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National Cholesterol Education Month — September 2001

High blood cholesterol is a major risk factor for heart disease, the leading cause of death in the United States. Lowering cholesterol levels reduces the incidence of heart disease and death among persons with or without coronary heart disease. To increase awareness of the importance of monitoring cholesterol levels and taking steps to achieve or maintain healthy levels, the National Cholesterol Education Program (NCEP) sponsors National Cholesterol Education Month every September. This year, the theme is "Know your cholesterol numbers; know your risk."

In May 2001, NCEP released the *Third Report of the Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults, Adult Treatment Panel III* (ATP III), which recommends that adults aged ≥20 years have their cholesterol checked at least once every 5 years. Cholesterol levels can be lowered through lifestyle changes such as dietary improvement, increased physical activity, weight control, drug therapy, or a combination of these (1).

During September, CDC-funded state cardiovascular health programs and their collaborators will conduct programs aimed at increasing awareness and understanding of high blood cholesterol and its impact on heart disease. For example, the Montana state health department and Blue Cross Blue Shield of Montana have developed and broadcast radio public service announcements providing cholesterol education. The Arkansas state health department will provide cholesterol educational information sheets to the public and health-care professionals.

Additional information about how cholesterol may affect health and about the new ATP III guidelines is available at http://www.americanheart.org/cld>, and http://www.cdc.gov/nccdphp/cvd.

References

 National Cholesterol Education Program Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults. Executive summary of the third report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (ATP III). JAMA 2001;285:2486–97.

^{*}Reference to non-CDC sites on the Internet are provided as a service to *MMWR* readers and do not constitute or imply endorsement of these organizations or their programs by CDC or the U.S. Department of Health and Human Services. CDC is not responsible for the content of pages found at these sites.

State-Specific Trends in High Blood Cholesterol Awareness Among Persons Screened — United States, 1991–1999

High blood cholesterol (HBC) is a major risk factor for heart disease. One of the national health objectives for 2010 is to reduce the percentage of adults aged ≥20 years with total blood cholesterol levels of ≥240 mg/dL (objective 12–14) (1). One strategy for achieving this objective is to increase awareness of HBC. State-specific data allow state health departments to monitor progress in educating the public about awareness of cholesterol levels and the need for persons to maintain low levels of blood cholesterol. To examine state-specific trends in the proportion of screened adults who reported that they were told that they had HBC, CDC analyzed data from the Behavioral Risk Factor Surveillance System (BRFSS) for 1991 through 1999. This report summarizes the results of that analysis and indicates that approximately one fourth of screened survey participants were aware that they had HBC; this proportion increased slightly from 1991 through 1999. Awareness of HBC is a necessary step to help persons take action to lower their cholesterol level and their risk for coronary heart disease.

BRFSS is a random-digit–dialed telephone survey of the noninstitutionalized U.S. population aged ≥18 years. For this report, BRFSS data from 1991, 1993, 1995, 1997, and 1999 were analyzed for 412,322 persons aged ≥20 years from 50 states and the District of Columbia (DC). Survey participants were asked whether they had ever had their blood cholesterol checked and, if so, had a physician or other health-care provider ever told them their blood cholesterol was high. Those who reported having ever had their blood cholesterol checked were included in the analysis and those who reported they had been told they had HBC were classified as being aware they had HBC (n=120,450). Data were weighted to account for the age, race/ethnicity, and sex distribution and nonresponse in each state. Analyses were conducted using SUDAAN 7.0 to account for the complex sampling design and to obtain accurate variance estimates. To allow for comparisons between states, the results were age-standardized with the direct method using the U.S. 2000 standard population (2). Participation rates in BRFSS ranged from 71.4% in 1993 to 55.2% in 1999. The prevalence of cholesterol screening during the preceding 5 years increased from 67.3% in 1991 to 70.8% in 1999 (3).

Among all 50 states and DC that participated in BRFSS during 1999, the agestandardized prevalence of persons screened who were ever told that they had HBC ranged from 20.5% in Oklahoma to 33.7% in Nevada (Table 1). For the 47 states that participated in BRFSS in all years from 1991 through 1999, the age-standardized prevalence of HBC awareness among persons screened increased from 25.7% in 1991 to 28.6% in 1999 (Table 1). The age-standardized prevalence of HBC awareness among persons screened increased in DC and 38 states and ranged from a 0.1 percentage point increase in Delaware to a 7.3 percentage point increase in Florida. The increase in HBC awareness was significant in Alabama, Arkansas, California, Florida, Georgia, Iowa, Maryland, Minnesota, Mississippi, Missouri, New York, North Carolina, Ohio, South Dakota, Tennessee, Texas, and West Virginia. For eight states (Alaska, Arizona, Connecticut, Hawaii, Oklahoma, Rhode Island, South Carolina, and Vermont), the prevalence of persons screened who reported HBC decreased from 1991 to 1999 and ranged from a 5.8 percentage point decline in Oklahoma to a 0.7 percentage point decline in Connecticut. The decrease was significant in Oklahoma. In Virginia, the prevalence of reported HBC among persons who ever had their cholesterol tested remained constant at 31.0% during 1991-1999.

TABLE 1. Prevalence of screened persons who were ever told they had high blood cholesterol, by reporting area — Behavioral Risk Factor Surveillance System, United States, 1991-1999

Officed States, 1.						% point change	
Area	1991*	1993†	1995⁵	1997 [¶]	1999**	1991 to 1999	(95% CI ^{††})
Alabama	25.3	27.6	26.1	27.9	31.3	5.9	(2.6- 9.3)
Alaska	32.9	30.4	28.6	27.5	29.3	-3.6	(-7.5- 0.3)
Arizona	26.4	24.2	26.5	30.7	23.4	-3.0	(-6.6- 0.6)
Arkansas	25.0	27.5	26.7	28.7	29.7	4.7	(1.2- 8.3)
California	25.7	28.4	27.8	29.3	28.9	3.3	(0.8- 5.8)
Colorado	24.9	26.7	28.4	27.8	25.1	0.1	(-2.9- 3.2)
Connecticut	27.4	29.0	24.8	23.2	26.7	-0.7	(-3.7- 2.3)
Delaware	29.6	29.2	29.4	27.8	29.7	0.1	(-3.4- 3.6)
District of Columbia	20.3	18.2	NA	18.2	22.1	1.8	(-1.8- 5.4)
Florida	22.8	30.4	28.6	29.8	30.1	7.3	(4.8– 9.8)
Georgia	23.4	26.9	22.5	24.3	28.7	5.3	(2.0- 8.6)
Hawaii	29.7	33.2	26.7	30.0	26.7	-3.0	(-6.4- 0.3)
Idaho	25.3	29.1	26.7	28.4	28.1	-3.0 2.8	(-0.4- 0.3) (-0.2- 5.7)
Illinois	25.3 27.1	27.8	26.6	31.7	29.6	2.5	(-0.2- 5.7) (-1.2- 6.3)
Indiana	27.1	30.3	29.6	27.6	30.3	3.1	(-0.1- 6.3)
			27.4			4.2	
lowa	24.3	28.2		26.5	28.5		(1.0– 7.5)
Kansas	NA 29.5	31.5	31.0 29.1	26.1	25.8	NA 1.7	(-1.2- 4.5)
Kentucky		31.3		29.0	31.1	1.7	
Louisiana	25.4	26.6	25.9	26.4	26.2	0.8	(-2.9- 4.5)
Maine	26.3	27.4	28.9	31.1	29.9	3.6	(-0.3- 7.5)
Maryland	24.8	26.6	25.2	28.3	29.5	4.7	(1.8– 7.7)
Massachusetts	26.5	27.9	31.0	24.6	28.8	2.3	(-0.9- 5.5)
Michigan	30.6	29.6	30.8	30.0	31.2	0.6	(-2.4- 3.7)
Minnesota	24.8	26.8	26.8	29.5	29.4	4.6	(2.3– 6.9)
Mississippi	25.1	31.0	23.9	27.5	29.4	4.3	(0.7– 7.9)
Missouri	24.7	30.8	27.8	28.8	28.4	3.8	(0.6– 6.9)
Montana	27.5	25.8	27.2	29.2	28.1	0.6	(–3.4– 4.7)
Nebraska	23.9	26.2	26.9	28.6	25.7	1.8	(-1.6- 5.2)
Nevada	NA	31.6	28.8	26.7	33.7	NA	
New Hampshire	29.5	29.3	26.8	30.6	32.0	2.5	(–1.5– 6.6)
New Jersey	24.7	27.8	24.9	27.3	25.4	0.7	(-2.5- 3.8)
New Mexico	22.3	28.8	28.3	27.1	25.7	3.4	(-0.2- 7.0)
New York	24.1	28.4	25.6	26.8	27.8	3.7	(0.6– 6.7)
North Carolina	24.9	25.7	23.9	25.7	30.2	5.4	(2.2– 8.5)
North Dakota	26.0	30.6	28.5	28.0	28.3	2.3	(-1.1- 5.8)
Ohio	23.4	27.0	27.3	26.0	31.0	7.6	(3.7–11.4)
Oklahoma	26.4	27.9	27.2	21.7	20.5	-5.8	(-9.1-[-2.6])
Oregon	26.1	28.5	27.9	30.3	26.5	0.4	(-2.6- 3.4)
Pennsylvania	25.6	26.2	28.8	24.3	26.1	0.5	(-2.2- 3.2)
Rhode Island	28.1	26.8	27.1	27.3	27.3	-0.8	(-3.7- 2.2)
South Carolina	27.2	27.1	26.5	23.9	26.3	-0.9	(-3.8- 2.1)
South Dakota	24.1	25.7	23.2	24.3	27.1	3.0	(0.1- 6.0)
Tennessee	24.0	28.4	26.7	28.9	28.1	4.1	(1.4– 6.8)
Texas	26.5	28.7	33.4	28.4	29.7	3.3	(0.1– 6.5)
Utah	24.8	28.2	23.0	26.7	27.7	2.9	(-0.3- 6.1)
Vermont	28.7	26.1	27.5	24.8	26.2	-2.5	(-5.7- 0.8)
Virginia	31.0	27.3	29.1	29.3	31.0	0.0	(-3.4- 3.3)
Washington	26.5	28.8	28.7	24.7	26.8	0.4	(-2.5- 3.2)
West Virginia	29.6	32.0	29.7	29.8	34.2	4.6	(1.5- 7.7)
Wisconsin	26.4	31.4	28.9	25.0	29.4	3.0	(-0.7- 6.7)
Wyoming	NA	NA	27.2	29.0	29.5	NA	, 5.7 5.77
Total ^{§§}	25.7	28.3	27.8	27.6	28.6		(2.5- 3.4)
* Sample sizes for ind							

^{*} Sample sizes for individual states ranged from 686 to 2387 adults aged ≥20 years who had their cholesterol screened in 1991.

† Sample sizes for individual states ranged from 770 to 3083 adults aged ≥20 years who had their cholesterol

screened in 1993.

§ Sample sizes for individual states ranged from 830 to 3810 adults aged ≥20 years who had their cholesterol

sample sizes for individual states ranged from 330 to 3610 adults aged ≥20 years who had their cholesterol screened in 1995.

Sample sizes for individual states ranged from 1024 to 3449 adults aged ≥20 years who had their cholesterol screened in 1997.

** Sample sizes for individual states ranged from 958 to 5274 adults aged ≥20 years who had their cholesterol screened in 1999.

** Tonfidence interval.

55 Includes 47 states with complete data from 1991 to 1999.

From 1991 to 1999, HBC awareness increased among all demographic groups (Table 2). The percentage of persons who had ever had their cholesterol tested and who reported having been told that they had HBC was consistently higher for successive age groups (from 18.6% among those aged 20–44 years to 42.7% among those aged ≥65 years for 1999). Reported HBC awareness was higher in 1999 than in 1991 among non-Hispanic whites, non-Hispanic blacks, and Hispanics. Numbers for American Indians/Alaska Natives and Asians/Pacific Islanders were too low for meaningful analysis. Awareness of HBC was higher among women than men until 1999 and increased for both men and women.

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Editorial Note: The findings in this report indicate that among persons who had their cholesterol level screened, the percentage who were told by a health-care provider that they had HBC increased significantly from 1991 to 1999. BRFSS data on cholesterol screening trends indicated an increase in the proportion of U.S. adults aged ≥20 years who were screened during the preceding 5 years for HBC from 67.3% in 1991 to 70.8% in 1999 (3). Possible reasons for this increase include improved efforts by public health

TABLE 2. Prevalence of screened persons who were ever told they had high blood cholesterol, by selected characteristics — Behavioral Risk Factor Surveillance System, United States*, 1991–1999

					9	6 point chang	е
Characteristic	1991	1993	1995	1997	1999	1991–1999	(95% CI [†])
Age group (yrs)							
20-44	17.9	19.7	19.2	18.3	18.6	0.7	(0.2-1.4)
45-64	33.8	36.2	35.4	35.7	37.0	3.2	(2.3-4.0)
≥65	34.0	38.8	38.9	40.3	42.7	8.7	(7.7–9.7)
Race/Ethnicity [§]							
Non-Hispanic white	25.8	28.2	28.1	27.6	28.9	3.1	(2.6-3.6)
Non-Hispanic black	24.6	28.2	25.9	26.1	27.2	2.6	(1.0-4.3)
Hispanic	23.7	28.5	26.5	29.6	27.4	3.7	(1.6-5.7)
Other [¶]	28.9	30.1	26.8	27.5	30.6	1.7	(-0.8-4.3)
Sex [§]							
Women	25.7	28.3	27.6	27.8	28.1	2.4	(1.8-2.9)
Men	25.4	27.9	27.7	27.2	29.0	3.6	(2.9-4.3)

^{*}Included 47 states with no missing data (excluded District of Columbia, Kansas, Nevada, and Wyoming).

[†] Confidence interval.

[§] Age-standardized to the 2000 population.

Numbers for other race groups were too small for meaningful analysis.

programs to increase awareness of cholesterol levels, increased counseling by health-care providers, or an increase in HBC prevalence. However, data from the National Health and Nutrition Examination Survey (NHANES) suggest that cholesterol levels are declining (4).

No national data allow state-level estimates of HBC based on actual blood cholesterol measurements. NHANES used directly measured cholesterol and observed decreasing cholesterol levels among adults between the 1971–1974 and 1988–1994 surveys (4). More recent data from NHANES are not available. The differences in reported HBC across demographic variables (age, sex, and race/ethnicity) in BRFSS are consistent with those measured in NHANES III (4).

The findings in this report are subject to at least two limitations. First, BRFSS data are self-reported, and some respondents may have over or underestimated their HBC status. Patients may not have been told that they had high cholesterol and may have underestimated their HBC status. However, the actual cut-point used by health-care providers is unknown, and patients with borderline high cholesterol may have been told that their cholesterol was high, which might have resulted in an overestimate of true prevalence. Second, because BRFSS is a telephone-based survey, and persons with lower socioeconomic status are less likely than more affluent persons to have a telephone, persons with lower socioeconomic status may be underrepresented.

Control of HBC requires successful implementation of multiple steps among both patients and health-care providers, including ongoing screening for HBC, knowing one's cholesterol levels, and treating and managing HBC through lifestyle changes (e.g., reduced dietary intake of saturated fat and cholesterol, increased dietary intake of viscous fiber, increased exercise, and weight control) and medical treatment as appropriate. The National Cholesterol Education Program of the National Heart, Lung, and Blood Institute recommends that all persons aged ≥20 years have their cholesterol checked at least once every 5 years (5). In May 2001, NCEP released the third Adult Treatment Panel (ATP III) Report, which includes updated clinical guidelines for cholesterol testing and management (6,7). The new features of ATP III focus on primary prevention among those with multiple risk factors, including an assessment of the 10-year risk for a heart attack, modifications in lipid and lipoprotein classification levels, and implementation of the treatment recommendations.

HBC is a modifiable risk factor for heart disease. The benefits of cholesterol lowering include a decrease in the incidence of coronary heart disease and a decline in mortality among those with or without coronary heart disease (8–10). HBC can be prevented or controlled with increased physical activity, adoption of diets low in saturated fats and cholesterol and high in fruits and vegetables, and with the use of drugs that lower cholesterol. References*

- US Department of Health and Human Services. Healthy people 2010 (conference ed, 2 vols). Washington, DC: US Department of Health and Human Services, 2000.
- Klein RJ, Schoenborn CA. Age adjustment using the 2000 projected U.S. population: healthy people statistical notes, no. 20. Hyattsville, Maryland: US Department of Health and Human Services, CDC, National Center for Health Statistics, 2001.
- CDC. State-specific cholesterol screening trends—Behavioral Risk Factor Surveillance System, 1991–1999. MMWR 2000;49:750–5.

^{*}All MMWR references are available on the Internet at http://www.cdc.gov/mmwr. Use the search function to find specific articles.

- National Center for Health Statistics. Health, United States, 2000 with adolescent health chartbook. Hyattsville, Maryland: US Department of Health and Human Services, CDC, National Center for Health Statistics, 2000.
- Cleeman JI, Lenfant C. The National Cholesterol Education Program: progress and prospects. JAMA 1998;280:2099–104.
- National Cholesterol Education Program Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults. Executive summary of the third report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (ATP III). JAMA 2001;285:2486–97.
- National Cholesterol Education Program Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults. Third report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (ATP III). Available at http://www.nhlbi.nih.gov>. Accessed September 2001.
- Scandinavian Simvastatin Survival Study Group. Randomised trial of cholesterol lowering in 4444 patients with coronary heart disease: the Scandinavian Simvastatin survival study. Lancet 1994;344:1383–9.
- Long-Term Intervention with Pravastatin in Ischaemic Disease (LIPID) Study Group.
 Prevention of cardiovascular events and death with pravastatin in patients with coronary heart disease and a broad range of initial cholesterol levels. N Engl J Med 1998;339:1349–57.
- Downs JR, Clearfield M, Weiss S, et al. Primary prevention of acute coronary events with lovastatin in men and women with average cholesterol levels: results of AFCAPS/TexCAPS. JAMA 1998;279:1615–22.

Prevalence of Healthy Lifestyle Characteristics — Michigan, 1998 and 2000

Most persons with chronic diseases such as cardiovascular disease, cancer, diabetes, and chronic lung disease share multiple common risk factors and lifestyle behaviors (1). Tobacco use, poor diet, and physical inactivity have been identified as the leading contributors to overall mortality in the United States, accounting for one third of all deaths (2); Michigan has a particularly high burden of chronic disease-related mortality (3). To characterize the prevalence of four healthy lifestyle characteristics (HLCs) (i.e., healthy weight, adequate fruit and vegetable consumption, regular leisure-time physical activity [LTPA], and not smoking) in Michigan residents, data were analyzed from Michigan's Behavioral Risk Factor Surveillance System (BRFSS) for 1998 and 2000. This report summarizes the results of the analysis, which indicate that the proportion of Michigan residents who engaged in all four healthy lifestyle practices was extremely low, and that the prevalence was influenced by sex, education and self-reported health status. The comprehensive assessment of HLCs may be a useful adjunct to chronic disease surveillance.

BRFSS is a random-digit–dialed telephone survey of the noninstitutionalized U.S. population aged ≥18 years. Data were analyzed from 4816 adults for 1998 and 2000 combined. Missing data from 502 persons resulted in a sample size of 4314. Healthy weight was defined as having a body mass index between 18.5 and 25.0. Adequate fruit and vegetable consumption was defined as eating five or more fruits and vegetables daily. Regular LTPA was defined as at least 30 minutes of physical activity five or more times per week. Not smoking was defined according to self-reported absence of current cigarette use (i.e., former or never versus current). Data were weighted to adjust for the probability of selection and the distribution of the state's population by age, race/ethnicity, and sex. Descriptive analyses, including age-adjusted prevalence estimates, were

Healthy Lifestyle Characteristics — Continued

generated for each demographic variable (age, race/ethnicity, education, and household income) and self-reported health status using SUDAAN. Data were standardized by age to the projected 2000 U.S. population. Significant differences in the adjusted odds ratios (AORs) for engaging in all four HLCs were identified using a multiple logistic regression model that contained all independent variables.

An estimated 37.9% (95% confidence interval [CI]=36.3%–39.5%) of Michigan adults had a healthy body weight, 22.8% (95% CI=21.4%–24.2%) ate the recommended amount of fruits and vegetables, 25.9% (95% CI=24.4%–27.4%) engaged in regular LTPA, and 72.3% (95% CI=70.8%–73.8%) did not smoke. Overall, 11.2% (95% CI=10.1%–12.3%) of adults engaged in none of these practices, 38.6% (95% CI=37.0%–40.2%) in one, 33.3% (95% CI=31.7%–34.9%) in two, 13.9 % (95% CI=12.8%–15.0%) in three, and 3.0% (95% CI=2.5%–3.5%) in all four.

The prevalence of engaging in all four HLCs was significantly different by sex, education, and health status (p<0.05) (Table 1). The prevalence of engaging in all four HLCs was lower in men (age-adjusted prevalence=1.6%) than in women (age-adjusted prevalence=4.5%; AOR=0.3; 95% Cl=0.2–0.5). The prevalence of engaging in all four HLCs increased with education. The prevalence in college graduates was more than three times higher than in those with a high school education or less (AOR=3.2; 95% Cl=1.7–6.1). However, the age-adjusted prevalences were still very low in all three education groups (Table 1). The prevalence of engaging in all four HLCs decreased with decreasing health status. Persons reporting excellent health had a much higher age-adjusted prevalence (7.1% [95% Cl=5.3%–8.9%]) than adults with fair or poor health (1.0% [95% Cl=0.1%–1.9%; AOR=0.1; 95% Cl=0.04–0.4]). However, the prevalence rates in all four groups were low (Table 1).

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Editorial Note: The findings in this report document the low prevalence of healthy lifestyles in Michigan. The prevalence of HLCs in this report is consistent with that in the Nurses Health Study for a similar grouping of five healthy lifestyle behaviors (4) and is indentical to that from the overall 2000 BRFSS data. When compared with other states, obesity and smoking in Michigan are higher than the national average (5). However, the daily consumption of five fruits and vegetables in Michigan is consistent with the national average, and Michigan ranked among the top 10 states for participation in regular and sustained physical activity in 2000.

Disease risk, especially that related to cardiovascular disease, has usually been examined separately. Some studies have measured disease risk more comprehensively by combining factors such as smoking, obesity, hypertension, and high blood cholesterol (6). This study used a similar approach by assessing the combination of healthy factors that reduce disease risk, which may be a useful adjunct to the more traditional risk factor surveillance method.

The findings in this report are subject to five limitations. First, data were self-reported and some responses may be considered socially undesirable. As a result, respondents may both underreport weight (7) and overreport LTPA or fruit and vegetable consumption. Second, BRFSS collects information about LTPA only and may underestimate total activity. Third, BRFSS estimates of daily fruit and vegetable consumption are similar to

Healthy Lifestyle Characteristics — Continued

TABLE 1. Age-specific and age-adjusted* prevalence of all four healthy lifestyle characteristics (HLCs)†, and adjusted prevalence odds ratios§ among persons aged 18–74 years — Behavioral Risk Factor Surveillance System, Michigan, 1998 and 2000

Characteristic	No.¶	(%)	(95% CI**)	Odds ratio	(95% CI)
Age group (yrs)					_
18–34	1313	(3.1)	(2.0-4.2)	_	
35–54	1996	(2.7)	(2.0-3.4)	0.8	(0.5-1.3)
55–74	1005	(3.6)	(2.4-4.8)	1.4	(0.8-2.4)
Sex ^{††}					
Women	2397	(4.5)	(3.6-5.4)	_	
Men	1917	(1.6)	(1.0-2.2)	0.3	(0.2-0.5)
Race					
White	3603	(3.2)	(2.6-3.8)	_	
Black	500	(1.7)	(0.6-2.8)	0.9	(0.4-1.9)
Education⁵⁵					
High school	1701	(1.2)	(0.7-1.7)	_	
Some college	1374	(4.0)	(2.8-5.2)	3.1	(1.7–5.9)
College graduate	1232	(4.9)	(3.6-6.2)	3.2	(1.7–6.1)
Household income					
<\$35,000	1634	(2.6)	(1.7–3.5)	_	
\$35,000-\$50,000	738	(2.6)	(1.4–3.8)	0.7	(0.4–1.4)
>\$50,000	1516	(3.5)	(2.5-4.5)	0.9	(0.5-1.4)
Health status [¶]					
Excellent	952	(7.1)	(5.3-8.9)	_	
Very good	1639	(2.6)	(1.8-3.4)	0.3	(0.2-0.6)
Good	1178	(1.5)	(0.8-2.2)	0.2	(0.1–0.3)
Fair or poor	538	(1.0)	(0.1–1.9)	0.1	(0.04-0.4)

^{*} Prevalence estimates were age-adjusted using the 2000 projected U.S. population.

estimates based on multiple records but are smaller than estimates based on more extensive food-frequency questionnaires (8). Fourth, the number of black respondents in this study was too small for meaningful analysis. Finally, noncoverage and nonresponse biases related to telephone survey data may affect estimates.

Findings from previous epidemiologic studies (6,9) underscore the need for comprehensive primary prevention activities to reduce the prevalence of common chronic disease risk factors. Primary prevention may be a useful strategy in promoting the adoption and maintenance of HLCs (10). Primary prevention includes addressing the underlying social determinants that lead to behavioral and physiologic risk factors by mobilizing both health-care providers and the general population to adopt new policies. These policies include regulatory, educational, and environmental changes designed to facilitate the implementation of prevention programs.

[†]The four HLCs were defined as having a healthy body weight (body mass index between 18.5 and 25.0), getting regular leisure-time physical activity (≥30 minutes, five or more times per week), eating fruits and vegetables five or more times per day, and not smoking.

[§] Based on results of a multiple logistic regression model containing age, sex, education, household income, and health status.

[¶]Unweighted sample size for subgroups and total.

^{**}Confidence interval.

^{††} HLC significantly different by sex after adjusting for all variables (p<0.0001).

[§] HLC significantly different by education after adjusting for all variables (p<0.0001).

^{**}Response to the question, "Would you say that in general your health is excellent, very good, good, fair, or poor?" HLC significantly different by health status after adjusting for all variables (p<0.0001).

Healthy Lifestyle Characteristics — Continued

In Michigan, two initiatives sponsored by the Michigan Department of Community Health (MDCH) and the Governor's Council on Physical Fitness promote physical activity and healthy weight. First, a voluntary Exemplary Physical Education Curriculum provides school-aged children with the fitness levels, motor skills, activity-related knowledge, and personal/social skills needed for an active life. Second, environmental changes that make it easier and safer for persons to be physically active are encouraged through the "Promoting Active Communities Award," which recognizes communities that enact policies to promote physical activity. To promote a healthy diet, MDCH's 5-A-Day program provides technical support, information, and materials to local agencies to assist them in conducting local programs. MDCH also works with grocery stores to provide education materials and grocery rewards to consumers to encourage them to eat fruits and vegetables.

References

- Brownson RC, Remington PL, Davis JR. Chronic disease epidemiology and control.
 2nd ed. Washington, DC: American Public Health Association, 1998.
- McGinnis JM, Foege WH. Actual causes of death in the United States. JAMA 1993;270:2207–12.
- 3. Hahn RA, Teutsch SM, Rothenberg RB, Marks JS. Excess deaths from nine chronic diseases in the United States, 1986. JAMA 1990;264:2654–9.
- 4. Stampfer MJ, Hu FB, Manson JE, Rimm EB, Willet WC. Primary prevention of coronary heart disease in women through diet and lifestyle. N Engl J Med 2000;343:16–22.
- CDC. Chronic diseases and their risk factors: the nation's leading causes of death: a report with expanded state-by-state information. Atlanta, Georgia: US Department of Health and Human Services, CDC, 1999.
- Yusuf HB, Giles WH, Croft JB, Anda RF, Casper ML. Impact of multiple risk factors profiles on determining cardiovascular disease risk. Prev Med 1998;27:1–9.
- 7. Rowland ML. Self-reported weight and height. Am J Clin Nutr 1990;52:1125-33.
- Serdula MK, Coates RC, Byers T, et al. Evaluation of a brief telephone questionnaire to estimate fruit and vegetable consumption in diverse study populations. Epidemiology 1993;4:455–63.
- 9. Stamler J, Dyer AR, Shekelle RB, Neaton J, Stamler R. Relationship of baseline major risk factors to coronary and all-cause mortality, and to longevity: findings from long term follow-up of Chicago cohorts. Cardiology 1993;82:191–222.
- 10. Rose G. The strategy of preventive medicine. Oxford: Oxford University Press, 1992.

Outbreak of Powassan Encephalitis — Maine and Vermont, 1999–2001

Powassan (POW) virus, a North American tickborne flavivirus related to the Eastern Hemisphere's tickborne encephalitis viruses (1), was first isolated from a patient with encephalitis in 1958 (1,2). During 1958–1998, 27 human POW encephalitis cases were reported from Canada and the northeastern United States (3). During September 1999–July 2001, four Maine and Vermont residents with encephalitis were found to be infected with POW virus. These persons were tested for other arbovirus infections found in the northeast after testing for West Nile virus (WNV) infection was negative. This report describes these four cases, summarizes the results of ecologic investigations, and discusses a potential association between ticks that infest medium-sized mammals and the risk for human exposure to POW virus. The findings underscore the need for personal protective measures to prevent tick bites and continued encephalitis surveillance.

Powassan Encephalitis — Continued

Case Reports

Case 1. In June 2001, a 70-year-old man from Kennebec County, Maine, was taken to a local hospital with generalized muscle weakness, somnolence, diarrhea, and anorexia. On clinical examination, he had a fever of 104.7 F (40.4 C), leukocytosis of 11,500/mm³ (normal: 4,300–10,800/mm³), decreased renal function, and anemia. He subsequently developed left-sided hemiplegia and marked confusion. Cerebrospinal fluid (CSF) contained 40 white blood cells (WBCs)/mm³ (normal: <4/mm³) (87% lymphocytes) with elevated protein (96 mg/dL; normal: 20–50 mg/dL). Magnetic resonance imaging (MRI) revealed parietal changes consistent with microvascular ischemia or demyelinating disease. No causes for his apparent stroke were found. After 22 days of hospitalization, he was discharged to a rehabilitation facility. Nearly 3 months after symptom onset, he remains in the facility and is unable to move his left arm or leg. Serum specimens and CSF collected 3 days after hospitalization revealed POW virus-specific lgM; neutralizing antibody (1:640 titer) also was found in serum specimens. Although some cross-reaction with WNV and St. Louis encephalitis (SLE) virus occurred in the lgM assay, no neutralizing antibody was found.

The patient had not left Maine for 25 years. On ecologic investigation, overgrown bushes, leaf piles, and stacks of old lumber and scrap metal covered his property. Family members reported seeing woodchucks, skunks, and squirrels on the property. During the 2 weeks before illness, the patient's main activities were lying on the ground repairing a boat hull and yard work. Approximately 6 weeks after illness onset, nine medium-sized mammals were trapped on or near the patient's property. Collections from these mammals and the grassy and brushy areas of the property yielded 31 ticks (*Ixodes cookei*). Tests for POW virus infection were conducted at CDC. Of the nine mammal serum samples, four (two woodchucks and two skunks) contained neutralizing antibody to POW virus, but no virus was isolated from the ticks.

Case 2. In September 2000, a 53-year-old woman from York County, Maine, sought medical care at a local hospital for loss of balance, visual disturbance, and fever of 103 F (39.4 C). Her clinical examination showed agitation without confusion, ataxia, bilateral lateral gaze palsy, and dysarthria. CSF contained 148 WBCs/mm³ (46% neutrophils, 40% lymphocytes). During hospitalization, she developed altered mental status, generalized muscle weakness, and complete ophthalmoplegia. An electroencephalogram (EEG) indicated diffuse encephalitis, and a MRI showed bilateral temporal lobe abnormalities consistent with microvascular ischemia or demyelinating disease. After 13 days, she was transferred to a rehabilitation facility where she remained for 2 months. Nine months after onset of symptoms, she was walking and had regained her strength, but the ophthalmoplegia continued. A serum specimen collected 19 days after illness onset was positive for POW virus-specific IgM and neutralizing antibody (1:640 titer) and negative for WNV and SLE virus antibodies.

The patient had not left Maine in several months before illness onset. During two visits to a rural vacation home in the month before illness onset, the patient removed several squirrel nests but reported no contact with ticks or rodents. One month after illness onset, an ecologic evaluation of her primary home noted a well-manicured suburban property near brush and woodlands. No evidence of medium-sized mammals was found, and only three *lx. scapularis* were collected; no POW virus was isolated. Nine months after illness onset, an ecologic evaluation of the patient's vacation home found several mammals, but none had ticks, and no serology samples were collected.

Powassan Encephalitis — Continued

Case 3. In July 2000, a 25-year-old man from Waldo County, Maine, sought medical care at a local hospital for fever of 101.3 F (38.5 C), headache, vomiting, somnolence, and confusion. On clinical examination, the patient had difficulty answering simple questions and was intermittently uncooperative. He had bilateral hand twitching, muscle weakness, and pronounced lip smacking. CSF contained 920 WBCs/mm³ (74% lymphocytes) with elevated protein (77 mg/dL). EEG showed diffuse background slowing consistent with encephalitis. After 11 days of hospitalization, he was transferred to a rehabilitation facility. When discharged home 44 days later, the patient required assistance to stand and perform daily activities. Serum specimens and CSF collected 3 days after illness onset were negative for antibody to WNV and SLE virus but positive for POW virus-specific IgM antibody. The serum sample also had neutralizing antibody (1:80 titer) to POW virus. At the time of illness onset, the patient worked as a logger and lived in rural Maine where he raised livestock.

Case 4. In September 1999, a 66-year-old man from Washington County, Vermont, sought medical care at a hospital for somnolence, severe headache, increasing confusion, and bilateral leg weakness that developed over 6 days. On clinical examination, he was afebrile but had slow speech, memory loss, a wide-based gait, and bilateral weakness in proximal lower extremities. CSF contained 54 WBCs/mm³ (95% lymphocytes) and elevated protein (67 mg/dL). An EEG showed diffuse background slowing consistent with encephalitis. When discharged home 11 days later, he could walk but had cognitive difficulties, including severe memory lapses. Serum specimens collected 19 days after illness onset contained POW virus-specific lgM and neutralizing antibody (1:640 titer) but no antibody to WNV and SLE virus. During the month before illness onset, the patient traveled frequently to a vacation home where he saw numerous squirrels and skunks.

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Editorial Note: These four cases of POW encephalitis are the first reported in Maine and Vermont and the first in the United States since 1994 (4). Since the introduction of WNV into the northeastern United States in 1999 (5), testing for POW virus and other arboviruses that cause encephalitis has increased (CDC, unpublished data, 2001). These cases were identified as a direct result of requests for WNV testing. As surveillance continues, knowledge of the epidemiology of POW virus in the United States may increase.

In North America, POW virus has been isolated from four tick species, including *lx. cookei, lx. marxi, lx. spinipalpus*, and *Dermacentor andersoni*; a variant POW virus also has been isolated from *lx. scapularis*; and evidence of infection has been found in 38 mammal species, primarily woodchucks (1,6). Unlike *lx. scapularis*, the primary vector for Lyme disease, *lx. cookei* rarely search for hosts on vegetation and are often found in or near the nests or burrows of medium-sized mammals. Infections have occurred from May to December, with a peak during June–September when ticks are most active (1). Although neither the first or second patients recalled tick bites, ecologic investigations suggest that their illnesses resulted from visiting or living in areas where ticks are common. As with many infectious agents transmitted by *lxodid* ticks, few infected persons recalled tick bites because these ticks are small and can be easily missed (3).

Powassan Encephalitis — Continued

POW encephalitis is associated with significant long-term morbidity and has a case-fatality rate of 10%–15% (1,3). Because there is no vaccine or specific therapy for POW encephalitis, the best means of prevention is protection from tick bite. This includes using insect repellents, wearing light-colored clothing with long sleeves and pants tucked into socks or boots, avoiding or clearing brushy areas, and removing ticks before they attach or as soon after attachment as possible. Checking family pets also can prevent ticks from entering the home. Because *lx. cookei* are often found on woodchucks and skunks and may be the primary vector of POW virus, environmental controls reducing human contact with small and medium-sized mammals should reduce risk for exposure to POW virus-infected ticks. Persons should keep areas adjacent to their home clear of brush, weeds, trash, and other elements that could support small and medium-sized mammals. When removing rodent nests, avoid direct contact with nesting materials and use sealed plastic bags for disposal and to prevent direct contact with ticks.

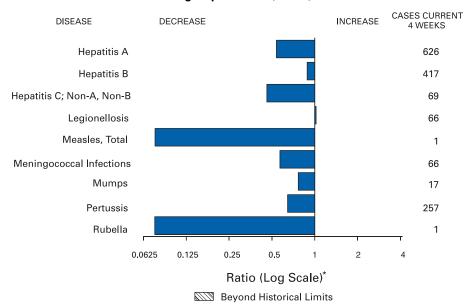
Because of the lack of awareness and the need for specialized laboratory tests to confirm diagnosis, the frequency of POW encephalitis may be greater than previously suspected. POW encephalitis should be included in the differential diagnosis of all encephalitis cases occurring in the northern United States, especially the northeast. Laboratory tests for POW virus infection are not commercially available but can be requested through state public health laboratories for testing at CDC. Awareness should be promoted among clinicians and public health staff, and tick-bite prevention strategies emphasized for the general public.

References*

- 1. Artsob H. Powassan encephalitis. In: Monath T, ed. The arboviruses: epidemiology and ecology. Volume IV. Boca Raton, Florida: CRC Press, 1988:29–49.
- McLean DM, Donohue WL. Powassan virus: isolation of virus from a fatal case of encephalitis. Can Med Assoc J 1958:80:708–11.
- Gholam BIA, Puksa S, Provias JP. Powassan encephalitis: a case report with neuropathology and literature review. Can Med Assoc J 1999;161:1419–22.
- 4. CDC. Arboviral disease—United States, 1994. MMWR 1995;44:641-4.
- 5. Nash D, Mostashari F, Fine A, Miller J, O'Leary D, Murray K. The outbreak of West Nile virus infection in the New York City area in 1999. N Engl J Med 2001:344:1807–14.
- Ebel GD, Spielman A, Telford SR. Phylogeny of North American Powassan virus. J Gen Vir 2001;82:1657–65.

^{*}Reference to non-CDC sites on the Internet are provided as a service to *MMWR* readers and do not constitute or imply endorsement of these organizations or their programs by CDC or the U.S. Department of Health and Human Services. CDC is not responsible for the content of pages found at these sites.

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



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

TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending September 1, 2001 (35th Week)*

		Cum. 2001		Cum. 2001
Anthrax		-	Poliomyelitis, paralytic	_
Brucellosis†		53	Psittacosis†	9
Cholera		3	Q fever [†]	15
Cyclosporiasis	s [†]	108	Rabies, human	1
Diphtheria		1	Rocky Mountain spotted fever (RMSF)	319
Ehrlichiosis:	human granulocytic (HGE)†	127	Rubella, congenital syndrome	-
	human monocytic (HME)†	54	Streptococcal disease, invasive, group A	2.572
Encephalitis:		26	Streptococcal toxic-shock syndrome [†]	44
•	eastern equine [†]	4	Syphilis, congenital 1	161
	St. Louis [†]	1	Tetanus	17
	western equine [†]	-	Toxic-shock syndrome	82
Hansen diseas	se (leprosy)†	51	Trichinosis	14
	ılmonary syndrome [†]	5	Tularemia [†]	71
Hemolytic ure	emic syndrome, postdiarrheal [†]	76	Typhoid fever	170
HIV infection,	pediatric†§	98	Yellow fever	-
Plague	•	2		

^{-:} No reported cases.

^{*}Incidence data for reporting year 2001 are provisonal and cumulative (year-to-date).

[†] Not notifiable in all states.

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 26, 2001.
Updated from reports to the Division of STD Prevention, NCHSTP.

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending September 1, 2001, and September 2, 2000 (35th Week)*

week	s endin	g Septe	ptembe	r 2, 200	U (35th	vveek)*				
			01.1						coli 0157:H	
	Cum.	Cum.	Chlan Cum.	Cum.	Cryptos Cum.	poridiosis Cum.	NET Cum.	Cum.	Cum.	LIS Cum.
Reporting Area	2001 ¹	2000	2001	2000	2001	2000	2001	2000	2001	2000
UNITED STATES	19,145	26,250	449,858	463,520	1,422	1,534	1,565	2,925	1,343	2,552
NEW ENGLAND Maine	746 20	1,420 25	14,977 668	15,576 965	69 11	84 13	164 21	260 18	157 22	286 25
N.H. Vt.	17 10	25 27	809 399	678 358	4 25	11 18	24 11	23 27	21 5	31 29
Mass.	411	890	6,857	6,616	22 3	27	84 9	123	77	126
R.I. Conn.	53 235	62 391	1,942 4,302	1,700 5,259	4	2 13	15	11 58	7 25	12 63
MID. ATLANTIC Upstate N.Y.	3,974 322	5,811 607	49,973 8,902	43,325 976	164 66	230 61	113 87	304 187	122 85	210 39
N.Y. City	1,996	3,136	19,132	17,836	65	120	8	19	8	14
N.J. Pa.	960 696	1,153 915	8,038 13,901	7,872 16,641	4 29	11 38	18 N	98 N	29 -	95 62
E.N. CENTRAL Ohio	1,408 237	2,458 388	67,466 13,962	79,649 20,725	418 103	433 68	374 97	706 144	282 84	541 157
Ind.	165	250	9,177	8,820	50	28	50	82	32	6 8
III. Mich.	665 261	1,365 331	17,349 19,591	22,500 16,741	1 106	59 59	93 56	145 86	80 50	117 73
Wis. W.N. CENTRAL	80 454	124 614	7,387 22.864	10,863 26.080	158 207	219 158	78 251	249 421	36 233	126 421
Minn.	85 47	115	4,261	5,308	99 53	21 46	92	101	98 39	127 109
lowa Mo.	218	66 286	1,858 9,007	3,566 8,892	26	22	50 33	118 85	49	77
N. Dak. S. Dak.	1 18	2 6	599 1,201	597 1,201	7 6	9 9	12 18	14 35	21 19	16 40
Nebr. Kans.	39 46	43 96	2,054 3,884	2,478 4,038	15 1	43 8	32 14	50 18	7	40 12
S. ATLANTIC	6,167	7,196	86,104	86,827	206	248	141	224 1	99 4	217
Del. Md.	116 751	131 842	1,912 7,634	1,916 9,226	2 28	5 9	3 14	19	1	1 1
D.C. Va.	465 501	499 461	1,838 11,965	2,122 10,768	10 15	6 11	38	47	U 30	U 45
W. Va. N.C.	49 402	42 431	1,544 13,769	1,434 14.905	1 19	3 18	5 30	11 50	6 26	7 54
S.C. Ga.	350 757	530 872	7,750 16,870	5,916 18,430	74	95	7 19	16 34	9 13	13 36
Fla.	2,776	3,388	22,822	22,110	57	101	25	46	10	60
E.S. CENTRAL Ky.	977 201	1,295 146	31,630 5,999	33,983 5,335	31 3	37 5	85 41	87 26	79 39	79 25
Ténn. Ala.	293 224	531 337	9,680 8,269	9,554 10,806	8 11	9 12	25 12	38 5	30 6	41 5
Miss.	259	281	7,682	8,288	9	11	7	18	4	8
W.S. CENTRAL Ark.	2,058 104	2,672 126	68,974 4,742	69,793 4,457	22 5	85 7	44 6	193 49	59 -	233 34
La. Okla.	472 107	445 219	11,314 7,147	12,345 5,821	7	10 7	3 18	13 13	24 20	37 11
Tex.	1,375	1,882	45,//1	47,170	2	61	17	118	15	151
MOUNTAIN Mont.	714 12	1,007 10	25,981 1,305	26,868 1,016	105 7	77 8	176 10	286 26	100	213
Idaho Wyo.	15 1	16 7	1,209 564	1,241 526	12 2	4 5	29 7	44 12	- 1	26 9
Colo. N. Mex.	140 56	239 107	5,284 3,622	7,988 3,278	29 18	33 7	69 10	108 15	54 8	77 14
Ariz.	295	319	9,684	8,642	6	7	20	36	12	27
Utah Nev.	63 132	97 212	1,279 3,034	1,569 2,608	27 4	10 3	22 9	37 8	24 1	50 10
PACIFIC Wash.	2,647 290	3,777 334	81,889 9,111	81,419 8,714	200 37	182 U	217 59	444 131	212 62	352 159
Oreg. Calif.	112 2,204	113 3,229	3,108 65,435	4,630 64,073	22 137	13 169	32 113	94 183	27 119	90 91
Alaska Hawaii	2,204 13 28	3,229 15 86	1,796 2,439	1,633 2,369	137	-	3 10	24 12	- 4	2 10
Guam	9	13	_	335	-	-	N	N	U	U
P.R. V.I.	580 2	759 25	1,764 53	Ü	-	-	1 -	5	Ü	Ü
Amer. Samoa C.N.M.I.	-	-	Ŭ 85	U	U	U U	U	U	Ŭ	Ŭ

N: Not notifiable.

U: Unavailable. -: No reported cases.

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

^{*}Incidence data for reporting year 2001 are provisonal and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

and cumulative (year-to-date).

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

Chlamydia refers to genital infections caused by *C. trachomatis*.

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TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending September 1, 2001, and September 2, 2000 (35th Week)*

week	s enaing s	September			Tember	2, 20	UU (35th		
	Gono	rrhea	Hepati Non-A,	tis C; Non-B	Legione	llosis	Listeriosis		yme sease
Reporting Area	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2001	Cum. 2000
UNITED STATES	207,338	233,931	2,323	2,205	606	658	292	6,829	10,932
NEW ENGLAND Maine	4,202 79	4,424 55	14	21 2	29 4	38 2	32	1,989	3,316
N.H.	107	69	-	-	7	2	2	88 4	41
Vt. Mass.	48 2,089	43 1,791	6 8	4 10	4 5	3 15	2 16	409	23 982
R.I. Conn.	501 1,378	412 2,054	-	5 -	2 7	3 13	1 11	233 1,255	213 2,057
MID. ATLANTIC Upstate N.Y.	25,210 5.458	25,017 4,677	978 40	485 25	122 39	178 47	45 19	3,492 1,912	5,761 2,174
N.Y. City N.J.	8,016 4,958	7,553 4,964	896	425	10 5	26 16	8 7	2 448	153 2,142
Pa.	6,778	7,823	42	35	68	89	11	1,130	1,292
E.N. CENTRAL Ohio	36,350 7,674	47,155 12,449	123 8	173 8	149 82	177 69	34 11	379 83	666 46
Ind.	3,842 10,832	4,128 14,062	1 11	- 17	14	26 24	4	16	19 33
Mich. Wis.	11,441 2,561	11,841 4,675	103	148	33 20	30 28	16 2	1 279	21 547
W.N. CENTRAL	9,762	11,575	472	398	40	44	9	248	178
Minn. Iowa	1,375 428	2,131 804	7 -	5 1	9 6	3 11	-	202 24	99 21
Mo. N. Dak.	5,364 19	5,636 44	455 -	382	15 1	21	5 -	17 -	41 -
S. Dak. Nebr.	186 695	194 959	3	3	3 5	2 3	1	3	3
Kans. S. ATLANTIC	1,695 53,225	1,807 61,002	7 81	7	1 132	4 108	3 50	2 585	14 834
Del.	1,122	1,120	81 - 14	67 2 8	3	5 40	-	31 379	163
Md. D.C.	4,231 1,644	6,271 1,649		8 3 3	27 7	-	9	8	495 3
Va. W. Va.	7,019 423	6,666 440	9	13	18 <u>N</u>	19 N	9 5	98 10	103 22
N.C. S.C.	11,257 5,344	12,173 5,572	16 5	13 1	7 6	9 4	2	27 3	35 3
Ga. Fla.	9,248 12,937	11,741 15,370	37	3 21	9 55	6 25	7 14	29	10
E.S. CENTRAL Ky.	20,278 2,371	24,433 2,334	159 6	328 29	42 9	25 14	15 4	33 18	33 6
Tenn. Ala.	6,436 6.415	7,689 8,298	51 2	67 7	21 10	8	6 5	9	19 5
Miss.	5,056	6,112	100	225	2	1	-	-	3
W.S. CENTRAL Ark.	33,805 2,932	36,583 2,538	162 3	547 7	5 -	20	6 1	7	58 5
La. Okla.	7,848 3,302	8,981 2,494	75 3	303 6	2	7 2	2	1 -	5 -
Tex.	19,723	22,570	81	231	-	11	3	6	48
MOUNTAIN Mont.	6,695 78	7,020 28	238 1	54 4	40	25 1	26	10	7
ldaho Wyo.	53 46	59 36	2 191	3 2	2	4	1 1	4	1 3
Colo. N. Mex.	2,054 592	2,110 726	16 11	11 11	11 2	8 1	6 6	1 -	-
Ariz. Utah	2,677 104	2,916 162	9 2	13 -	11 7	6 5	6 1	1	1
Nev.	1,091	983	6	10	3	-	5	1	2
PACIFIC Wash.	17,811 2,024	16,722 1,509	96 16	132 23 22	47 6	43 14	75 6	86 6	79 5
Oreg. Calif.	456 14,669	622 14,060	10 70	22 85	N 37	N 29	3 62	6 72	6 66
Alaska Hawaii	266 396	217 314	-	2	4	-	4	2 N	2 N
Guam P.R.	399	34 352	- 1	2 1	2	- 1	-	- N	N
V.I. Amer. Samoa	6 U	- U	Ü	Ü	Ū	Ü	-	Ū	Ü
C.N.M.I.	7	ŭ		ŭ	-	ŭ	-	-	ŭ

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

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending September 1, 2001, and September 2, 2000 (35th Week)*

week	s enaing	Septemi	Jer 1, 20	vi, and S	eptember	2, 2000 (3		K) "
	Mal	aria	Rabie	es, Animal	NE	Salmor TSS		HLIS
Panarting Area	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
Reporting Area UNITED STATES	2001 748	2000 936	2001 4,145	2000 4,739	2001 21,878	2000 24,679	2001 18,195	2000 21,635
NEW ENGLAND	39	49	460	537	1,536	1,547	1,518	1,594
Maine N.H.	4 2	5 1	47 16	90 9	140 129	92 94	121 120	77 97
Vt. Mass.	- 12	2 20	47 175	43 183	50 927	86 913	45 801	88 907
R.I.	3	5	41	36	82	83	114	115
Conn. MID. ATLANTIC	18 198	16 233	134 827	176	208	279	317 2,554	310 3,501
Upstate N.Y.	45	44	532	847 541	2,816 795	3,315 773	816	893
N.Y. City N.J.	104 21	124 39	20 130	8 113	719 589	844 806	790 527	873 673
Pa.	28	26	145	185	713	892	421	1,062
E.N. CENTRAL Ohio	70 20	103 13	88 33	113 33	3,146 942	3,375 799	2,690 795	2,374 1,014
Ind. III.	14 1	5 53	1 14	19	354 767	410 1,074	310 704	435 1
Mich. Wis.	22 13	21 11	34 6	50 11	551 532	609 483	566 315	654 270
W.N. CENTRAL Minn.	27 6	39 13	243 29	409 65	1,417 381	1,595 369	1,518 474	1,765 481
Iowa	5	2	55	60	216	230	209	240
Mo. N. Dak.	9	9 2	32 29	35 94	404 43	484 47	549 59	585 56
S. Dak. Nebr.	2	7	25 4	76 1	110 100	62 146	92	<i>7</i> 8 111
Kans.	5	6	69	78	163	257	135	214
S. ATLANTIC Del.	207 1	205 3	1,454 25	1,648 31	5,455 58	4,789 80	3,818 61	3,948 92
Md. D.C.	89 13	74 13	179 -	292	555 57	527 39	603 U	480 U
Va. W. Va.	40 1	41 2	292 102	398 89	931 80	653 107	678 92	640 106
N.C. S.C.	9 5	19 2	404 86	403 107	759 556	657 480	723 459	734 373
Ga.	12	8	224	218	855	796	884	1,188
Fla. E.S. CENTRAL	37 22	43 31	142 149	110 135	1,604 1 <i>.</i> 474	1,450 1,458	318 1,057	335 1,180
Ky. Tenn.	8 8	9	15 87	18 72	236 389	259 389	143 452	186 527
Ala.	4 2	13	47	44 1	426 423	393	328	387 80
Miss. W.S. CENTRAL	10	1 58	510	634	1,562	417 3,100	134 1,297	1,878
Ark. La.	3 4	2 10	20	20 3	468 270	433 514	92 458	362 416
Okla. Tex.	2 1	4 42	48 442	44 567	278 546	262 1,891	236 511	196 904
MOUNTAIN	35	35	180	196	1,444	1,841	1,080	1,769
Mont. Idaho	2 3	1 2	31 13	52 9	, 49 96	69 90	4	81
Wyo. Colo.	- 18	18	21	42	44 406	48 502	43 360	40 491
N. Mex.	3	-	11	16	181	164	146	152
Ariz. Utah	3	6 4	96 7	66 9	415 155	442 341	368 136	483 346
Nev. PACIFIC	3 140	4 183	1 234	2 220	98 3,028	185 3,659	23 2,663	176 3,626
Wash.	4	19	-	-	337	346	491	471
Oreg. Calif.	9 119	30 125	1 196	6 189	165 2,250	213 2,906	230 1,701	270 2,700
Alaska Hawaii	1 7	9	37 -	25 -	28 248	39 155	2 239	25 160
Guam	-	2	-	-	-	20	U	U
P.R. V.I.	3	4	67 	56 	405	427 	U U	U
Amer. Samoa C.N.M.I.	U -	U U	U -	U U	U 8	U U	U U	U U
Ni. Nież weżifielele	Hillian	- امامانه،	. N					

N: Not notifiable.

U: Unavailable.

-: No reported cases.

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

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

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending September 1, 2001, and September 2, 2000 (35th Week)*

weeks ending September 1, 2001, and September 2, 2000 (35th Week)* Shigellosis' Synhilis												
	NET			PHLIS	Sy (Primary)	philis & Secondary)	Tube	erculosis				
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.				
Reporting Area UNITED STATES	2001 10,648	2000 14,589	2001 5,227	2000 8,183	2001 3,721	2000 4,099	2001 7,908	9,408				
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	185 6 4 7 131 15 22	274 8 4 3 199 19	172 2 2 2 2 116 19 31	263 11 7 179 22 44	37 - 1 2 19 7 8	4,099 55 1 1 - 38 4 11	290 7 11 2 164 24 82	286 12 15 4 168 25 62				
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	933 378 258 146 151	1,890 526 776 398 190	582 93 267 157 65	1,209 177 514 332 186	314 19 161 77 57	190 7 81 46 56	1,555 221 811 337 186	1,539 209 823 359 148				
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	2,770 1,948 153 271 202 196	3,011 232 1,133 855 544 247	1,331 923 28 204 156 20	871 206 129 2 491 43	616 58 111 160 269 18	866 55 257 298 217 39	832 142 66 406 171 47	920 198 87 430 146 59				
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	1,051 286 316 199 20 122 54 54	1,621 518 361 505 12 4 76 145	851 341 261 140 21 59	1,375 594 260 356 21 3 61 80	50 21 1 11 - - 2 15	48 8 10 25 - - 2 3	303 158 18 91 3 8 25	339 108 25 129 2 13 14 48				
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	1,583 7 106 42 202 8 245 202 154 617	1,871 12 134 41 311 4 124 95 167 983	517 7 57 U 110 8 125 91 91 28	701 15 73 U 238 3 95 68 134 75	1,331 8 162 28 76 - 307 178 222 350	1,351 7 201 29 95 3 353 143 260 260	1,618 9 141 51 162 21 236 134 290 574	1,937 10 169 16 188 21 257 186 425 665				
E.S. CENTRAL Ky. Tenn. Ala. Miss.	932 340 66 175 351	656 236 249 37 134	400 175 75 124 26	361 52 278 28 3	406 30 215 87 74	595 59 358 83 95	499 78 192 164 65	596 70 224 197 105				
W.S. CENTRAL Ark. La. Okla. Tex.	1,062 415 112 32 503	2,337 147 199 77 1,914	714 155 132 15 412	712 43 125 30 514	478 26 100 48 304	560 75 150 82 253	712 100 - 100 512	1,404 143 122 109 1,030				
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	640 2 25 3 157 79 284 44 46	713 6 41 5 129 87 290 57 98	372 - 1 140 45 137 41 8	506 - 25 3 91 63 194 64 66	167 - - 1 31 13 111 7 4	156 - 1 1 7 12 130 1	301 6 8 2 78 21 115 24 47	346 10 6 2 56 29 139 32 72				
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	1,492 139 58 1,241 5 49	2,216 345 121 1,717 7 26	288 167 74 - 1 46	2,185 321 80 1,757 3 24	322 37 8 269	278 47 10 220 - 1	1,798 167 74 1,431 31 95	2,041 162 64 1,649 74 92				
Guam P.R. V.I. Amer. Samoa	- 8 - U	34 25 - U	U U U	U U U	172 - U	3 116 - U	76 - U	37 109 - U				
C.N.M.I.	4	U	U	U	-	U	20	U				

N: Not notifiable.

U: Unavailable. -: No reported cases.

^{*}Incidence data for reporting year 2001 are provisonal and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

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

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending September 1, 2001, and September 2, 2000 (35th Week)*

N: Not notifiable. U: Unavailable.

[:] No reported cases. *Incidence data for reporting year 2001 are provisonal and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

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

[§] Of 194 cases among children aged <5 years, serotype was reported for 95, and of those, 17 were type b.

TABLE III. (Cont'd) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending September 1, 2001, and September 2, 2000 (35th Week)*

		, 2000	(35th	Week)	*						
	Dis	gococcal ease		Mumps			Pertussis			Rubella	
Reporting Area	Cum. 2001	Cum. 2000	2001	Cum. 2001	Cum. 2000	2001	Cum. 2001	Cum. 2000	2001	Cum. 2001	Cum. 2000
UNITED STATES	1,560	1,558	2	150	247	78	2,988	4,194	-	17	107
NEW ENGLAND	83	92	-	-	4	-	263	1,085	-	-	11
Maine N.H.	1 10	7 9	Ū	-	-	Ū	25	31 79	Ū	-	2
Vt. Mass.	5 47	2 54	-	-	- 1	-	25 194	173 751	-	-	- 8
R.I.	2	7	-	-	1	-	5	14	-	-	-
Conn. MID. ATLANTIC	18 166	13 176	-	- 17	2 19	- 1	14 216	37 389	-	5	1 8
Upstate N.Y.	46	47	-	3	6	1	117	179		1	1
N.Y. City N.J.	31 39	35 33	-	9 2	6 3	-	34 13	54 30	-	3 1	7
Pa.	50	61	-	3	4	-	52	126	-	-	-
E.N. CENTRAL Ohio	200 70	268 63	-	14 1	18 7	5 1	367 217	486 228	-	3	1
Ind.	29	31	-	1	-	2	46	62	-	1	-
III. Mich.	20 46	68 76	-	10 2	6 4	2	41 39	54 55		2	1 -
Wis.	35	30	-	-	1	-	24	87	-	-	-
W.N. CENTRAL Minn.	104 15	109 17	-	8 3	14	14 11	168 58	306 180	-	3	1 -
lowa Mo.	21 39	22 51	-	-	6 4	-	17 69	34 48	-	1 1	-
N. Dak.	5	2	-	-		-	3	2	-		-
S. Dak. Nebr.	4 10	5 5	Ū	1	1	Ū	4	3 9	Ū	-	1
Kans.	10	7	-	4	3	3	17	30	-	1	-
S. ATLANTIC Del.	301 3	223	2	26	37 -	7 -	163	310 8	-	4	60
Md. D.C.	35	22	-	4	8	1	21 1	78 3	-	-	-
Va.	31	35	-	6	8	3	31	44 1	-	-	-
W. Va. N.C.	11 58	10 32	2	3	5	3	2 51	74	-	-	52
S.C. Ga.	31 36	18 37	-	2 7	10 2	-	26 7	23 27	-	2	6
Fla.	96	69	-	4	4	-	24	52	-	2	2
E.S. CENTRAL Ky.	103 18	109 23	-	3 1	4	5 1	85 18	88 44	-	-	5 1
Tenn. Ala.	44 30	45 30	-	-	2	2 2	37 27	25 16	-	-	i 3
Miss.	11	11	-	2	-	-	3	3	-	-	-
W.S. CENTRAL	176	166 11	-	8 1	25 1	2	248	223	-	-	7 1
Ark. La.	16 56	38	-	2	1 5	2	11 2	31 15	-	-	1
Okla. Tex.	24 80	22 95	Ū	5	19	Ū	1 234	12 165	Ū	-	- 5
MOUNTAIN	76	71	-	9	16	38	1,039	494	_	1	2
Mont. Idaho	3 7	4 6	U	1 1	1	U	21 165	24 45	U	-	-
Wyo.	6 27	23	-	i 1	1	-	2 205	3 271	-	- 1	- 1
Colo. N. Mex.	11	6	-	2	1	12 2	89	76	-	-	-
Ariz. Utah	11 7	22 7	-	1 1	4 4	23 1	491 57	51 15	-	-	1 -
Nev.	4	3	U	1	5	U	9	9	U	-	-
PACIFIC Wash.	351 53	344 36	-	65 1	110 5	6 5	439 104	813 239	-	1	12 7
Oreg.	30	45	N	N	Ň 77	1	35	88	Ū	-	-
Calif. Alaska	257 2	249 6	U -	29 1	8	U -	268 3	437 18	-	-	5
Hawaii	9	8	-	34	20	-	29	31	-	1	-
Guam P.R.	4	8	U	-	12	U	2	3 5	U	-	1 -
V.I. Amer. Samoa	Ū	Ū	U U	Ū	Ū	U U	Ū	Ū	U	Ū	Ū
C.N.M.I.		Ŭ	Ŭ		Ŭ	Ŭ		Ŭ	Ŭ		ŭ

N: Not notifiable. U: Unavailable.

^{-:} No reported cases. *Incidence data for reporting year 2001 are provisonal and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulat (year-to-date).

TABLE IV. Deaths in 122 U.S. cities,* week ending September 1, 2001 (35th Week)

		All Cau	ıses, By	Age (Y			P&I	OUT (SSEIT W		All Cau	ses, By	Age (Y	ears)		
Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	P&I [†] Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn Cambridge, Mass Fall River, Mass. Hartford, Conn. Lowell, Mass. Lynn, Mass. New Bedford, Mass. New Bedford, Mass. New Haven, Conn. Providence, R.I. Somerville, Mass Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Erie, Pa.Š. Jersey City, N.J.	. 12 26 45 5 5 5 5 5 5 5 5 5 5 3 3 4 43 3 1,608 49 19 605 11 11 3 24	363 88 U 9 9 23 35 20 5 18 30 39 22 23 29 42 1,131 33 35 6 8 8 8 4 16	3 2 9 4	33 12 U - 1 1 - 1 3 6 - 4 2 2 3 3 122 1 1 3 4 2 1 1 1	6 2 U 1 1 1 28 2 1 3 2 1 - 1	9 5 U - - 1 1 1 23 5 - -	51 18 10 11 13 13 13 13 14 14 15 16 16 18 17 18 18 19 19 19 19 19 19 19 19 19 19 19 19 19	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, Fla. Washington, D.d Wilmington, D.d Wilmington, D.d Birmingham, Alc Chattanooga, Te Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, A Nashville, Tenn.	97 55 74 50 Fla. 68 180 C. 104 I. U 706 a. 148 enn. 64 90 27 . 169 94 Ia. U	731 96 104 99 96 59 22 40 30 49 118 68 U 488 100 44 62 22 124 63 U 7	260 47 344 18 20 16 16 19 11 13 45 21 U 145 31 12 24 4 30 19 U 25 312	114 20 18 5 13 17 5 5 11 12 U 42 10 4 3 1 7 9 U 8	39 5 4 4 4 3 2 4 9 1 1 1 5 1 U 18 4 3 1 - 4 2 U 4 71	43 9 5 5 5 4 3 8 8 3 3 · 1 2 U 13 3 1 · · · 4 1 U 4 2	57 54 12 7 - 2 - 5 4 2 U 60 18 5 5 2 17 4 U 9
New York City, N.' Newark, N.J. Paterson, N.J. Philadelphia, Pa. Pittsburgh, Pa. S Reading, Pa. Rochester, N.Y. Schenectady, N.Y. Scranton, Pa. S Syracuse, N.Y. Trenton, N.J. Utica, N.Y. Yonkers, N.Y.	U 15 U 29 15 99	742 U 9 U 19 12 79 17 31 53 12 U	U 4 U 6 3 14 - 8 11 3	90 U 1 U 3 - 6 1 2 4 2	16 U 1 U 1 - - - - U	16 U - U - - - - - U	43 U · U 1 2 8 1 11 1 U	W.S. CENTRAL Austin, Tex. Baton Rouge, La Corpus Christi, Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La San Antonio, Te Shreveport, La. Tulsa, Okla.	Tex. 59 200 89 103 336 53 . U	883 38 31 40 107 53 70 203 33 U 132 71 105	19 11 15 55 21 22 65 10 U 37 26 31	134 11 9 3 17 5 7 47 5 U 14 8	71 4 4 12 7 3 13 4 U 15 3 5	33 1 1 9 3 1 8 1 U 3 3 3	79 4 11 4 21 4 U 12 9
E.N. CENTRAL Akron, Ohio Canton, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Gary, Ind. Grand Rapids, Mi Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohi W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans Kansas City, Kans Kansas City, Mo. Lincoln, Nebr. Minneapolis, Min Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	183 34 120 32 50 48 76 0 63 798 1 27 27 21 110 46	986 244 424 0 589 1366 82 U 33 511 123 777 25 35 90 90 52 29 102 54 94 105 105 105 105 105 105 105 105 105 105	37 44 18 U 5 11 6 7 50 8 27 4 8 6 10	86 3 1 U 5 14 15 6 U 2 4 1 1 10 2 3 2 3 3 5 5 6 4 1 2 7 2 12 6 10 2 9	34 11 10 29 44 11 10 - - 2 - 2 1 2 - - - - - - - - - - - - -	34 1 - U 3 4 4 5 1 U - 1 - 2 8 1 3 1 2 1 1 1 - 16 3 - 3 1 - 1 1 4 3	80 2 4 U 2 6 10 5 U 4 3 1 4 8 4 13 2 3 1 7 1 4 6 1 4 7 1 9 3 · 2 8	MOUNTAIN Albuquerque, N Boise, Idaho Colo. Springs, C Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, U Tucson, Ariz. PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawa Long Beach, Cal Los Angeles, Cal Pasadena, Calif. Portland, Oreg. Sacramento, Cal San Diego, Calif. San Francisco, C San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. TOTAL	. 40 . 40 . 200 . 55 . 104 . 209 . 209 . 160 . 124 . 124 . 1,652 . 133 . 37 . 11 . 65 . 147 . 201 . 201 . 20	647 73 29 41 130 20 101 20 77 95 1,172 10 88 29 9 360 0 20 U 132 97 U 119 21 22 49 6,943	191 21 10 4 224 51 3 29 8 8 24 17 309 - 18 7 7 U 26 31 1 26 9 9 1 22 9 26 1 27 28 29 29 20 20 31 4 4 20 31 4 4 4 4 4 4 4 4 4 4 5 1 5 1 6 1 6 1 6 1 7 1 8 1 8 1 8 1 8 1 8 1 8 1 8 1 8 1 8	95 14 1 7 7 3 121 4 4 3 10 0 0 10 1 1 1 1 1 1 1 1 1 1 1 1 1 1	31 1 1 4 5 1 1 8 2 2 7 7 2 4 4 - 2 15 5 5 U 0 6 5 5 U 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	19 2 2 2 2 2 2 2 1 7 7 - 1 2 2 7 2 9 2 2 U 3 4 4 U 2 2 2 2 1 7 2 2 1 7	64 10 3 2 8 12 2 7 1 10 9 124 10 3 3 3 3 4 - U17 16 U13 3 6 11 5 6 11 5 6 6 11 5 6 6 6 7 7 7 8 7 8 8 8 8 8 8 8 8 8 8 8 8

U: Unavailable. -: No reported cases.

^{*}Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of ≥100,000. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included.

[†] Pneumonia and influenza.

⁵ 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. ¹ Total includes unknown ages.

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