilator-associated pneumonias (1; CDC, unpublished data, 1994). Numerous outbreaks of *B. cepacia* infection have been reported among cystic fibrosis patients (1–3). In December 1995, a similar outbreak involving *B. cepacia* in respiratory cultures from patients without cystic fibrosis was traced to intrinsically contaminated alcohol-free mouthwash prepared by a different manufacturer (4). An investigation by the Food and Drug Administration (FDA) suggested an association with the deionization procedure of the water used to prepare the product (R. Johnson, FDA, personal communication, 1998).

Potential pathogens may be present in low numbers in many nonsterile products used in hospitals. Mechanically ventilated patients are vulnerable to pathogens in their mouths and upper airways because of their inability to maintain the mucociliary and cough mechanisms that normally protect the lower respiratory tract (5). These outbreaks of *B. cepacia* related to mouthwash highlight the increased risk for respiratory colonization and infection among patients on ventilators. Hospital surveillance and investigation of unusual clusters are crucial to promptly identifying unexpected sources of these pathogens and protecting patients at risk.

Clinicians who detect ventilator-associated pneumonia or respiratory colonization with *B. cepacia* associated with the use of nonalcohol containing mouthwash are encouraged to report such episodes through local and state health departments to CDC's Hospital Infections Program, National Center for Infectious Diseases, telephone (404) 639-6413; fax (404) 639-6459; and to MedWatch, the FDA Medical Products Reporting Program, telephone (800) 332-1088.

References

- Mangram A, Jarvis WR. Nosocomial Burkholderia cepacia outbreaks and pseudo-outbreaks. Infect Control Hospital Epidemiol 1996;17:718–20.
- 2. Ederer GM, Matsen JM. Colonization and infection with *Pseudomonas cepacia*. J Infect Dis 1972;125:613–8.
- 3. Tablan OC, Martone WJ, Jarvis WR. The epidemiology of *Pseudomonas cepacia* in patients with cystic fibrosis. European J Epidemiol 1987;3:336–42.
- 4. Bernstein B, Dineen T, Kehl S, Wilson P, Sohnle P. Outbreak of *Burkholderia cepacia* colonization and infection related to contaminated oral mouthwash [Abstract]. In: Program and

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abstracts of the 34th Annual Meeting of the Infectious Diseases Society of America, New Orleans, Louisiana, September 1996.

 Beck-Sague CM, Sinkowitz RL, Chinn RY, Vargo J, Kaler W, Jarvis WR. Risk factors for ventilator-associated pneumonia in surgical intensive care unit patients. Infect Control Hospital Epidemiol 1996;17:374–6.

Neonatal Tetanus — Montana, 1998

Neonatal tetanus (NT) is a severe, often fatal disease caused by a toxin of *Clostrid-ium tetani*, a ubiquitous spore-forming bacterium found in high concentrations in soil and animal excrements. NT is associated with nonsterile delivery and umbilical cord-care practices for newborns of mothers with antitoxin levels insufficient to protect the newborn by transplacental transfer of maternal antibody. In 1997, NT accounted for an estimated 277,400 deaths worldwide (1) but is rare in the United States. During 1995–1997, of 124 tetanus cases reported in the United States, only one occurred in a neonate (*2,3*). This report summarizes the investigation in March 1998 of an NT case by the Missoula City-County Health Department (MCCHD) and the Montana Department of Health and Human Services (MDHHS). The findings indicated that tetanus in a newborn of an unvaccinated mother occurred after application of nonsterile clay to the umbilical cord.

On March 21, 1998, a 9-day-old newborn, who had no previous medical problems, was taken to a hospital by her parents who reported a 10-hour history of an inability to nurse and difficulty in opening her jaw. Her parents also had noticed a foul-smelling discharge from her umbilical cord during the preceding 1–2 days. No other symptoms were noted by the parents. On admission, the newborn had trismus, increased general muscle tone, and hyperresponsiveness to external stimuli. The umbilical cord was covered with dried clay, which when retracted revealed a foul-smelling yellow-green discharge. Culture from the umbilical cord grew several anaerobic (*C. perfringens, C. sporogenes*) and aerobic (*Staphylococcus, Streptococcus,* and *Bacillus* sp.) bacterial species. NT was diagnosed based on the clinical characteristics.

The newborn was treated with tetanus immune globulin (500 units intramuscularly) and penicillin G (300,000 U/kg/day intravenously) for 10 days. On March 24, she required mechanical ventilation and remained ventilated for 12 days. She was discharged on April 10, with no apparent neurologic sequelae and was developing normally on follow-up at age 7 months.

The mother, a 32-year-old non-Hispanic white woman born in the United States, had never been vaccinated because of her family's philosophic beliefs. She had no complications during her pregnancy and was attended throughout her pregnancy by a licensed "direct-entry" midwife* from her community. The newborn was delivered in a local hospital by cesarean section. While in the hospital, she received standard umbilical cord care with isopropyl alcohol. The newborn was discharged at 3 days of age. For home umbilical cord care, the parents applied a "Health and Beauty Clay" powder provided by the midwife. This clay powder was applied to the umbilical cord up to three times daily with a clean cotton-tipped swab. The family lived in a rural area

^{*}Direct-entry midwives are a group distinct from certified nurse midwives; in Montana, they are licensed to attend women during uncomplicated pregnancies, labor, and postpartum periods.

Neonatal Tetanus — Continued

in a house adjacent to a horse pasture. Although the newborn and her mother stayed primarily indoors, the family's dog often ran between the house and the pasture.

The "Health and Beauty Clay" was a bentonite clay from Death Valley, California. According to the manufacturer, it had been sold for 21 years as a cosmetic product without reported adverse health outcomes. The manufacturing process of the clay did not include sterilization. The clay was shipped in 2-lb. containers, sold by weight in a local store, and dispensed to local midwives in smaller containers. The midwives would further aliquot the clay into 2-oz., presumably clean vials for distribution to their patients. The use of the clay for umbilical cord care was common among local direct-entry midwives because they believed it accelerated drying of the umbilical cord.

On April 9, MCCHD distributed a health-care advisory to more than 60 health-care providers in the area that emphasized the importance of tetanus toxoid vaccination, particularly for pregnant women, and cautioned against using nonsterile products for umbilical cord care. Following this case, use of clay for umbilical cord care was discontinued by midwives in the community. The mother of the case-patient has since been vaccinated with tetanus and diphtheria toxoids (Td), but as of October 1998 has not initiated vaccination for her infant because of concern about potential adverse effects. *Reported by: B Goode, Missoula City-County Health Dept, Missoula; K Caruso, Community Medical Center, Missoula; J Murphy, A Weber, J Burgett, Montana Dept of Health and Human Svcs. Child Vaccine-Preventable Disease Br, Epidemiology and Surveillance Div, National Immunization Program; and an EIS Officer, CDC.*

Editorial Note: In the United States, NT is rare. Tetanus-associated deaths among children aged <1 year, an indicator for NT deaths (most tetanus deaths in this age group are caused by NT), declined from 64.0 per 100,000 population in 1900 to 4.5 by the 1940s. By 1967 in the United States, NT incidence was <0.01 per 1000 live-born infants.[†] This decline is associated with improvements in birth practices and increased levels of population immunity following the initiation of routine tetanus toxoid vaccination since the 1940s. Since 1972, 31 cases of NT have been reported to CDC. Of these cases, only five (16%) mothers had a history of ever having received tetanus toxoid, and only one was known to have received more than one dose.

Factors contributing to this case include the lack of maternal vaccination, the anaerobic conditions and *C. tetani* contamination of the umbilical cord resulting from the application of a nonsterile clay, and the potential exposure to *C. tetani* spores from the nearby horse pasture. The case described in this report is the first since 1984 in an infant of a mother born in the United States and with philosophic objections to vaccination. Since 1984, only two other cases of NT have been reported, both in infants of unvaccinated or inadequately vaccinated mothers born outside of the United States (*3,4*). The case in this report was the first NT case and one of only four tetanus cases reported from Montana since 1965.

Vaccination with tetanus toxoid during pregnancy is safe and effective in preventing NT (5). The Advisory Committee on Immunization Practices recommends giving a booster dose of Td to previously vaccinated pregnant women who have not received a Td vaccination within the preceding 10 years, and unvaccinated or partially vaccinated pregnant women should complete the primary series of three doses of Td (6,7).

To prevent NT cases in the United States, health-care professionals should review and update the vaccination status of childbearing-aged women and particularly those who are pregnant. In addition, targeted education regarding the importance and

[†]Data on NT incidence per 1000 live-born infants were not available until the 1960s.

Neonatal Tetanus — Continued

safety of tetanus vaccination is needed among parents and direct-entry midwifery groups, and parents and health-care providers should avoid applying nonsterile products to the umbilical cord of newborns, including products that create anaerobic conditions. Unless all women giving birth are vaccinated appropriately with tetanus toxoid, even hospital-born infants in the United States are at risk for developing NT, especially if unconventional practices of umbilical cord care are followed.

References

- 1. Expanded Program on Immunization. WHO/EPI Information System. Geneva, Switzerland: World Health Organization, September, 1998; document no. WHO/EPI/GEN/98.10.
- 2. Bardenheier B, Prevots DR, Khetsuriani N, Wharton M. Tetanus surveillance—United States, 1995–1997. In: CDC surveillance summaries (July). MMWR 1998;47(no. SS-2):1–13.
- 3. Craig AS, Reed GW, Mohon RT, et al. Neonatal tetanus in the United States: a sentinel event in the foreign-born. Pediatr Infect Dis J 1997;16:955–9.
- 4. Kumar S, Malecki LM. A case of neonatal tetanus. South Med J 1991;84:396-8.
- Galazka AM. Tetanus: the immunological basis for immunization. Geneva, Switzerland: World Health Organization, Expanded Program on Immunization, 1990; document no. WHO/EPI/GEN/93.13.
- 6. CDC. General recommendations on immunization: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 1994;43(no. RR-1):20–1.
- 7. CDC. Update on adult immunization: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 1991;40(no. RR-12):12–3.

Risk Factors for Short Interpregnancy Interval — Utah, June 1996–June 1997

The Utah Medicaid program provides pregnancy-related coverage for women whose household incomes are \leq 133% of the federal poverty level.* For women who are not otherwise eligible, Medicaid coverage of family-planning and other services ends after the second calendar month following delivery. To assess whether increased access to family-planning services would benefit Medicaid recipients, the interpregnancy intervals (IPIs) of Utah residents whose most recent pregnancies were covered by Medicaid (Medicaid-recipient mothers) were compared with those of all other Utah resident mothers. This report summarizes the results of that study, which indicate that Medicaid-recipient mothers aged \geq 20 years were at increased risk for having short IPIs, and may therefore benefit from extended Medicaid coverage or other means of assuring access to family-planning services.

Data for live-born infants of Utah resident mothers from June 30, 1996, through June 29, 1997, were matched to the Medicaid eligibility database by using the mother's date of birth, last name, first name, and middle initial. IPI was defined as the time between delivery dates of consecutive live-born infants minus the gestational age (1) of the most recent child. A short IPI was defined as an IPI of <12 months; this cutoff was based on a parallel study that showed that IPIs of <12 months were associated with significantly elevated risks for adverse perinatal outcomes (Utah Department of Health, unpublished data, 1998). The risk for having a short IPI was examined

^{*}Poverty statistics are based on definitions developed by the Social Security Administration in 1964 (which subsequently were modified by federal interagency committees in 1969 and 1980) and prescribed by the Office of Management and Budget as the standard to be used by federal agencies for statistical purposes.

Risk Factors for Short Interpregnancy Interval — Continued

in relation to maternal Medicaid status and age, marital status, education, and race/ethnicity. A mother's education was categorized as age-appropriate if she had completed high school or the usual number of grades for her age. The educational status of mothers aged \geq 20 years also was evaluated based on the number of years of school completed (0–11, 12, 13–15, 16, and >16 years).

Of the 42,429 live-born infants of Utah resident mothers from June 30, 1996, through June 29, 1997, 15,810 (37.3%) were ineligible for the study because they were first-born infants. An additional 4773 (11.2%) infants were excluded because the date of the previous delivery of a live-born infant or information for estimating gestational age were missing on the birth certificates.

Of the 21,846 (51.5%) infants eligible for study, 3916 (17.9%) were born after a short IPI (Table 1). Medicaid-recipient mothers (22.9% of all mothers in the study population) were nearly twice as likely as other mothers to have short IPIs. Young maternal age correlated strongly with short IPIs. Short IPIs were more common among unmarried mothers (22.5%) than married mothers (17.4%), and among mothers with less than age-appropriate education (25.2%) than among those with age-appropriate education (17.1%). American Indian/Alaskan Native mothers and Asian/Pacific Islander mothers were more likely to have short IPIs than non-Hispanic mothers. Mothers who had either one or five or more previous live-born infants were more likely to have short IPIs than mothers.

When stratified analyses were performed, the association between Medicaid status and short IPI differed by maternal age. Among mothers aged \geq 20 years at delivery (21,207 [97.1%] of all mothers studied), Medicaid recipients were more likely to have short IPIs than others (relative risk [RR]=1.7, 95% confidence interval [CI]=1.6–1.8). However, among mothers aged 15–19 years (631 [2.9%] of all mothers), no such association was found (RR=1.0, 95% CI=0.9–1.2).

To evaluate risk factors for having short IPIs while simultaneously controlling for other covariates, logistic regression analyses were performed. Among mothers aged ≥20 years (Table 2), Medicaid-recipient mothers were more likely to have short IPIs than other mothers (odds ratio=1.6, 95% CI=1.5–1.8). Mothers from racial/ethnic minority groups had a higher risk for short IPI than white mothers. Married mothers were more likely to have short IPIs than unmarried mothers. The risk for short IPI was inversely correlated with maternal age.

Reported by: J Duncan, B Nangle, Bur of Vital Records; N Streeter, L Bloebaum, Div of Community and Family Health Svcs; DC Tingey, JA Olson, Div of Health Care Financing, Utah Dept of Health. State Br, Div of Applied Public Health Training; Div of Public Health Surveillance and Informatics, Epidemiology Program Office; and an EIS Officer, CDC.

Editorial Note: Short IPIs have been associated with an increased risk for adverse birth outcomes (2–5). Pregnancies that are too closely spaced often are unintended, which can place substantial financial and psychologic burdens on the mother and her family (6). The findings in this study indicate that Medicaid-recipient mothers aged \geq 20 years were more likely to have short IPIs than other mothers of the same age. Young mothers and mothers of racial/ethnic minority groups had increased risk for short IPIs. Utah data were unavailable to evaluate whether Medicaid-recipient mothers tended to have a shorter duration of breastfeeding or greater desire to build their families quickly than other mothers —two factors that may contribute to short IPIs. Utah women whose de-

Risk Factors for Short Interpregnancy Interval - Continued

liveries were covered by Medicaid, most of whom lose Medicaid coverage shortly after delivery, may have less access to family-planning services. Improving these women's access to family-planning services might help them prevent unintended pregnancies, and improve birth outcomes. Access could be improved by removing financial barriers through extending Medicaid coverage for all women after a Medi-

		Shor	rt IPI		
Maternal risk factor	N*	No.	(%)	Relative risk	(95% CI†)
Delivery covered by Medicaid					
Yes No	4,995 16,851	1,348 2,568	(27.0) (15.2)	1.8 referent	(1.7– 1.9)
Maternal age at delivery (vrs)					
15–17	92	53	(57.6)	7.8	(5.5–11.2)
18–19	547	240	(43.9)	6.0	(4.3- 8.3)
20–24	5,024	1,392	(27.7)	3.8	(2.7- 5.2)
25–29	7,814	1,330	(17.0)	2.3	(1.7– 3.2)
30–34	5,536	639	(11.5)	1.6	(1.1– 2.2)
35–39	2,344	226	(9.6)	1.3	(0.9– 1.8)
40–47	489	36	(7.4)	referent	
Education [§] Less than age-					
appropriate	2,248	567	(25.2)	1.5	(1.4– 1.6)
Age-appropriate	19,441	3,319	(17.1)	referent	
Marital status					
Unmarried	2,262	510	(22.5)	1.3	(1.2- 1.4)
Married	19,583	3,406	(17.4)	referent	
Race					
American Indian/					
Alaskan Native	329	105	(31.9)	1.8	(1.6- 2.1)
Asian/Pacific Islander	562	135	(24.0)	1.4	(1.2- 1.6)
Black	113	28	(24.8)	1.4	(1.0- 2.0)
White	20,671	3,615	(17.5)	referent	
Ethnicity					
Hispanic	1,682	402	(23.9)	1.4	(1.3– 1.5)
Non-Hispanic	20,105	3,505	(17.4)	referent	
No. previous live-born infants					
1	9,997	1,930	(19.3)	1.2	(1.1- 1.3)
2	5,903	968	(16.4)	1.0	(0.9- 1.1)
3	3,160	503	(15.9)	referent	
4	1,461	240	(16.4)	1.0	(0.9- 1.2)
≥5	1,309	267	(20.4)	1.3	(1.1– 1.5)
Overall	21,846	3,916	(17.9)		

TABLE 1. Risk factors for short	interpregnancy	interval	(IPI)	(<12	months)	 Utah,
June 30, 1996–June 29, 1997						

*Numbers may not equal overall number because of missing data.

[†]Confidence interval.

[§]A mother's education was considered age-appropriate if she had completed high school or the usual number of grades for her age.

Risk Factors for Short Interpregnancy Interval - Continued

Variable	Odds ratio [†]	(95% Cl [§])	
Delivery covered by Me	dicaid		
Yes	1.6	(1.5–1.8)	
No	referent		
Maternal age at delivery	y (yrs)		
20–24	4.7	(3.3–6.8)	
25–29	2.6	(1.8–3.7)	
30–34	1.7	(1.2 - 2.4)	
35–39	1.4	(0.9–2.0)	
40–47	referent		
Education (vrs)			
0–11	referent		
12	1.0	(0.8–1.1)	
13–15	1.1	(1.0–1.3)	
16	1.1	(0.9–1.3)	
>16	1.4	(1.2–1.7)	
Marital status			
Unmarried	0.7	(0.6–0.8)	
Married	referent		
Bace			
American Indian/			
Alaskan Native	1.9	(1.5–2.5)	
Asian/Pacific Islander	1.6	(1.3–2.0)	
Black	1.6	(1.0-2.5)	
White	referent		
Ethnicity			
Hispanic	1.3	(1.1–1.5)	
Non-Hispanic	referent		

TABLE 2. Logistic regression analysis of risk factors* for short interpregnancy interval (<12 months) among mothers aged 20–47 years (n=20,982) — Utah, June 30, 1996–June 29, 1997

*The number of previous live-born children is not included in the model because of a multicolinearity problem.

[†]Controlling for all of the other variables in the table.

[§]Confidence interval.

caid-covered delivery, or through increasing availability of family-planning services for low-income women. In Utah, 74 publicly funded clinics provide family-planning services to low-income women. However, only 26% of all such women are served; Utah ranks 49th among all states for providing access to family-planning services (7).

The higher risk for having short IPIs among mothers from racial/ethnic minority groups may have been due to cultural or socioeconomic differences or to unequal access to health care. An example of such a factor was observed in one study that found the length of breastfeeding differed by race (*8*).

Because the sociodemographic characteristics of the Utah population are different from those of other states, caution should be used in generalizing the results of this study. In addition, the findings of this study are subject to at least three methodologic limitations. First, Medicaid status was based on the most recent pregnancy. The num-

Risk Factors for Short Interpregnancy Interval — Continued

ber of women in the study population whose Medicaid status had changed since the previous pregnancy was unknown; an implicit assumption has been made that mothers on Medicaid for the most recent pregnancy were on Medicaid for the previous one. Second, this study included women whose Medicaid eligibility was independent of pregnancy and those who were eligible only because they were pregnant and met the program's income requirements. Finally, the retrospective approach of this study may have overestimated the risk for having short IPIs among young women, although such bias is unlikely to account for much of the elevated risk among those women.

In Utah, extending Medicaid coverage of family-planning services and improving use of family-planning programs should be considered to help low-income women prevent unintended pregnancies and improve birth outcomes. Public health programs for preventing short IPIs also should target young mothers regardless of their Medicaid-eligibility status. The analysis described in this report may be repeated to monitor the success of efforts to improve reproductive health of women in Utah.

References

- 1. Ventura SJ, Martin JA, Curtin SC, Mathews TJ. Report of final natality statistics, 1995. Hyattsville, Maryland: US Department of Health and Human Services, CDC, National Center for Health Statistics, 1997. (Monthly vital statistics report, vol 45, no. 11, suppl).
- 2. Brody DJ, Bracken MB. Short interpregnancy interval: a risk factor for low birthweight. Am J Perinatol 1987;4:50–4.
- 3. Erickson JD, Bjerkedal T. Interpregnancy interval. Association with birthweight, still birth, and neonatal death. J Epidemiol Community Health 1978;32:124–30.
- 4. Bakketeig LS, Hoffman HJ, Oakley ART. Perinatal mortality. In: Bracken MB, ed. Perinatal epidemiology. New York, New York: Oxford University Press, 1984.
- 5. Institute of Medicine. Preventing low birthweight. Washington, DC: National Academy Press, 1985.
- 6. Institute of Medicine. The best intentions: unintended pregnancy and the well-being of children and families. Washington, DC: National Academy Press, 1995.
- 7. Alan Guttmacher Institute. Contraception counts: Utah information. New York, New York: The Alan Guttmacher Institute, 1998.
- 8. Statistical Analysis Division, Center for Health Statistics. PRAMS surveillance report: Alabama, 1996. Montgomery, Alabama: Alabama Department of Public Health, 1998.

Notice to Readers

FDA Approval of a Fourth Acellular Pertussis Vaccine for Use Among Infants and Young Children

On July 29, 1998, the Food and Drug Administration (FDA) licensed North American Vaccine, Inc. (Beltsville, Maryland) to distribute a combined diphtheria and tetanus toxoids and acellular pertussis vaccine (DTaP) (CertivaTM *)[†] for the first four doses of

^{*}Diphtheria and Tetanus Toxoids and Acellular Pertussis Vaccine, Adsorbed, Certiva[™], manufactured and distributed by North American Vaccine, Inc., Beltsville, Maryland. Marketed by Ross Products Division, Abbott Laboratories, Inc. The diphtheria and tetanus toxoid components are produced by the Statens Seruminstitut, Copenhagen, Denmark. Final formulation and release of Certiva[™] are conducted by North American Vaccine, Inc.

[†]Use of trade names and commercial sources is for identification only and does not imply endorsement by CDC or U.S. Department of Health and Human Services.

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the diphtheria and tetanus toxoids and pertussis vaccination series administered to infants and children aged 6 weeks–6 years. CertivaTM is the fourth acellular pertussis vaccine to be licensed for use in infants and young children in the United States (1).

Vaccine doses should be administered at ages 2, 4, 6, and 15–20 months. Data are insufficient to evaluate the use of Certiva[™] as a fifth dose among children aged 4–6 years who have received Certiva[™] for the previous four doses. Additional information about the immunogenicity and safety of a fifth dose following four previous doses of the same acellular pertussis vaccine is being collected and should be available before these infants are aged 4–6 years and require a fifth dose.

The Advisory Committee on Immunization Practices (ACIP), the Committee on Infectious Diseases, the American Academy of Pediatrics, and the American Academy of Family Physicians recommend that children routinely receive a series of five doses of vaccine against diphtheria, tetanus, and pertussis before age 7 years (1,2). The first four doses should be administered at ages 2, 4, 6, and 15–18 months, and the fifth dose at age 4–6 years.

The following evidence supports the use of Certiva[™] for the first four doses of the diphtheria, tetanus, and pertussis vaccination series:

- The rates of local reactions, fever, and other common systemic symptoms following receipt of Certiva[™] inoculations were significantly lower than those following whole-cell pertussis vaccination (administered as diphtheria and tetanus toxoids and pertussis vaccine [DTP]) for doses one through three in controlled clinical studies (1,3).
- 2. Efficacy of three doses of Certiva[™] against pertussis disease was assessed in a double-blind, randomized, placebo-controlled trial in Sweden using CertivaTM -EU, a vaccine containing the same amount of pertussis toxoid (40 μ g) per dose as CertivaTM but more diphtheria toxoid (25 Lf versus 15 Lf) and more tetanus toxoid (7 Lf versus 6 Lf) (4). Infants were randomly assigned to be vaccinated with either Certiva[™] -EU or DT (Diphtheria and Tetanus Toxoids Adsorbed Vaccine, Statens Seruminstitut, Copenhagen, Denmark) at ages 3, 5, and 12 months. The main observation period started 30 days after the third dose of vaccine and lasted a mean of 17 months. In this trial, pertussis was defined according to the World Health Organization case definition (i.e., a paroxysmal cough illness lasting \geq 21 days and confirmed by culture, serology, or epidemiologic link to a culture-positive household contact). Starting 1 month after the third dose, the vaccine efficacy of Certiva[™] -EU against WHO-defined pertussis was 72% (95% confidence interval=62%-78%) (3). Although a serologic correlate of protection for pertussis has not been established, the antibody response to pertussis toxoid in U.S. infants after doses of Certiva[™] at 2, 4, 6, and 15–20 months of age was comparable to that achieved in a previous trial among infants in Sweden in whom efficacy was demonstrated after three doses at 3, 5, and 12 months of age.

Because of the reduced frequency of adverse reactions and demonstrated efficacy, the ACIP recommends a licensed DTaP for all five doses of the routine diphtheria, tetanus, and pertussis vaccination series and for the remaining doses in the series for children who have started the vaccination series with whole-cell DTP vaccine (1). The ACIP considers the data to be insufficient in terms of safety and efficacy to express a preference between different acellular pertussis vaccine formulations.

Notice to Readers — Continued

Whenever feasible, the same DTaP vaccine should be used throughout the entire vaccination series. No data exist on the safety, immunogenicity, or efficacy of different DTaP vaccines when administered interchangeably in the primary or booster vaccination of a child. However, if the vaccine provider does not know or have available the type of DTaP vaccine the child to be vaccinated had received previously, any of the licensed DTaP vaccines may be used to complete the vaccination series (1).

References

- CDC. Pertussis vaccination: use of acellular pertussis vaccines among infants and young children: recommendations of the Immunization Practices Advisory Committee (ACIP). MMWR 1997;46(no. RR-7).
- American Academy of Pediatrics. Pertussis. In: Peter G, ed. 1997 Red book: report of the Committee on Infectious Diseases. 24th ed. Elk Grove Village, Illinois: American Academy of Pediatrics 1997:394–407.
- 3. Diphtheria and tetanus toxoids and acellular pertussis vaccine adsorbed (Certiva[™]) [Package insert]. Beltsville, Maryland: North American Vaccine, Inc., 1998.
- 4. Trollfors B, Taranger J, Lagergard T, et al. A placebo-controlled trial of a pertussis-toxoid vaccine. N Engl J Med 1995;333:1045–50.



FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending October 31, 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 October 31, 1998 (43rd Week)

	Cum. 1998		Cum. 1998
Anthrax Brucellosis Cholera Congenital rubella syndrome Cryptosporidiosis* Diphtheria Encephalitis: California* eastern equine* St. Louis* western equine* Hansen Disease Hantavirus pulmonary syndrome* ¹ Hemolytic uremic syndrome, post-diarrheal* HIV infection, pediatric* [§]	47 7 3 2,723 1 79 4 22 - 92 19 71 178	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	7 1 39 278 1,813 44 351 35 110 11 278 -

-: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 from reports to the Division of HIV/AIDS Prevention–Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP), last update September 27, 1998. [¶] Updated from reports to the Division of STD Prevention, NCHSTP.

					Esche	erichia			Honatitic		
	All	DS	Chlar	nydia	NETSS [†]	PHLIS [§]	Gono	rrhea	Hepa C/NA	ititis A,NB	
Reporting Area	Cum. 1998*	Cum. 1997	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1998	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1997	
UNITED STATES	35,486	47,056	443,269	388,515	2,513	1,681	268,692	245,076	3,379	2,912	
NEW ENGLAND	1,381	2,059	15,480	14,879	292	238	4,422	4,965	81	49	
Nane N.H.	24 28	50 34	850 779	830 675	33 41	42	59 78	59 79	-	-	
Vt.	17	32	350	349	19	17	33	44	- 70	3	
R.I.	94	119	1,910	1,669	130	130	320	378	3	39 7	
Conn.	506	1,097	4,614	5,324	52	40	2,065	2,648	-	-	
Upstate N.Y.	9,642 1,102	13,920 2,134	50,259 N	47,355 N	248 189	/0	29,829 5,334	31,407 5,372	303 239	262 191	
N.Y. City	5,457	7,287	28,897	22,597	7	12	12,939	11,767	-	-	
Pa.	1,765	2,892	13,086	16,458	52 N	48 10	5,312 6,244	6,302 7,966	64	71	
E.N. CENTRAL	2,567	3,637	72,052	52,489	393	278	51,990	33,630	432	472	
Ohio Ind.	540 414	722 459	21,208 4.656	18,508 7.664	106 90	60 43	13,831 3.924	12,090 5.062	77	17 12	
III.	993	1,514	20,054	U	96	39	16,975	U	31	80	
Wis.	468 152	25	8,454	9,115	N	62 74	3,670	3,981	387	338	
W.N. CENTRAL	664	984	24,711	26,947	469	345	12,571	11,833	261	52	
Minn. Iowa	136 58	156 85	5,183 2 063	5,514 3 753	222 89	181 49	1,980 660	1,932 964	9 8	3 25	
Mo.	312	506	10,049	9,973	40	56	7,313	6,089	238	10	
N. Dak. S. Dak.	4 13	10 8	616 1,286	708 1 <i>.</i> 107	10 29	15 31	51 192	58 124	-	2	
Nebr.	59	83	1,520	2,208	53	-	510	973	4	2	
Kans.	82 9 235	130	3,994 91 877	3,684	26	13	75 979	76.082	2 151	204	
Del.	112	194	2,148	14	-	2	1,249	1,022	-	- 204	
Md. D.C.	1,304 691	1,714 879	6,142 N	5,932 N	29 1	12	7,939 3.010	9,611 3,686	9	7	
Va.	688	965	10,950	9,469	Ň	42	7,482	6,921	11	24	
vv. va. N.C.	70 638	105 681	2,123 18,084	2,388 14,354	8 52	6 46	669 15,764	/67 14,260	6 19	16 43	
S.C.	604	644	13,833	10,426	11	8	8,720	9,599	5	35	
Fla.	4,156	5,218	19,798	21,613	43	24	15,263	15,273	92	79	
E.S. CENTRAL	1,444	1,677	32,373	28,911	104	39	31,857	28,957	173	302	
Ky. Tenn.	222 522	319 656	5,264 11,213	5,230 10,426	30 49	- 33	3,069 9,755	3,390 9,099	19 147	12 203	
Ala.	395	453	8,513	7,081	22	2	10,924	9,906	5	10	
WISS.	305 4 202	249 5 156	7,303	58 880	3 104	4 23	39 948	0,002 38 053	2 360	427	
Ark.	159	193	3,073	2,429	10	10	3,165	4,109	9	13	
La. Okla.	708 238	879 255	12,440 8,112	7,987 6,171	5 13	7	10,576 4,458	7,800 4,006	85 12	186 7	
Tex.	3,097	3,829	42,210	42,293	76	-	21,749	22,138	254	221	
MOUNTAIN Mont	1,230	1,365	26,570	24,336	298 15	208	7,511	6,607	307	266	
Idaho	19	48	1,674	1,340	36	22	145	123	87	59	
Wyo. Colo	1 230	13 315	583 6.685	492 5.966	53 73	54 57	27 1.931	44 1.863	60 28	65 29	
N. Mex.	179	146	2,965	3,123	17	13	723	717	83	49	
Ariz. Utah	499 101	318 122	9,426 1,779	8,586 1,432	21 72	26 21	3,391 187	2,872 229	8 23	25 4	
Nev.	178	367	2,415	2,432	11	15	1,075	709	11	14	
PACIFIC Wash	5,121	6,591 528	64,115 9 027	57,727 7 547	395 89	340 104	14,585 1 615	13,542 1 602	1,311 21	878 23	
Oreg.	138	249	4,930	4,007	96	95	695	605	5	3	
Calif. Alaska	4,500 17	5,688 43	46,624 1.538	43,427 1.270	204 6	127	11,606 257	10,596 316	1,230 1	705	
Hawaii	131	83	1,996	1,476	Ň	14	412	423	54	147	
Guam PB	-	2 1 5 10	201	193	N	-	24	27 /90	-	-	
V.I.	24	84	N	N	N	Ŭ	U	403 U	U	Ų	
Amer. Samoa C.N.M.I.	-	- 1	U N	U N	N N	U U	U 28	U 20	U -	U 2	

TABLE II. Provisional cases of selected notifiable diseases, United States,
weeks ending October 31, 1998, and October 25, 1997 (43rd Week)

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

*Updated monthly from reports to the Division of HIV/AIDS Prevention–Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention, last update September 27, 1998. [†]National Electronic Telecommunications System for Surveillance. [§]Public Health Laboratory Information System.

	Legion	ellosis	Lyme Disease		Ma	laria	Syp (Primary &	hilis Secondary)	Tubero	ulosis	Rabies, Animal
Reporting Area	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1997	Cum. 1998*	Cum. 1997	Cum. 1998
UNITED STATES	1,029	848	10,199	10,258	1,111	1,531	5,748	7,112	11,885	14,649	5,751
NEW ENGLAND Maine N.H. Vt. Mass. R.I.	69 1 5 28 19	73 3 7 12 25 9	2,498 11 39 9 713 561	2,720 8 33 8 275 356	53 5 1 16 8	76 1 8 2 29 5	64 1 2 4 38 1	116 - - 58 2	371 10 9 2 203 51	364 18 13 5 202 31	1,232 186 72 57 438 83
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	215 76 25 11 103	17 172 48 18 23 83	1,165 6,387 3,563 25 1,420 1,379	2,040 5,895 2,450 156 1,696 1,593	18 270 83 119 44 24	31 451 64 282 81 24	18 225 34 62 71 58	56 335 31 74 135 95	96 2,335 300 1,243 498 294	95 2,571 350 1,305 536 380	396 1,308 931 U 177 200
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	339 113 99 27 69 31	275 100 45 28 67 35	136 74 54 7 1 U	528 36 29 13 24 426	110 14 11 35 43 7	144 17 16 57 39 15	876 119 202 344 160 51	543 184 153 U 111 95	1,036 83 100 524 311 18	1,461 229 128 756 258 90	123 54 9 15 35 10
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans	67 6 10 22 - 3 19 7	50 9 15 2 15 5	180 148 21 - - 3 6	117 88 5 17 - 1 2 4	84 50 8 15 2 - 1 8	46 19 9 3 1 1	107 8 - 81 - 1 4 13	154 16 7 102 - 3 26	330 124 40 91 8 16 18 33	461 120 46 189 10 10 20 66	600 106 136 24 122 130 7 75
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Ela	123 12 26 6 18 N 11 10 8 30	103 10 18 4 22 N 13 7 1 28	737 34 526 4 58 11 49 5 5 45	678 109 440 7 53 8 32 2 1 26	266 3 75 16 49 2 23 6 34 58	271 5 77 18 64 - 16 16 30 45	2,116 20 551 66 125 2 618 240 236 258	2,896 17 780 95 208 3 773 318 452 250	1,662 18 243 85 222 32 365 207 420 70	2,787 28 256 80 275 47 346 278 495 982	1,664 17 397 490 65 136 121 259 179
E.S. CENTRAL Ky. Tenn. Ala. Miss.	58 24 22 5 7	47 11 26 3 7	79 21 41 16 1	83 15 37 10 21	27 4 15 6 2	34 12 7 10 5	1,023 88 480 239 216	1,471 116 638 369 348	893 136 289 302 166	1,072 155 369 344 204	239 29 122 86 2
W.S. CENTRAL Ark. La. Okla. Tex.	39 - 3 12 24	29 1 6 2 20	23 6 4 2 11	81 24 3 20 34	26 1 13 4 8	48 5 12 6 25	890 95 363 107 325	1,151 133 308 108 602	1,782 120 200 140 1,322	2,083 154 185 171 1,573	131 31 100 -
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	64 2 1 16 2 17 21 3	56 1 2 1 18 3 12 12 7	17 5 1 5 4 - 2	11 3 2 - 1 2 1 2	50 1 8 19 12 8 1 1	62 2 27 8 11 3 9	205 2 1 11 22 155 4 10	153 - 13 8 116 5 10	352 18 12 4 U 56 155 46 61	474 6 10 2 71 57 207 26 95	193 47 55 39 5 19 26 2
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	55 9 - 44 1 1	43 7 35 1	142 7 20 114 1	145 8 17 118 2	225 17 16 187 2 3	399 19 21 347 3 9	242 27 5 208 1 1	293 9 9 273 1 1	3,124 177 118 2,653 42 134	3,376 246 123 2,798 63 146	261 7 231 23
Guam P.R. V.I. Amer. Samoa C.N.M.I.	2 - U U	- - U -	U U U	U U U	1 - U U -	5 U U	1 156 U U 164	3 212 U U 10	36 68 U U 77	13 164 U U 7	47 U U

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States,weeks ending October 31, 1998, and October 25, 1997 (43rd Week)

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

	H. influ	ienzae,	Hepatitis (Viral), by type						Measl	es (Rubec	ola)	
	inva	sive		A	E	3	Indi	ndigenous		orted [†]	То	tal
Reporting Area	Cum. 1998*	Cum. 1997	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1997	1998	Cum. 1998	1998	Cum. 1998	Cum. 1998	Cum. 1997
UNITED STATES	867	895	18,137	23,620	7,204	7,838	2	58	-	21	79	127
NEW ENGLAND	59	52	221	580	152	148	-	1	-	2	3	19
Maine N.H.	3	5 9	17 10	53 31	2 16	6 15	-	-	-	-	-	1
Vt.	7	3	14	12	4	8	-	-	-	1	1	-
Mass. R.I.	34 5	2	90 14	126	47 58	62 14	-	-	-	-	2	16
Conn.	1	2	76	121	25	43	-	-	-	-	-	1
MID. ATLANTIC	122	141	1,189	1,763	881	1,124	-	8	-	5	13	26
N.Y. City	26	38	310	789	235	406	-	-	-	-	-	10
N.J. Pa	42 5	41 17	278 295	258 426	168 238	204 268	U	7	U	1	8	3
E.N. CENTRAL	143	144	2,958	2,471	1.295	1,234	-	11	-	3	14	10
Ohio	45	76	265	269	68	65	-	-	-	1	1	-
Ind. III.	36 49	14 36	291 547	257 686	660 151	88 234	2	2	-	1	3	- 7
Mich.	7	17	1,713	1,095	387	360	-	9		1	10	2
WN CENTRAL	79	1 15	142	1 8/3	29	407 396	0	-	0	-	-	1 17
Minn.	62	33	112	166	42	35	-	-	-	-	-	8
lowa Mo	2	5 4	384 543	394 944	58 210	32 283	- U	1	- U	-	1	- 1
N. Dak.	-	-	3	10	4	5	-	-	-	-	-	-
S. Dak. Nebr.	- 1	2 1	30 38	19 76	2 14	1 13	-	-	-	-	-	8
Kans.	6	-	87	234	22	27	U	-	U	-	-	-
S. ATLANTIC	176	134	1,638	1,579	968	1,022	-	3	-	5	8	13
Md.	49	48	272	169	135	142	-	-	-	1	1	2
D.C. Va	- 16	- 12	53 181	21 195	11 88	28 106	-	-	-	- 2	- 2	1
W. Va.	5	3	6	10	8	14	-	-	-	-	-	-
N.C. S.C.	23	21	35	95	35	215 88	-	-	-	-	-	2
Ga.	41	26	513	424	128	110	-	1	-	1	2	1
ES CENTRAL	48	20 48	325	400 524	346	583	_	-	_	2	2	1
Ky.	7	7	20	66	39	34	-	-	-	-	-	-
Ienn. Ala.	27 12	27 12	199 63	323 74	238 67	373 60	-	-	2	1	1	- 1
Miss.	2	2	43	61	2	116	U	-	U	-	-	-
W.S. CENTRAL	51	42	3,372	4,862	1,085	1,085	-	1	-	-	1	8
La.	22	11	94	194	128	131	-	1	-	-	1	-
Okla. Tex	26	27	513 2 676	1,251 3 227	71 801	40 841	-	-	-	-	-	1 7
MOUNTAIN	92	73	2,737	3,625	693	735	1	1	-	-	1	8
Mont.	-	-	88	65	5	9	U	-	U	-	-	-
Wyo.	1	4	35	28	30	22	-	-	-	-	-	-
Colo.	18	13	282	347 201	99 287	131	-	-	-	-	-	-
Ariz.	53	29	1,714	1,895	158	172	1	1	-	-	1	5
Utah Nev	5 7	3 16	171 94	498 374	63 35	77 66	- U	-	- U	-	-	1
PACIFIC	, 97	216	4,500	6,373	1.432	1.511	1	32	-	4	36	25
Wash.	9	5	848	548	99	60	-	-	-	1	1	2
Calif.	44	167	3,282	5,344	1,211	1,331	-	5	-	2	7	19
Alaska Hawaii	1 7	8 7	16 36	27 135	12 6	12 10	1	27	-	1	28	- 4
Guam	-	-	-	-	2	3	U	-	U	-	-	-
P.R.	2		49	244	323	663	-	-	-	-	-	-
v.ı. Amer. Samoa	U U	U U	U U	U U	U U	U U	U	U	U	U	U U	U
C.N.M.I.	-	6	3	1	53	41	U	-	U	-	-	1

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination,
United States, weeks ending October 31, 1998,
and October 25, 1997 (43rd Week)

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

 * Of 204 cases among children aged <5 years, serotype was reported for 101 and of those, 39 were type b.

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

	Mening Dise	jococcal ease		Mumps			Pertussis		Rubella			
Reporting Area	Cum. 1998	Cum. 1997	1998	Cum. 1998	Cum. 1997	1998	Cum. 1998	Cum. 1997	1998	Cum. 1998	Cum. 1997	
UNITED STATES	2,200	2,698	7	397	524	112	4,894	4,449	-	323	157	
NEW ENGLAND	94	172	-	7	8	10	778	796	-	38	1	
Maine N H	6 4	17 13	-	-	-	-	5 95	14 113	-	-	-	
Vt.	5	4	-	-	-	2	67	204	-	-	-	
Mass. B I	47	84 18	-	4	2	8	563 9	423	-	8	1	
Conn.	25	36	-	2	1	-	39	26	-	29	-	
MID. ATLANTIC	193	281	-	21	49	6	459	329	-	130	33	
N.Y. City	20	46	-	6 4	3	- -	266	60	-	14	28	
N.J.	50	58 105	U	2	7	U	5	13	U	4	-	
F N CENTRAL	328	411	5	69	20 65	29	530	481	-	-	- 6	
Ohio	124	146	1	27	25	14	246	140	-	-	-	
Ind. III.	57 82	45 125	- 1	6 11	9 10	5 10	118 90	50 69	-	-	- 2	
Mich.	37	59	3	25	17	-	59	51		-	-	
WIS.	28	36	U	-	4	U	1/	1/1	U	-	4	
Minn.	29	34	-	12	14 5	6 5	466 275	221	-	- 27	-	
lowa Mo	38	40	-	10	7	1	66	46		- 2	-	
N. Dak.	5	2	-	2	-	-	2	1	-	-	-	
S. Dak. Nebr	7 11	5 10	-	-	- 1	-	8 17	4	-	-	-	
Kans.	27	21	U	-	1	U	66	, 21	U	25	-	
S. ATLANTIC	382	459	1	45	61	3	276	377	-	19	78	
Del. Md.	2	5 41	-	-	-	- 1	5 51	107	-	- 1	-	
D.C.	1	8	-	-	- 10	-	1	3	-	-	1	
W. Va.	13	40	-	o -	-	-	30 1	42	-	-	-	
N.C.	52 51	84 49	-	10	10 10	1	90 25	109 25	-	13	59 15	
Ga.	84	90	-	1	10	-	24	13	-	-	-	
Fla.	117	118	-	20	20	-	49	71	-	4	2	
E.S. CENTRAL Ky.	207	204 42	-	14	26	-	45	125 56	-	2	-	
Tenn.	65	68	-	1	5	1	33	34	-	2	- 1	
Miss.	24	24	Ū	5 5	10	Ū	20	10	Ū	-	-	
W.S. CENTRAL	267	264	1	57	73	13	330	230	-	88	4	
Ark. La.	28 56	31 47	1	11 10	1 12	9	82 7	39 18	-	-	-	
Okla.	38	36	-	-	-	-	29	31	-	-	-	
	145	150	-	30	60 54	4 25	212	142	-	88	4	
Mont.	4	8	Ū	- 52	- 54	35 U	912	16	Ū	-	-	
Idaho Wyo.	10 5	10 3	-	4	3	-	240 8	500 7	-	-	2	
Colo.	26	42	-	6	3	6	190	299	-	-	-	
N. Mex. Ariz.	24 41	24 39	N -	N 6	N 32	1 12	87 198	88 35	-	1	- 5	
Utah	13	12		5	8	16	146	17		2	-	
Nev.	0 //15	556	0	10	/ 17/	9	34 1 036	21 769	0	1/1	- 27	
Wash.	57	74	-	9	18	5	275	322	-	9	5	
Oreg. Calif.	75 275	105 368	N	N 92	N 123	4	96 636	39 374	-	- 3	14	
Alaska	3	2	-	2	8	-	14	16	-	-	-	
Hawaii	5	/	-	22	25	-	15	18	-	2	8	
P.R.	6	8	-	∠ 1	7	-	3	-	-	-	-	
V.I. Amer Samoa	U	U	U	U	U	U	U	U	U	U	U	
C.N.M.I.	-	-	Ŭ	2	4	Ŭ	1	-	Ŭ	-	-	

TABLE III. (Cont'd.) Provisional cases of selected notifiable diseases preventable
by vaccination, United States, weeks ending October 31, 1998,
and October 25, 1997 (43rd Week)

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

	All Causes, By Age (Years)						P&I [†]			All Cau	ises, By	y Age (Years)			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 Haven, Conn. Providence, R.I. Somerville, Mass. Springfield, Mass.	529 140 37 12 23 42 17 11 11 28 49 42 6 37 26	385 92 29 8 17 27 16 8 22 36 32 5 27 27	92 29 4 2 5 8 1 3 5 9 5 - 10 3	34 11 4 2 - 3 - 1 4 3 1 - 1	15 6 - 1 3 - - 2 - 2	32	42 15 4 - 2 3 2 3 - 6 2 3	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.	1,265 128 264 100 122 45 58 39 66 227 101 13	779 77 154 64 74 60 28 36 25 55 55 147 50 9	277 23 63 18 28 25 10 14 12 7 47 30	138 18 34 15 11 10 3 4 1 4 19 15 4	36 5 4 1 5 5 3 3 1 - 6 3 -	35 5 9 2 4 2 1 1 - 8 3 -	74 3 21 10 4 - 3 5 6 19 3 2
Marchard, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Erie, Pa.	20 59 2,191 43 32 U 26 6 53	44 1,574 32 30 U 17 2 46	8 416 8 2 U 4 3 7	4 132 2 U 2 1	3 38 1 - U -	31 - - - - - - - - - - -	97 2 - U - 3	E.S. CENTRAL Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala. Nashville, Tenn.	910 158 94 100 49 205 94 77 133	617 111 66 76 33 129 63 55 84	179 32 19 18 8 42 19 10 31	69 4 6 5 19 10 9	26 8 2 1 8 1 3 3	19 3 1 2 7 1 5	44 5 17 2 6 2 3 3
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.	38 1,181 U 12 400 48 19 131 19 37 84 45 45 17 U	28 819 U 273 40 19 105 105 32 62 34 15 U	6 239 U 4 86 7 - 20 3 4 16 6 1 U	2 87 0 28 - 3 - 1 3 - 1 3 - U	1 25 U - 4 - 1 - 1 2 2 1 U	1 11 1 9 1 - 2 - 3 - 3 - U	48 U 21 4 2 5 4 5 3 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.	1,468 70 54 180 50 124 381 76 75 224 87 98	934 48 36 41 97 37 82 221 40 42 169 58 63	316 11 12 4 41 8 24 96 23 21 36 18 22	130 6 4 3 24 2 12 43 7 5 10 8 6	41 2 9 1 2 10 4 5 2	47 3 1 9 2 4 11 2 3 4 3 5	82 3 1 6 4 8 25 7 15 7 6
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind.	1,620 51 43 U 115 123 199 118 190 41 62	1,154 43 30 U 79 85 148 78 112 34 46	284 5 8 U 19 31 30 48 6 13	114 2 4 U 7 10 16 6 21 2	42 1 U 4 4 1 6 1	26 - - - - - - - - - - - - - - - - - - -	93 2 14 14 12 4 1 4	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.	890 77 31 . 73 108 145 30 148 31 121 126	624 60 27 53 73 95 21 97 26 82 90	166 12 14 20 33 6 28 4 26 22	69 4 2 5 10 11 2 16 1 8 10	16 - 1 4 - 3 - 4 3	14 1 - 4 2 1 4 - 1	63 4 2 1 7 8 3 13 - 19 6
Gary, Ind. Grand Rapids, Mich Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio	8 194 49 124 41 52 43 101 U	7 51 123 36 92 28 42 38 82 U	8 39 20 7 6 4 12 U	1 21 2 6 4 2 1 5 U	- 9 1 5 1 2 - 1 U	3 2 1 1 - 1 U	27403227U	PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawaii Long Beach, Calif. Los Angeles, Calif. Pasadena, Calif. Portland, Oreg. Sacramento, Calif.	1,761 19 108 23 61 89 457 22 111 155	1,246 13 77 20 48 61 323 20 89 104	328 3 19 - 8 16 87 2 15 30	123 1 9 1 2 6 36 5 17	30 1 1 2 3 7 - 3	33 1 2 1 3 4 - 2 1	137 5 2 7 12 21 6 12 9
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.	742 31 25 42 115 43 150 90 94 67 85	538 22 29 76 34 116 58 63 53 65	114 9 2 4 21 7 20 16 17 7 11	43 165196735	21 3 4 1 3 1 4 2 3	18 - - 1 - 2 9 3 2 1	50 6 - 8 5 9 6 - 5 1	San Diego, Calif. San Francisco, Calif San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash. TOTAL	134 150 160 22 113 55 82 11,376 [¶]	88 103 112 16 72 42 58 7,851	25 34 5 28 10 15 2,172	13 9 10 9 1 4 852	3 1 3 1 1 265	4 3 4 1 1 4 226	14 19 11 6 6 6 682

TABLE IV. Deaths in 122 U.S. cities,* week ending October 31, 1998 (43rd 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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