

Weekly

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National Diabetes Awareness Month — November 2004

November is National Diabetes Awareness Month. An estimated 18.2 million persons in the United States (i.e., 6.3% of the population) have diabetes. However, 5.2 million (29%) of these persons have not had their condition diagnosed. Persons with diabetes have a risk of premature death that is approximately two times greater than that of persons without the disease. From 1980 to 2002, the number of persons with diabetes in the United States more than doubled. In 2000, diabetes was the sixth leading cause of death in the United States and cost the nation more than \$132 billion dollars in healthcare expenditures. Additional information about diabetes.

During November, CDC, along with 59 state and territorial diabetes-control programs and other partners, will highlight activities that increase awareness about diabetes and women's health. More than 9.3 million women in the United States are now living with diabetes. CDC is a major partner in the *Initiative on Diabetes and Women's Health*, which will release a 30-minute video that emphasizes the healthy behaviors and coping skills that women have learned from adolescence through their older years to help manage their disease.

Throughout this month, *MMWR* will publish reports related to diabetes, including reports on diabetes among certain racial/ethnic populations, diabetes and obesity, diabetes and vision impairment, and psychological distress associated with the disease. Influenza and Pneumococcal Vaccination Coverage Among Persons Aged <u>>65</u> Years and Persons Aged 18–64 Years with Diabetes or Asthma — United States, 2003

Vaccination of persons at risk for complications from influenza and pneumococcal disease is a key public health strategy for preventing associated morbidity and mortality in the United States. Risk factors include older age and medical conditions that increase the risk for complications from infections. During the 1990–1999 influenza seasons, more than 32,000 deaths each year among persons aged \geq 65 years were attributed to complications from influenza infection (1). National health objectives for 2010 call for 90% influenza and pneumococcal vaccination coverage among noninstitutionalized persons aged \geq 65 years and 60% coverage among noninstitutionalized persons aged 18–64 years who have risk factors (e.g., diabetes or asthma) for complications from infections (2) (objective nos. 14.29a–d). To estimate influenza and pneumococcal vaccination coverage among these

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Notifiable Disease Morbidity and 122 Cities Mortality Data

Robert F. Fagan Deborah A. Adams Felicia J. Connor Lateka Dammond Rosaline Dhara Donna Edwards Patsy A. Hall Pearl C. Sharp populations, CDC analyzed data from the 2003 Behavioral Risk Factor Surveillance System (BRFSS) survey^{*}. This report summarizes the results of that analysis, which indicated that 1) influenza vaccination levels among adults aged 18–64 with diabetes or asthma, 2) pneumococcal vaccination levels among adults aged 18–64 years with diabetes, and 3) influenza and pneumococcal vaccination levels among adults aged ≥ 65 years all were below levels targeted in the national health objectives for 2010. Moreover, vaccination coverage levels varied among states for both vaccines and both age groups. Innovative approaches and adequate, reliable supplies of vaccine are needed to increase vaccination coverage, particularly among adults with high-risk conditions.

BRFSS is a state-based, random-digit-dialed telephone survey of the U.S. civilian, noninstitutionalized population aged \geq 18 years. All 50 states, the District of Columbia (DC), and three U.S. territories participate in the survey. Respondents were asked, "During the past 12 months, have you had a flu shot?" and "Have you ever had a pneumonia shot?" Persons with diabetes were defined as respondents who answered "yes" to the question, "Have you ever been told by a doctor that you have diabetes?" Women who were told that they had diabetes only during pregnancy were not defined as having diabetes. Participants were also asked, "Have you ever been told by a doctor, nurse, or other health professional that you had asthma?" Those who responded "yes" were then asked, "Do you still have asthma?" Respondents who answered affirmatively to both questions were classified as having asthma. For the 2003 BRFSS, the median state/area response rate was determined to be 53.2% (range: 34.4%-80.5%) by using the CASRO method. A total of 266,346 persons responded, of whom 207,735 (83.0%) were aged 18-64 years and 56,547 (17.0%) were aged ≥ 65 years. Among respondents aged 18-64 years, 17,084 (7.8%) reported having asthma and 12,412 (5.7%) reported having diabetes. Respondents with unknown influenza (0.3%) or pneumococcal (7.0%) vaccination status were excluded from the analysis. Vaccination levels were estimated for the 50 states, DC, Guam, Puerto Rico, and the U.S. Virgin Islands (USVI). Data were weighted by age, sex, and, in certain states/areas, race/ethnicity to reflect the estimated adult population. Statistical software was used to calculate point estimates and 95% confidence intervals.

In 2003, of respondents aged \geq 65 years, influenza vaccination coverage levels during the preceding 12 months ranged from 34.9% (USVI) to 80.3% (Minnesota), with a median of 69.9% (Table 1). Among respondents aged \geq 65 years, the

^{*} Conditions ascertained by BRFSS that are indicated for vaccination include asthma (indicated for influenza vaccine) and diabetes (indicated for influenza and pneumococcal vaccines).

	Influe ac	enza vaccination ar dults aged <u>≥</u> 65 yea	nong rs	Pneumococcal vaccination among adults aged ≥65 years			
Area	%	(95% CI*)	% change [†]	%	(95% CI)	% change	
Alabama	70.2	(±3.6)	5.4 [§]	61.4	(±4.0)	3.0	
Alaska	66.5	(±7.4)	-3.0	59.6	(±8.3)	-0.2	
Arizona	68.9	(±4.5)	-0.9	65.5	(±4.8)	-2.5	
Arkansas	71.0	(±3.2)	1.9	61.9	(±3.4)	3.1	
California	72.5	(±3.8)	1.0	65.2	(±4.1)	-1.5	
Colorado	74.2	(+3.9)	0.9	69.1	(+4.1)	0.9	
Connecticut	74.3	(± 3.0)	29	64.5	(+3.3)	0.0	
Delaware	70.0	(± 0.0)	-1.6	67.4	(+4 3)	3.0	
District of Columbia	63.0	(+6.2)	4.4	50.1	(+6.7)	2.2	
Florida	65.9	(+3.7)	9 <u>∩</u> §	64.5	(+3.8)	7 3§	
Georgia	67.0	(+3.3)	7.7§	60.5	(±0.0)	3.3	
Hawaii	71.6	(±3.5)	-23	44.5	(±3.0)	-15 0§	
Idaha	71.0	(±3.1)	-2.3 5 28	67.2	(± 3.4)	-10.0*	
Illinois	62.2	(± 3.3)	0.20	56.7	(± 3.4)	9.00	
Indiana	03.3	(± 4.2)	2.2	50.7 61 F	(± 4.3)	0.0	
	77.5	(± 3.0)	-0.2	71.4	(± 3.2)	0.3 E 28	
IOwa	77.0	(± 2.0)	4.0	71.4	(±2.9)	0.23	
Kansas	70.8	(±3.0)	2.1	60.3	(±3.3)	-1.8	
Кептиску	69.1	(±2.9)	3.3	59.6	(±3.3)	3.0	
Louisiana	68.3	(±3.3)	11.08	64.2	(±3.4)	7.98	
Maine	74.8	(±4.3)	1.0	64.8	(±4.7)	-1.9	
Maryland	68.4	(±4.2)	2.5	62.0	(±4.3)	-1.4	
Massachusetts	74.9	(±2.9)	2.4	69.4	(±3.1)	6.0 ⁹	
Michigan	67.5	(±3.6)	-0.2	62.7	(±3.8)	-0.3	
Minnesota	80.3	(±2.8)	3.7	73.0	(±3.3)	2.6	
Mississippi	69.0	(±3.2)	6.0 [§]	61.8	(±3.5)	2.8	
Missouri	69.9	(±3.7)	1.3	61.1	(±3.9)	0.3	
Montana	72.8	(±3.8)	5.1	69.1	(±4.0)	1.8	
Nebraska	73.6	(±2.7)	5.4 [§]	64.8	(±2.9)	3.5	
Nevada	60.0	(±5.9)	-0.4	63.2	(±6.0)	-1.8	
New Hampshire	73.9	(±3.2)	1.6	69.3	(±3.3)	5.5 [§]	
New Jersey	67.2	(±2.2)	-1.9	62.4	(±2.3)	-0.7	
New Mexico	72.4	(±2.8)	5.8 [§]	63.9	(±3.1)	1.2	
New York	68.0	(±3.4)	3.4	61.7	(±3.5)	-0.7	
North Carolina	68.8	(±2.9)	0.6	66.6	(±2.9)	3.6	
North Dakota	73.0	(±3.5)	-0.9	71.2	(±3.6)	-1.3	
Ohio	68.0	(± 4.2)	1.4	64.7	(± 4.5)	1.0	
Oklahoma	75.8	(±2.1)	3.1	68.6	(±2.3)	3.1	
Oregon	70.5	(± 3.3)	2.5	71.7	(± 3.4)	6.7 [§]	
Pennsylvania	69.2	(+3.4)	-1.4	66.1	(+3.6)	2.6	
Rhode Island	76.2	(+3.3)	2.5	69.3	(+3.6)	17	
South Carolina	69.3	(± 3.0)	-0.1	63.0	(± 3.2)	-1.8	
South Dakota	77.9	(± 2.0)	3.8§	63.7	(+2.8)	7.0§	
Tennessee	69.1	(± 4.5)	-2.6	60.8	(+4.8)	-0.6	
Tevas	67.7	(±4.0)	6.7§	62.0	(+3.2)	5.1§	
litab	7/ 8	(± 0.0)	3.7	66.2	(±0.2)	1.2	
Vormont	74.0	(±3.0)	0.5	66.1	(±2.4)	0.2	
Virginio	60.6	(± 3.0)	0.5	65.2	(± 3.4)	-0.2	
Weehington	09.0	(± 3.7)	4.5	0J.Z	(±3.9)	4.4 5.6 [§]	
	73.4	(± 1.7)	0.3%	00.0	(±1.0)	0.03	
	09.1	(±3.3)	3.3	03.0	(±3.0)	2.1	
	72.1	(±3.6)	-1.9	00./	(±3.8)	-3.9	
vvyoming	72.6	(±3.4)	1.9	/0.4	(±3.5)	2.2	
Guam	59.7	(±13.5)	15.6	37.0	(±13.1)	10.0	
Puerto Rico	40.2	(±4.1)	4.8	32.9	(±4.0)	7.88	
U.S. Virgin Islands	34.9	(±7.6)	2.7	31.6	(±7.6)	1.2	
Median	69.9			64.2			
Range	34.9-80.3			31.6–73.0			

TABLE 1. Percentage of persons aged \geq 65 years who reported receiving influenza vaccine during the preceding year and receiving pneumococcal vaccine ever, by area — Behavioral Risk Factor Surveillance System, United States, 2003

* Confidence interval. † Change in vaccination coverage from 2002 to 2003. § Statistically significant at p<0.05.

proportion reporting ever having received pneumococcal vaccine ranged from 31.6% (USVI) to 73.0% (Minnesota), with a median of 64.2%. Compared with 2002, a total of 41 and 38 states/areas experienced increases in influenza and pneumococcal coverage among those aged \geq 65 years, respectively; 11 of these increases were statistically significant for each vaccine.

Among adults aged 18-64 years with asthma or diabetes, substantial variation in vaccination coverage by area also was observed. For respondents with asthma, median influenza coverage was 34.0% and ranged from 22.5% (Puerto Rico) to 46.6% (Wyoming) (Table 2). Influenza vaccination rates among persons with asthma were higher among persons aged 50-64 years (median: 53.4%; range: 27.6%-74.9%) than among persons aged 18-49 years (median: 27.7%; range: 16.6%-41.1%). For respondents with diabetes, median influenza coverage was 49.0% and ranged from 26.5% (Puerto Rico) to 62.4% (South Dakota); the median pneumococcal coverage was 37.1% and ranged from 19.5% (Puerto Rico) to 58.2% (Montana). For persons with diabetes, vaccination rates were higher among those aged 50-64 years (for influenza, median: 56.5%; range: 23.7%-73.1% and for pneumococcal, median: 42.6%; range: 19.7%-68.1%) than among persons aged 18-49 years (for influenza, median: 37.8%; range: 22.2%–59.9% and for pneumococcal, median: 28.3%; range: 13.3%-56.7%).

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Editorial Note: The findings in this report indicate an increase in influenza and pneumococcal vaccination coverage for the majority of areas from 2002 to 2003 among adults aged \geq 65 years; however, coverage among persons indicated for these vaccinations remains below the national health objectives for 2010. In addition, almost half of the states reported >50% influenza coverage levels for participants aged 18–64 years with diabetes; however, the median coverage level of influenza vaccination among participants with asthma and the median coverage level of pneumococcal vaccines among participants with diabetes were below the 2010 target of 60% for noninstitutionalized adults at high risk. Among respondents with asthma and diabetes, those aged 18–49 years had substantially lower vaccination coverage than those aged 50–64 years.

Lack of awareness of the need for vaccination is common among adults aged <65 years with high-risk conditions, such as diabetes or asthma. In a 2003 survey, approximately 75% of unvaccinated persons aged 18–64 years with diabetes reported that they were unaware of the need for influenza vaccine (CDC, unpublished data, 2003). Although use of preventive health services by adults with diabetes has increased since 1995 (3), a substantial proportion of generalist and subspecialist physicians did not strongly recommend influenza and pneumococcal vaccinations to their patients who are elderly or at high risk (4). Low vaccination rates among persons with high-risk conditions might reflect the challenge of targeting patients for vaccinations on the basis of high-risk conditions instead of age (1). Although a majority of patients seen by subspecialists might be those who most need vaccination, subspecialists might not perceive the provision of preventive services as their role. Primary care physicians and subspecialists should work together to ensure that persons at high risk receive appropriate vaccinations. In addition, strategies to increase awareness among young adults of the need for vaccinations could be emphasized by diabetes- and asthmacare programs (3,5). The Diabetes Quality Improvement Project, a collaborative effort between public and private organizations to improve preventive care for persons with diabetes, has been ongoing since 1995 (6); this effort is one possible reason for the higher influenza vaccination rates among those with diabetes compared with those with asthma.

The findings in this report are subject to at least three limitations. First, vaccination status (influenza and pneumococcal) was based on self-report and not validated. The validity of self-reported pneumococcal vaccination is lower than that of influenza vaccination (7). Second, the median BRFSS response rate (53.2%) in this survey was low. BRFSS results have been compared with results from the National Health Interview Survey (NHIS), a household-based, face-to-face interview survey with higher response rates. Comparisons demonstrate similar trends and subgroup differences; however, BRFSS vaccination estimates are consistently higher than NHIS estimates (8). Finally, because BRFSS does not systematically assess other medical conditions for which influenza and pneumococcal vaccines are recommended, vaccine coverage for all persons with high-risk conditions was not examined.

The variation in influenza and pneumococcal vaccination coverage observed among areas suggests that vaccination coverage can be improved. Previous studies have indicated that organizational changes, such as nurse standing orders, combined with teamwork and collaboration, are effective intervention measures for increasing adult vaccination services (9). Effective measures to promote the use of such measures are needed for vaccination rates to increase.

Because of the 2004 influenza vaccine shortage, vaccine providers have been asked to direct available inactivated influenza vaccine to persons with chronic conditions, such as diabetes and asthma, and other priority groups. Further analysis of influenza vaccine coverage data will be needed to assess

TABLE 2. Percentage of persons aged 18–64 years with asthma or diabetes who reported receiving influenza vaccine during the pre-ceding year and persons aged 18–64 years with diabetes reporting receiving pneumococcal vaccine ever, by area — Behavioral Risk Factor Surveillance System, United States, 2003

	Influenza vaccination among adults aged 18–64 years with asthma Influenza vaccination among adults aged 18–64 years with diabetes		ccination among d 18–64 years diabetes	Pneumococcal vaccinatio among adults aged 18–64 yo with diabetes		
Area	%	(95% CI*)	%	(95% CI)	%	(95% CI)
Alabama	33.8	(25.7-41.8)	47.4	(39.4-55.4)	34.8	(26.9-42.7)
Alaska	38.7	(29.0-48.3)	50.4	(34.9–65.9)	41.2	(25.3–57.0)
Arizona	33.9	(24.5-43.4)	54.4	(41 8–66 9)	33.8	(22.0-45.6)
Arkansas	40.2	(33.5-46.8)	45.2	(37.8–52.6)	30.7	(23.8 - 37.7)
California	28.7	(22.9-34.5)	40.2	(31.9 - 48.5)	29.5	(21.7 - 37.3)
Colorado	39.0	(32.5-45.5)	52.7	(43.2-62.2)	41.2	(31.2–51.2)
Connecticut	39.9	(344 - 454)	54.8	(46.3-63.3)	33.5	(25.0-42.0)
Delaware	34.8	(27.5-42.0)	44.0	(35.2-52.8)	27.9	(20.5 - 35.2)
District of Columbia	24.7	(16.8–32.7)	41.8	(28.8–54.7)		(2010 0012)
Florida	28.8	(20.1 - 37.6)	43.4	(34.0–52.7)	43.7	(34.2-53.3)
Georgia	31.6	(24.0-39.2)	38.2	(31.9–44.4)	26.2	(21.0-31.5)
Hawaii	42.0	(33.8–50.1)	57.5	(48.6–66.3)	26.4	(18.3–34.4)
Idaho	31.3	(25.2–37.4)	54.6	(45.9–63.4)	38.6	(30.2–47.0)
Illinois	32.4	(26.1–38.7)	38.1	(29.8 - 46.4)	29.4	(21.3-37.5)
Indiana	33.7	(28.4–39.0)	46.6	(40.0-53.2)	40.5	(34.0-47.1)
lowa	31.3	(24.3-38.3)	62.2	(54.3 - 70.1)	48.5	(40.2 - 56.9)
Kansas	30.4	(24.2–36.6)	49.8	(41.4–58.2)	33.9	(25.9–42.0)
Kentucky	29.7	(24.1-35.2)	46.6	(40.0-53.3)	33.8	(27.6 - 40.0)
Louisiana	36.6	(29.7-43.6)	40.9	(34.4–47.4)	31.6	(25.6-37.7)
Maine	39.3	(31.4–47.3)	49.0	(38.4–59.7)	35.0	(24.9 - 45.2)
Maryland	38.4	(30 7-46 0)	46.6	(37 6–55 6)	38.0	(28.9-47.0)
Massachusetts	36.8	(31.9–41.6)	49.7	(42.2–57.3)	39.1	(31.5-46.8)
Michigan	34.3	(27.8–40.9)	42.1	(34.6 - 49.7)	38.0	(30.4-45.5)
Minnesota	40.1	(32.4–47.8)	56.3	(47.2–65.4)	33.6	(25.1–42.2)
Mississippi	30.4	(23.5–37.4)	39.8	(33.6-46.1)	22.6	(17.0–28.3)
Missouri	31.9	(24.4-39.5)	48.6	(39.0–58.1)	35.2	(26.2 - 44.2)
Montana	46.6	(38.4–54.8)	58.8	(48.7–69.0)	58.2	(47.7–68.7)
Nebraska	43.1	(36.6–45.1)	57.0	(48.8–65.2)	37.7	(29.7–45.8)
Nevada	27.8	(18.6–37.1)	29.0	(17.7–40.3)	40.0	(26.4–53.6)
New Hampshire	36.8	(31.0-42.6)	61.9	(54.2–69.6)	50.6	(42.6–58.7)
New Jersey	31.6	(27.4–35.8)	41.9	(36.4–47.4)	29.6	(24.6–34.6)
New Mexico	39.7	(33.2–46.1)	61.3	(53.4–69.2)	46.1	(37.6–54.6)
New York	38.6	(32.9–44.3)	53.5	(45.7–61.4)	43.6	(35.5–51.7)
North Carolina	34.0	(27.9–40.1)	46.1	(39.7–52.4)	38.3	(31.9–44.7)
North Dakota	38.8	(30.6–46.9)	56.3	(45.4–67.3)	36.4	(26.1–46.7)
Ohio	30.4	(23.3–37.5)	38.0	(29.6–46.5)	41.8	(32.3–51.2)
Oklahoma	38.0	(32.8–43.3)	53.9	(48.1–59.6)	41.3	(35.6–46.9)
Oregon	34.4	(28.2–40.7)	54.5	(45.3–63.6)	48.4	(38.9–57.9)
Pennsylvania	33.6	(26.5–40.6)	59.3	(50.7–67.9)	37.1	(28.2-45.9)
Rhode Island	42.0	(35.7-48.3)	58.9	(49.4–68.3)	46.6	(37.2–56.0)
South Carolina	38.9	(32.4–45.4)	52.1	(45.9–58.2)	34.9	(29.1–40.8)
South Dakota	45.8	(37.6–54.0)	62.4	(54.9–69.9)	37.7	(30.1–45.3)
Tennessee	32.8	(24.9-40.8)	47.4	(38.6-56.1)	28.1	(21.0-35.3)
Texas	31.9	(25.8-37.9)	40.8	(34.9-46.7)	29.2	(23.5-34.8)
Utah	30.9	(23.4-38.3)	53.1	(42.5-63.7)	53.4	(42.6-64.1)
Vermont	30.7	(24.5-36.9)	56.0	(47.2-64.7)	40.7	(31.8-49.6)
Virginia	32.9	(25.5-40.2)	45.1	(37.1–53.0)	34.8	(27.0-42.5)
Washington	36.5	(33.4-39.7)	50.5	(46.2-54.8)	43.8	(39.4-48.1)
West Virginia	37.5	(30.5-44.5)	52.4	(44.8–60.0)	40.3	(32.9–47.7)
Wisconsin	34.3	(27.2-41.3)	58.0	(47.9–68.2)	55.5	(45.3–65.8)
Wyoming	46.6	(39.2–54.0)	47.3	(38.6–56.0)	46.0	(37.2–54.9)
Guam	_		_		_	
Puerto Rico	22.5	(17.3–27.8)	26.5	(20.3–32.7)	19.5	(13.9–25.1)
U.S. Virgin Islands			28.2	(18.7–37.7)	_	
Median	34.0		49.0		37.1	
Range	22.5-46.6		26.5-62.4		19.5–58.2	

* Confidence interval. [†]Number of respondents too small for meaningful analysis.

the impact of this shortage on influenza vaccine coverage and efforts to redirect vaccine to persons at greatest risk for influenza complications. Ensuring adequate amounts of influenza vaccine is critical if vaccination rates of persons at high risk are to continue improving. Pneumococcal vaccine supplies appear to be adequate to meet expected demand. Pneumococcal vaccination should be encouraged for populations at high risk, both to reduce the risk for invasive pneumococcal disease itself and to reduce complications of influenza infection.

References

- CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 2004;53(No. RR-6).
- US Department of Health and Human Services. Healthy People 2010, 2nd ed. With understanding and improving health and objectives for improving health (2 vols.). Washington, DC: US Department of Health and Human Services; November 2000.
- CDC. Preventive-care practices among persons with diabetes—United States, 1995 and 2001. MMWR 2002;51:965–9.
- 4. Nichol KL, Zimmerman R. Generalist and subspecialist physicians' knowledge, attitudes, and practices regarding influenza and pneumococcal vaccinations for elderly and other high-risk patients: a nationwide survey. Arch Intern Med 2001;161:2702–8.
- Task Force on Community Preventive Services. Recommendations regarding interventions to improve vaccination coverage in children, adolescents, and adults. Am J Prev Med 2000;18:92–6.
- Fleming BB, Greenfield S, Engelgau MM, Pogach LM, Clauser SB, Parrott MA. The Diabetes Quality Improvement Project: moving science into health policy to gain an edge on the diabetes epidemic. Diabetes Care 2001;24:1815–20.
- MacDonald R, Baken L, Nelson A, Nichol KL. Validation of self-report of influenza and pneumococcal vaccination status in elderly outpatients. Am J Prev Med 1999;16:173–7.
- Nelson DE, Powell-Griner E, Town M, Kovar MG. A comparison of national estimates from the National Health Interview Survey and the Behavioral Risk Factor Surveillance System. Am J Public Health 2003; 93:1335–41.
- Stone EG, Morton SC, Hulscher ME, et al. Interventions that increase use of adult immunization and cancer screening services: a meta-analysis. Ann Intern Med 2002;136:641–51.

Influenza Vaccination and Self-Reported Reasons for Not Receiving Influenza Vaccination Among Medicare Beneficiaries Aged <u>></u>65 years — United States, 1991–2002

Annual influenza vaccination of the U.S. elderly population has been demonstrated as safe and effective in reducing the risks of illness, hospitalization, and death (1). The Medicare Current Beneficiary Survey (MCBS) has measured annual influenza vaccination rates since 1991; the latest data available are for the 2001–02 influenza season. Since 1996, self-reported reasons for not receiving influenza vaccine also have been measured. During 1991–2002, MCBS indicated a steady upward trend in vaccination coverage among Medicare beneficiaries, with the exception of the 2000–01 influenza season, when vaccine distribution was delayed. The most frequently cited reasons for not receiving influenza vaccine were 1) not knowing that influenza vaccination was needed and 2) concerns that vaccination might cause influenza or side effects. During the 2000–01 influenza season, vaccine shortage or unavailability was noted for the first time as an important reason for nonvaccination. Further efforts are needed to educate the elderly regarding the benefits of influenza vaccination and to address any concerns regarding the safety of the vaccine.

MCBS is a nationally representative survey of the Medicare population, conducted by the Centers for Medicare & Medicaid Services (CMS). Beneficiaries sampled from Medicare enrollment files (or appropriate proxies) are interviewed in person. Primary sampling units (PSUs) consist of persons in 107 geographic areas chosen to represent the nation; beneficiaries residing in these PSUs are selected by systematic random sampling within age strata. Data for this report were analyzed by using statistical software to account for sampling weights in calculating point estimates of proportions; analyses were restricted to Medicare beneficiaries aged ≥ 65 years who resided in a noninstitutional setting.

Each year, MCBS asks respondents, "Did you have a flu shot for last winter?" The percentage reporting receipt of influenza vaccination increased each influenza season from 1991–92 through 1999–2000, and especially in 1993–94 (Figure 1), when influenza vaccination first became a Medicare benefit. However, during the 2000–01 influenza season, the vaccination rate declined instead of maintaining an annual increase; 67.0% (20.5 million of 30.6 million



FIGURE 1. Percentage of Medicare beneficiaries aged ≥65 years who reported influenza vaccination, by influenza season — United States, 1991–2002

a-ware: *adj*

(ə-'wâr) 1 : marked by comprehension, cognizance, and perception; see

also MMWR.



know what matters.



Medicare beneficiaries aged ≥ 65 years living in the community) reported receiving the vaccine, compared with a record high of 70.0% (21.2 million out of 30.3 million) in 1999– 00. For the 2001–02 influenza season, 68.8% (21.3 million of 31.0 million) reported receiving influenza vaccine.

The MCBS also asks about reasons for not getting influenza vaccination. The question asked is "Why didn't you get a flu shot for last winter?" Respondents are free to give any reason or reasons, with open-ended responses recorded by interviewers into prespecified categories. This question was omitted for the 1999–00 influenza season.

The leading reasons for nonvaccination reported for 1997–98, 1998–99, 2000–01, and 2001–02 were not knowing that influenza vaccination was needed and concerns that vaccination might cause influenza or side effects (Figure 2). In 2000–01, for the first time, one of the leading reasons was that vaccine was unavailable or in short supply. For the 2000–01 season,

12.7% of unvaccinated respondents reported vaccine unavailability as a reason for not receiving influenza vaccine. This equates to approximately 1.25 million persons, or 4.2% of the total elderly Medicare population living in the community, which amounts to roughly the difference between the expected annual increase and the actual decline for 2000–01 in selfreported influenza vaccination. By contrast, during the 2001–02 influenza season, an estimated 7.5% of unvaccinated respondents (approximately 707,000 persons, or 2.3% of the total elderly Medicare population living in the community) reported vaccine unavailability as a reason for nonvaccination. **Reported by:** *GS Adler, MPhil, Office of Research, Development, and Information, Centers for Medicare & Medicaid Svcs. CA Winston, PhD, Immunization Svcs Div, National Immunization Program, CDC.*

Editorial Note: For the 2000–01 influenza season, production delays created shortages of influenza vaccine, especially at the beginning of the vaccination period (i.e., October and

FIGURE 2. Percentage of Medicare beneficiaries aged \geq 65 years who reported reasons for not receiving influenza vaccination, by reason — United States, 1997–98, 1998–99, 2000–01, and 2001–02 influenza seasons



* Not listed as a reason before 2000-01 influenza season.

November), when demand was greatest (2). Delays in vaccine production continued for the 2001-02 influenza season but were considered less severe (3). The 2000-01 decline and subsequent rebound of vaccination coverage in 2001-02 was observed by other government health surveys (4-6). However, the self-reported MCBS data establish an association between vaccine shortages and reduced vaccination rates among the Medicare population aged ≥ 65 years, one of the groups at high risk for influenza complications.

The findings in this report are subject to at least two limitations. First, vaccination status during the preceding influenza season is self-reported and subject to recall and social desirability bias. Second, the results are subject to survivor bias (i.e., persons who died could not be interviewed about their vaccination status the previous winter). Finally, participant reasons for nonvaccination are categorized during the interview, rather than recorded verbatim. Although "other" reasons are captured as

a category and reviewed for retrospective categorizing of responses, interviewer coding might result in misclassification or in a different distribution of reasons than would be obtained by providing respondents a list of answers from which to choose.

Consequences from the vaccine shortage in 2000-01 were limited because of the mildness of the influenza virus that season (7). CDC has previously estimated that for each 1 million elderly persons vaccinated, approximately 900 deaths and 1,300 hospitalizations are prevented (8). Influenza vaccination rates are used as indicators of progress toward achieving the national health objectives for 2010. CMS, together with CDC, has conducted a long-term, structured campaign to promote the benefits of vaccination to Medicare beneficiaries and to improve provider performance. These efforts have resulted in large increases in vaccination during the preceding decade. However, even the strongest efforts of government agencies to promote vaccination are subject to the constraints of limited vaccine supply. During the current influenza vaccine shortage, vaccine is prioritized for populations at high risk, including the elderly.

During 1997–2002, other reasons for nonvaccination were cited more often than reduced availability of vaccine. The most common reasons for nonvaccination were lack of knowledge about the need for vaccination and misconceptions about influenza vaccination and disease or side effects (9). These reasons remain important modifiers of elderly Medicare beneficiaries' behavior and can be further addressed through communications about influenza vaccination. Evidence-based strategies should be developed and used to 1) educate the public and vaccination providers regarding the benefit of influenza vaccine for the elderly and 2) address concerns about the safety and efficacy of the vaccine.

References

- US Preventive Services Task Force. Guide to clinical preventive services. 2nd ed. Washington, DC: US Department of Health and Human Services; 1996.
- 2. US General Accounting Office. Flu vaccine: supply problems heighten need to ensure access for high-risk people (report GAO-01-624). Washington, DC: US General Accounting Office; 2001.
- 3. Fukuda K, O'Mara D, Singleton J. How the delayed distribution of influenza vaccine created shortages in 2000 and 2001. Pharmacy and Therapeutics 2002;27:235–42.
- 4. CDC. Influenza and pneumococcal vaccination levels among persons aged ≥65 years—United States, 2001. MMWR 2002;51:1019–24.
- National Center for Health Statistics. Early release: Figure 4.1. Percent of adults aged 18 years and over who had received an influenza shot during the past 12 months, by age group and quarter, 1997–2004. Hyattsville, MD: US Department of Health and Human Services, CDC; 2004. Available at http://www.cdc.gov/nchs/data/nhis/earlyrelease/ 200409_04.pdf.
- 6. CDC. Influenza vaccination coverage among adults aged ≥50 years and pneumococcal vaccination coverage among persons aged ≥65 years— United States, 2002. MMWR 2003;52:987–92.

- CDC. Update: influenza activity—United States and worldwide, 2000–01 season, and composition of the 2001–02 influenza vaccine. MMWR 2001;50:466–70.
- CDC. Updated recommendations from the Advisory Committee on Immunization Practices in response to delays in supply of influenza vaccine for the 2000–01 season. MMWR 2000;49:888.
- 9. CDC. Reasons reported by Medicare beneficiaries for not receiving influenza and pneumococcal vaccinations—United States, 1996. MMWR 1999;48:886–90.

Acute Hepatitis B Among Children and Adolescents — United States, 1990–2002

Since the 1991 adoption of a comprehensive strategy to eliminate hepatitis B virus (HBV) transmission in the United States (1), the incidence of acute hepatitis B cases has declined steadily. Declines have been greatest among children born after the 1991 recommendations for universal infant hepatitis B vaccination were implemented. In 1995, the elimination strategy was expanded to include routine vaccination of all adolescents aged 11-12 years and, in 1999, to include children aged ≤ 18 years who had not been vaccinated previously (2). To describe the epidemiology of acute hepatitis B in children and adolescents in the United States, CDC analyzed notifiable disease surveillance data collected during 1990-2002 and data collected during 2001-2002 through enhanced surveillance of reported cases of acute hepatitis B in children born after 1990. This report summarizes the results of that analysis, which indicated that the rate of acute hepatitis B in children and adolescents decreased 89% during 1990-2002 and that racial disparities in hepatitis B incidence have narrowed. Many confirmed cases in persons born after 1990 occurred among international adoptees and other children born outside the United States. Continued implementation of the hepatitis B elimination strategy and accurate surveillance data to monitor the impact of vaccination are necessary to sustain the decline of acute hepatitis B among children.

Cases of acute hepatitis B were reported weekly to CDC by all 50 states and the District of Columbia. Acute hepatitis B rates were calculated per 100,000 population by using population denominators from the U.S. Census Bureau. Acute hepatitis B was defined as an acute illness with 1) discrete onset of symptoms and jaundice or elevated serum aminotransferase levels and 2) laboratory evidence of either IgM antibody to hepatitis B core antigen (IgM anti-HBc) or hepatitis B surface antigen (HBsAg). Since March 2001, CDC has conducted enhanced hepatitis B surveillance, contacting states to confirm all reported cases of acute hepatitis B in persons born after 1990. State surveillance staff members were asked to verify each of the items in the case definition and provide information regarding vaccination history and country of birth. If errors were identified during this process, states were asked to correct the information in an updated submission to CDC.

National Surveillance

During 1990–2002, a total of 13,829 cases of acute hepatitis B were reported in the United States among persons aged \leq 19 years. The incidence of reported cases declined steadily during this period, from 3.03 per 100,000 population in 1990 to 0.34 in 2002, representing a decline of 89%. The incidence among adolescents aged 15–19 years was consistently higher than the incidence among younger age groups (Figure 1), ranging from 8.69 per 100,000 population in 1990 to 1.13 in 2002. Children and adolescents in all age groups experienced steep declines in incidence during 1990–2002; incidence declined 94% among children aged 0–4 years, 92% among children aged 5–9 years, 93% among those aged 10–14 years, and 87% among adolescents aged 15–19 years.

Among children and adolescents aged \leq 19 years in 1990, incidence per 100,000 population was highest among Asian/ Pacific Islanders (A/PIs) (6.74) and blacks (4.29); whites had the lowest race-specific incidence (1.39). Differences in incidence between whites and A/PIs and between whites and blacks were 5.34 and 2.90, respectively. From 1990 to 2002, rates declined 92% among A/PIs, 88% among whites, 88% among blacks, and 84% among American Indians/Alaskan Natives (AI/ANs) (Figure 2). In 2002, the highest incidence per 100,000 population was among A/PIs (0.55), followed by blacks (0.51), AI/ANs (0.43), and whites (0.16); since 1990, differences in incidence between whites and A/PIs and whites and blacks declined by 93% and 88%, respectively.





* Per 100,000 population.

FIGURE 2. Rate* of acute hepatitis B in persons aged \leq 19 years, by race and year — United States, 1990–2002



* Per 100,000 population.

Case Investigations

Follow-up investigations conducted by CDC and state and local health departments verified 19 case reports from 2001 and 2002 as cases of acute hepatitis B among children born after 1990 (Table). Of the verified case reports, 12 (60%) involved males, eight (42%) involved children aged <2 years, and 11 (58%) involved children born in the United States. Seven (37%) reported race as A/PI, five (26%) as white, four (21%) as black, and three (16%) as unknown. Eight (42%)

TABLE. Acute hepatitis B cases* among U.S. residents born after 1990, by year and selected characteristics — United States, 2001–2002

Reporting year	Age	Race	Birth country or continent	International adoptee
2001	8 yrs	White	United States	No
	1 yr	Asian/Pacific		
		Islander (A/PI)	United States	No
	7 yrs	A/PI	Asia	No
	1 yr	A/PI	United States	No
	5 yrs	A/PI	China	Yes
	8 yrs	White	Bulgaria	Yes
	5 yrs	White	United States	No
	9 yrs	A/PI	United States	No
	10 mos	Unknown	Russia	Yes
	1 yr	A/PI	Phillipines	Yes
	5 yrs	White	United States	No
	9 mos	White	Ukraine	Yes
	10 mos	Black	United States	No
	5 yrs	Black	United States	No
2002	7 yrs	Black	United States	No
	9 mos	A/PI	Vietnam	Yes
	1 yr	Unknown	Tanzania	Unknown
	9 yrs	Black	United States	No
	11 yrs	Unknown	United States	No

* Confirmed by follow-up investigation.

cases were reported in children born outside the United States, including six international adoptees (32%). Receipt of ≥ 1 dose of hepatitis B vaccine was confirmed in three (16%) cases. Vaccination status was unknown for 12 cases (63%).

Reported by: State and local health departments. C Shepard, MD, L Finelli, DrPH, B Bell, MD, J Miller, MPH, Div of Viral Hepatitis, National Center for Infectious Diseases, CDC.

Editorial Note: The incidence of acute hepatitis B cases in U.S. children and adolescents decreased during the era of universal childhood vaccination. This decline coincided with an increase in hepatitis B vaccination coverage among children aged 19–35 months, from 16% in 1992 to 90% in 2002, and among adolescents aged 13–15, from nearly 0 in 1992 to 67% in 2002 (*3*,*4*).

Declines in incidence were observed for children of all races. including A/PIs, whose rates historically have been higher than the national average. Because of the disproportionate burden of hepatitis B in A/PI communities, A/PI children were among the first groups for whom hepatitis B vaccination was recommended (3). The reduction of the disparity between A/PIs and other children is consistent with recent observations noting a decline in seroprevalence of HBV infection and successful implementation of routine hepatitis B vaccination among Asians who have recently immigrated to the United States (5). However, of the 11 verified cases during 2001-02 of acute hepatitis B among children born in the United States, three (27%) involved A/PIs (Table). Although the national origins of these children's household members are unknown, the substantial proportion of A/PIs suggests that horizontal transmission of HBV among first-generation Asians might be a persistent problem (6).

The higher incidence among older adolescents (aged 15–19 years) likely is attributable to their having been born before universal infant hepatitis B vaccination was recommended in 1991. Incidence among older adolescents is expected to decline further as the vaccinated cohort ages and as 1999 recommendations to vaccinate all previously unvaccinated persons aged 0–18 years are fully implemented. The expected decline in rates among adolescents also might be augmented by laws in 32 states requiring proof of hepatitis B vaccination before entry into middle school (7).

Follow-up information obtained through surveillance of reported cases suggests that children born outside the United States, especially international adoptees, represent a substantial proportion of cases. Cases of acute hepatitis B among international adoptees might result from undervaccination and increased risk for exposure while living in areas with high prevalence of chronic HBV infection. International adoptees are exempt from U.S. regulations^{*} that bar entry to immigrants without documentation of hepatitis B vaccination. Studies have demonstrated that international adoptees exhibit low rates of protective titers of antibodies to vaccine-preventable diseases upon arrival in the United States, including adoptees with written evidence of age-appropriate vaccination provided by the birth country (8,9). Appropriate evaluation and remediation of the immunization status of international adoptees has been promoted through national guidelines (10); however, the extent to which these guidelines have been implemented is unknown.

Despite the decline in acute hepatitis B cases among children in the United States, the presence of confirmed cases highlights the importance of infant vaccination and timely completion of the 3-dose vaccination series. The vaccination series should be started at birth, preferably before the newborn is discharged from the hospital. Infants born to women who are HBsAg positive or who have not had prenatal HBsAg testing should receive the first fose of hepatitis B vaccine within 12 hours of birth (*I*). Beginning the vaccination series at birth decreases the risk for perinatal HBV transmission and predicts successful completion of the series.

Although enhanced surveillance data from verified case reports suggest that international adoptees and other children born outside the United States might particularly benefit from future prevention efforts, many case reports lacked risk factor information. As the incidence of acute hepatitis B among children and adolescents declines, accurate surveillance data become increasingly important to monitor the effect of immunization recommendations. Continued efforts of local, state, and national surveillance staff to improve data quality are critical to eliminating HBV transmission in the United States.

*U.S. Code title 8, chapter 12, subchapter II, Part II, §1182, (a)(1)(C).

References

- 1. CDC. Hepatitis B virus: a comprehensive strategy for eliminating transmission in the United States through universal childhood vaccination: recommendations of the Immunization Practices Advisory Committee (ACIP). MMWR 1991;40(No. RR-13).
- CDC. Update: recommendations to prevent hepatitis B virus transmission—United States. MMWR 1999;48:33–4.
- CDC. Achievements in public health: hepatitis B vaccination, United States, 1982–2002. MMWR 2002;51:549–52,563.
- 4. CDC. National immunization survey. Atlanta, GA: US Department of Health and Human Services, CDC; 2004. Available at http:// www.cdc.gov/nip/coverage/#NIS.
- 5. Fiore A, Neeman R, Lee S, et al. Seroprevalence of hepatitis B virus (HBV) infection among Asian immigrants and their US-born children in Georgia. In: Abstracts of the 41st Annual Meeting of the Infectious Diseases Society of America, San Diego, CA; October 2003.
- Franks AL, Berg CJ, Kane MA, et al. Hepatitis B virus infection among children born in the United States to southeast Asian refugees. N Engl J Med 1989;321:1301–5.

- Immunization Action Coalition. Hepatitis B prevention mandates. St. Paul, MN: Immunization Action Coalition; 2004. Available at http:// www.immunize.org/laws/hepb.htm.
- Miller LC. Internationally adopted children—immunization status. Pediatrics 1999;103:1078.
- Hostetter MK. Infectious diseases in internationally adopted children: findings in children from China, Russia, and Eastern Europe. Adv Pediatr Infect Dis 1999;14:147–61.
- 10. CDC. General recommendations on immunization: recommendations of the Advisory Committee on Immunization Practices (ACIP) and the American Academy of Family Physicians (AAF). MMWR 2002;51(No. RR-2):19–21.

Blood Mercury Levels in Young Children and Childbearing-Aged Women — United States, 1999–2002

Exposure to high levels of mercury (Hg) can cause neurologic and kidney disorders (1-3). Because methylated Hg (methyl-Hg) in the aquatic environment accumulates in animal tissues up the food chain, persons in the United States can be exposed by eating freshwater fish, seafood, and shellfish. Exposure of childbearing-aged women is of particular concern because of the potential adverse neurologic effects of Hg in fetuses. To determine levels of total blood Hg in childbearing-aged women and in children aged 1-5 years in the United States, CDC's National Health and Nutrition Examination Survey (NHANES) began measuring blood Hg levels in these populations in 1999. This report summarizes NHANES results for 1999-2002 and updates previously published information (4,5). The findings confirmed that blood Hg levels in young children and women of childbearing age usually are below levels of concern. However, approximately 6% of childbearing-aged women had levels at or above a reference dose, an estimated level assumed to be without appreciable harm (\geq 5.8 µg/L). Women who are pregnant or who intend to become pregnant should follow federal and state advisories on consumption of fish.

NHANES is a continuous survey of the health and nutritional status of the civilian, noninstitutionalized U.S. population; data are released and reported in 2-year cycles (6). Each participant undergoes a household interview and a physical examination. During the physical examination, blood is collected by venipuncture from all persons aged ≥ 1 year. For this analysis, whole-blood specimens were analyzed for total and inorganic Hg for children aged 1–5 years and women aged 16–49 years by automated, cold-vapor atomic absorption spectrophotometry in CDC's inorganic toxicology laboratory. The analytic method detection limit was 0.14 µg/L (ppb) for total Hg and 0.4 µg/L (ppb) for inorganic Hg (7). Blood Hg levels less than the limit of detection were assigned a value equal to the detection limit divided by the square root of 2 for the calculation of geometric mean (GM) values.

During 1999–2002, the GMs of total blood Hg concentrations for all childbearing-aged women and for children aged 1–5 years were 0.92 µg/L and 0.33 µg/L, respectively; the 95th percentiles of blood Hg for women and children were 6.04 µg/L and 2.21 µg/L, respectively (Table 1). The percentage of all women aged 16–49 years with Hg levels \geq 5.8 µg/L (the Environmental Protection Agency's [EPA] Reference Dose [RfD]) was 5.66% (95% confidence interval [CI] = 4.04–7.95) (Table 2).

Among children aged 1–5 years, the estimated percentage who had blood Hg levels \geq 5.8 µg/L during 1999–2002 could not be reported because the observed percentage was too low for the given sample size to calculate a statistically reliable national population estimate. Almost all inorganic blood Hg levels were undetectable, indicating that total blood Hg greater than or equal to the EPA RfD mostly reflected exposure to organic Hg (especially methyl-Hg).

Reported by: *RL Jones, PhD, T Sinks, PhD, SE Schober, PhD, M Pickett, MPH, National Center for Environmental Health; National Center for Health Statistics, CDC.*

Editorial Note: This report updates NHANES 1999–2000 estimates of blood Hg levels (5), the first nationally representative estimates of U.S. women's and children's exposures to Hg based on biologic measures. The findings indicate that blood Hg levels in young children and childbearing-aged women usually are below levels of concern.

Among childbearing-aged women, for the 4-year period 1999–2002, estimates of the GM of blood Hg and the proportion with levels \geq 5.8 µg/L were lower than estimates for the 2-year period 1999–2000, reflecting apparent declines in these values for the 2-year period 2001–2002. However, when these differences were evaluated by comparing estimates for the two 2-year periods, the declines were not statistically significant: the GM of blood Hg for 2001–2002 was 0.83 µg/L (CI = 0.73–0.93), compared with 1.02 µg/L (CI = 0.80–1.24) for 1999–2000, and the percentage of women with blood Hg levels \geq 5.8 µg/L was 3.9% in 2001–2002 (CI = 2.40–6.43), compared with 7.8% in 1999–2000 (CI = 4.70–12.83). At least 2 more years of data are needed to best determine whether Hg exposure has declined among women of childbearing age in the United States.

Although NHANES data are released and often analyzed as 2-year periods, the estimates of blood Hg levels for 1999– 2002 are the most reliable estimates of current exposure. The 4-year period provides greater geographic coverage, and estimates and sample errors are more stable, thus reducing vari-

				Selected percentile (95% CI)					
Variable	No.	GM	(95% CI*)	5th	(95% CI)	10th	(95% CI)	25th	(95% CI)
Women									
Race/Ethnicity									
Mexican American	1,106	0.74	(0.64-0.84)	0.10	(0.08–0.15)	0.17	(0.12-0.23)	0.34	(0.27-0.45)
White, non-Hispanic	1,377	0.87	(0.76-0.99)	0.09	(0.08–0.10)	0.15	(0.13–0.18)	0.37	(0.34-0.45)
Black, non-Hispanic	794	1.18	(1.00–1.36)	0.17	(0.12–0.25)	0.30	(0.24–0.38)	0.60	(0.55–0.73)
Age group (yrs)									
16–29	2,004	0.68	(0.60-0.76)	0.08	(0.07-0.09)	0.11	(0.09-0.14)	0.29	(0.25-0.37)
30–49	1,633	1.10	(0.97-1.24)	0.13	(0.10-0.16)	0.24	(0.20-0.29)	0.52	(0.45-0.60)
Pregnancy status									
Pregnant	629	0.75	(0.60-0.90)	0.08	([†] –0.10)	0.10	(0.08-0.20)	0.32	(0.24-0.44)
Not pregnant	2,978	0.94	(0.84-1.04)	0.10	(0.09-0.11)	0.18	(0.15-0.21)	0.41	(0.38-0.47)
Total	3,637	0.92	(0.82–1.02)	0.09	(0.09–0.11)	0.17	(0.15–0.20)	0.40	(0.36–0.47)
Children									
Race/Ethnicity									
Mexican American	526	0.35	(0.30-0.40)			0.08	(–0.09)	0.13	(0.10-0.16)
White, non-Hispanic	447	0.29	(0.24–0.33)			0.07	(–0.08)	0.09	(0.09–0.10)
Black, non-Hispanic	424	0.50	(0.44–0.57)	0.08	(–0.10)	0.10	(0.09–0.13)	0.22	(0.18–0.26)
Total	1,577	0.33	(0.30–0.37)			0.07	(–0.08)	0.10	(0.09–0.12)

TABLE 1. Geometric means (GMs) and selected percentiles of total blood mercury (Hg) concentrations (μ g/L) for women aged 16–49 years and children aged 1–5 years, by selected variables — National Health and Nutrition Examination Survey, United States, 1999–2002

*Confidence interval.

[†]Below the limits of detection.

TABLE 1. (*Continued*) Geometric means (GMs) and selected percentiles of total blood mercury (Hg) concentrations (μ g/L) for women aged 16–49 years and children aged 1–5 years, by selected variables — National Health and Nutrition Examination Survey, United States, 1999–2002

Selected percentile (95% CI)								
Variable	50th	(95% CI)	75th	(95% CI)	90th	(95% CI)	95th	(95% CI)
Women								
Race/Ethnicity								
Mexican American	0.73	(0.67-0.83)	1.27	(1.16–1.48)	2.38	(2.05-2.95)	3.60	(3.03-6.48)
White, non-Hispanic	0.81	(0.76-0.92)	1.69	(1.51–2.15)	3.73	(2.84–5.14)	6.17	(4.64–9.30)
Black, non-Hispanic	1.15	(1.05–1.41)	2.12	(1.86–2.70)	3.89	(3.24–5.03)	5.54	(4.27–11.08)
Age group (yrs)								
16–29	0.64	(0.55-0.77)	1.34	(1.24–1.54)	2.58	(2.28-3.13)	3.87	(3.32-7.80)
30–49	1.02	(0.91 - 1.19)	2.10	(1.79-2.69)	4.56	(3.74-5.76)	6.97	(5.73-11.62)
Pregnancy status								
Pregnant	0.73	(0.63-0.97)	1.50	(1.38–1.90)	3.11	(2.14-4.79)	4.86	(3.00-8.02)
Not pregnant	0.88	(0.80-1.00)	1.83	(1.65-2.11)	3.93	(3.26-4.93)	6.11	(5.12-10.90)
Total	0.86	(0.80–0.98)	1.81	(1.62–2.16)	3.89	(3.20–4.88)	6.04	(5.08–10.74)
Children								
Race/Ethnicity								
Mexican American	0.28	(0.24-0.33)	0.63	(0.56-0.81)	1.36	(1.05–1.57)	1.85	(1.60-2.66)
White, non-Hispanic	0.20	(0.17–0.25)	0.49	(0.38–0.63)	1.15	(0.80–1.49)	1.78	(1.18–2.69)
Black, non-Hispanic	0.47	(0.40–0.58)	0.88	(0.78–1.02)	1.54	(1.312.04)	2.37	(1.75–3.64)
Total	0.26	(0.23–0.29)	0.61	(0.56–0.70)	1.29	(1.08–1.69)	2.21	(1.80–3.66)

ability caused by differing exposures to Hg across survey site locations. Accordingly, the National Center for Health Statistics advises users of these data that the most reliable estimates of current exposure are obtained when the 1999–2002 data are analyzed together (*6*).

The EPA RfD is based on measures of Hg in cord blood and is a level assumed to be without appreciable harm. The RfD was determined by applying an uncertainty factor of 10 to a dose (58 μ g/L) that was the lower 95% confidence limit of a dose associated with an increased proportion of abnor-

TABLE 2. Percentage of women aged 16–49 years with blood
mercury (Hg) levels >5.8 µg/L, by race/ethnicity - National Health
and Nutrition Examination Survey, United States, 1999–2002

Race/Ethnicity	No.	% with Hg levels <u>></u> 5.8 µg/L	(95% CI*)
Mexican American	1,106	1.70	(1.04–2.79)
White, non-Hispanic	1,377	5.77	(3.71-8.97)
Black, non-Hispanic	794	4.82	(2.55-9.11)
Total	3,637	5.66	(4.04–7.95)

* Confidence interval.

mal scores on the Boston Naming Test for children exposed in utero (2). All women and children in the 1999–2002 NHANES survey period had blood Hg levels below 58 μ g/L. The harm to a fetus from levels of exposure as measured by cord blood levels between 5.8 μ g/L and 58 μ g/L is uncertain.

The findings in this report are subject to at least two limitations. First, NHANES does not include an adequate sampling of women (e.g., sport fishers) who might eat large amounts of fish to characterize the distribution of total blood Hg in this group. Second, the ratio of Hg in cord to maternal blood (i.e., equivalent to NHANES measures) is uncertain (2,8). Therefore, NHANES values might not be directly comparable to the EPA RfD, which is based on cord blood Hg levels.

Fish are an important part of a diet, high in protein and nutrients and low in saturated fatty acids and cholesterol. The short-term strategy to reduce Hg exposure is to eat fish with low Hg levels and avoid or reduce consumption of fish with high Hg levels. Because exposure to methyl-Hg can harm fetuses, the Food and Drug Administration (FDA) advises that women who are or might become pregnant not eat shark, swordfish, king mackerel, and tile fish (9). In addition, EPA and the Agency for Toxic Substances and Disease Registry have established daily consumption levels of Hg considered to be without harm (1). State-based fish advisories and bans identify fish species contaminated by Hg and their locations and provide safety advice (10). The NHANES program continues to collect Hg measurements in human tissue to monitor the effectiveness of efforts to reduce Hg exposure in the U.S. population.

References

- Agency for Toxic Substances and Disease Registry. Toxicological profile for mercury. Atlanta, GA: US Department of Health and Human Services, Agency for Toxic Substances and Disease Registry; March 1999.
- 2. National Academy of Sciences. Toxicologic effects of methylmercury. Washington, DC: National Research Council; 2000.
- Clarkson N. Current concepts: the toxicology of mercury—current exposures and clinical manifestations [Review]. N Engl J Med 2003;349:1731–7.
- CDC. Blood and hair mercury levels in young children and women of childbearing age—United States, 1999. MMWR 2000;50:140–3.

- 5. Schober S, Sinks T, Jones R, et al. Blood mercury levels in US children and women of childbearing age, 1999–2000. JAMA 2003;289: 1667–74.
- National Center for Health Statistics. NHANES analytic guidelines, June 2004. Atlanta, GA: US Department of Health and Human Services, National Center for Health Statistics; 2004. Available at http:// www.cdc.gov/nchs/data/nhanes/nhanes_general_guidelines_june_04.pdf.
- Chen HP, Paschal DC, Miller DT, Morrow J. Determination of total and inorganic mercury in whole blood by on-line digestion with flow injection. Atomic Spectroscopy 1998;19:176–9.
- Stern A, Smith A. An assessment of the cord blood:maternal blood methylmercury ratio: implications for risk assessment. Environ Health Perspect 2003;111:1465–70.
- 9. US Department of Health and Human Services, US Environmental Protection Agency. What you need to know about mercury in fish and shellfish. 2004 EPA and FDA advice for: women who might become pregnant, women who are pregnant, nursing mothers, young children. Washington, DC: US Department of Health and Human Services, US Environmental Protection Agency; 2004. Available at http:// www.cfsan.fda.gov/~dms/admehg3.html.
- US Environmental Protection Agency. State fish advisories. Washington, DC: US Environmental Protection Agency; 2004. Available at http://www.epa.gov/ost/fish/states.htm.

Outbreak of Histoplasmosis Among Industrial Plant Workers — Nebraska, 2004

In February 2004, the Nebraska Health and Human Services System (NHHSS) notified CDC about an outbreak of histoplasmosis among workers at a local agricultural processing plant (plant A). Three workers at the plant had acute, febrile, respiratory illness; two had serologic evidence of histoplasmosis. NHHSS and CDC conducted an investigation to determine the source of transmission and the extent of the outbreak. This report summarizes the findings of that investigation, which confirmed occupationally acquired histoplasmosis. Additional measures might be necessary to minimize risk for histoplasmosis among persons who work in the agricultural industry in areas where it is endemic.

Plant A is located in an area with historically low endemicity for histoplasmosis. However, in September 2003, NHHSS had investigated an outbreak of histoplasmosis at plant A related to excavation of soil and repair of an underground outdoor pipe. Approximately 3 months later, on January 2, 2004, the excavated soil (i.e., spoil pile) was moved, under standard protocol and appropriate precautions, to an off-site landfill. None of the plant workers with suspected histoplasmosis in the 2004 outbreak had participated in removal of the spoil pile, nor had they been implicated in the 2003 outbreak.

To identify workers with symptomatic acute pulmonary histoplasmosis acquired during the 2004 outbreak, a cohort study was conducted among plant workers. To better identify risk factors for disease, a nested case-control study was performed among workers who had laboratory testing for histoplasmosis. For the cohort study, all workers were instructed by plant safety managers to complete a self-administered, webbased questionnaire in late February 2004. A clinical case of histoplasmosis was defined as fever plus at least one of the following four symptoms in a plant A worker reported since January 1, 2004: headache, cough, chest pain, or shortness of breath. Workers whose symptoms were consistent with the clinical case definition had histoplasmosis serology testing performed. A laboratory-confirmed case was defined as the presence of a complement fixation (CF) titer \geq 1:32 and/or the presence of an H or M band by immunodiffusion test from a single serum sample obtained from a plant A worker, drawn at least 6 weeks after onset of illness. Controls for the case-control study were randomly selected from workers without any symptoms of histoplasmosis identified during the cohort study. These workers were asked to have a serum sample drawn for histoplasmosis testing and were found to have no serologic evidence of recent Histoplasma capsulatum infection.

Of 979 plant workers, 724 (74%) completed the cohort questionnaire; 108 (16%) had symptoms consistent with the clinical case definition. The most commonly reported symptoms were headache (93%), cough (77%), and shortness of breath (44%). No workers were hospitalized. Symptomatic workers (clinical cases) were as likely as asymptomatic workers (nonclinical cases) to report working outside, seeing bird droppings, and performing grounds work. Symptomatic workers were more likely to have worked in building complex X (the complex in closest proximity to the spoil pile) than asymptomatic workers (44 [41%] versus 141 [23%]; risk ratio = 2.0; p<0.001). Building complex X was not located in an area known to be heavily contaminated with bird droppings.

Of the 108 symptomatic workers, 90 (83%) had sera available for testing; 25 (28%) had laboratory-confirmed histoplasmosis. Analysis of 22 workers with laboratory-confirmed histoplasmosis with specified dates of symptom onset indicated a cluster of cases during mid-January (Figure). Workers with laboratory-confirmed histoplasmosis were further categorized as clustered cases (n = 18) (symptom onset during January 7–25) and outlying cases (all others, n = four). For the case-control study, the 22 workers with laboratoryconfirmed histoplasmosis were compared with 31 unmatched controls. Workers categorized as clustered cases were more likely to have worked in building complex X than controls (12 [67%] versus eight [26%]; odds ratio [OR] = 5.8; 95% confidence interval [CI] = 1.6–20.4); no specific activities, dates reporting to work, or relative amount of outside activity FIGURE. Number* of laboratory-confirmed cases of histoplasmosis among plant A workers, by date of symptom onset — Nebraska, January 7–February 25, 2004



* n = 22.

during days of any reported soil disruptions (i.e., December 30, January 2, and January 15) were associated with increased risk for acquiring histoplasmosis. In contrast, workers categorized as outlying cases were as likely to work in building complex X as controls (one [25%] versus eight [26%]; OR = 1.0; p = 1.0; no specific occupation was associated with workers with outlying case status.

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Editorial Note: The findings in this report confirm that an outbreak of occupationally acquired histoplasmosis occurred among at least 25 workers from plant A in January 2004. Histoplasmosis usually is an acute, self-limited respiratory illness with an incubation period of 1-2 weeks after inhalation of H. capsulatum spores (1). Previous occupation-related outbreaks of histoplasmosis occurred among workers in a paper factory and courthouse and among bridge workers (2-4), in which disruption of bird or bat droppings, known sources of transmission for H. capsulatum spores, had occurred. As in previous outbreaks, this investigation identified cases clustered in time and location. The majority of patients reported illness onset in mid-January, suggesting a point-source exposure in early January. In addition, workers with laboratory-confirmed infection and onset of illness in mid-January were more likely to work at building complex X than controls, supporting the hypothesis of a common source of exposure.

The likely source of this outbreak was the disruption of the spoil pile from the 2003 outbreak. The spoil pile was known to be contaminated with *H. capsulatum*; the workers closest in proximity to it worked at building complex X, and the timing until symptom onset for workers categorized as clustered cases was consistent with histoplasmosis. However, implication of the spoil pile as the source of the outbreak was surprising for several reasons. First, building complex X was approximately 950 feet from the spoil pile. Second, appropriate precautions were taken during removal of the spoil pile (e.g., limiting the number of workers in the area, using appropriate personal protective equipment [PPE], soaking the spoil pile with water before manipulation, and using both a plastic liner and a tarp to cover the soil once it was transferred to a dumpster).

Onset of illness after the mid-January period indicates that some illness might not be associated with the point-source exposure at the plant, but rather might reflect ongoing lowlevel exposure to *H. capsulatum* either through other workrelated or nonwork-related activities. Further investigation will be necessary to identify all activities placing workers at increased risk for disease.

This outbreak underscores the highly infectious nature of H. capsulatum spores and the need to protect workers when engaging in work-related activities involving exposure to H. capsulatum. In particular, these data suggest that manipulation of soil known to be contaminated with H. capsulatum can pose a risk to persons who are not engaged in the activity directly but who might be hundreds of feet away. In addition to following CDC guidelines for the prevention of histoplasmosis among workers (5), the recommendations given to prevent further outbreaks of histoplasmosis at this site included assigning job activities according to three levels of risk: higher, lower, and minimal/no risk. Activities identified as higher risk include those involved in disturbing soil obviously contaminated with bird droppings or in disturbing accumulations of bird droppings. The level of PPE required for these activities includes the use of disposable coveralls such as Tyvek[®], rubber boots over normal work shoes, and a respirator providing a higher level of protection, such as a powered air-purifying respirator with high-efficiency particulate air (HEPA) filters or a full-facepiece respirator with HEPA filter. For lower-risk activities (e.g., those disturbing soil that has not been contaminated by bird droppings or during the changing of air filters from buildings or equipment), the level of PPE might be decreased to a NIOSH-certified N-95 filtering facepiece respirator. For activities designated as minimal risk (e.g., those in which no soil or bird droppings are disturbed), no respiratory protection is required, but N-95 filtering facepiece respirators should be available on a voluntary-use basis.

References

- 1. Lehan PH, Furculow ML. Epidemic histoplasmosis. J Chronic Dis 1957;5:489–503.
- 2. Stobierski MG, Hospedales CJ, Hall WN, et al. Outbreak of histoplasmosis among employees in a paper factory—Michigan, 1993. J Clin Microbiol 1996;34:1220–3.
- Dean AG, Bates JH, Sorrels TC, et al. An outbreak of histoplasmosis at an Arkansas courthouse, with five cases of probable reinfection. Am J Epidemiol 1978;108:36–46.
- Sorley DL, Levin ML, Warren JW, Flynn JP, Gerstenblith J. Bat-associated histoplasmosis in Maryland bridge workers. Am J Med 1979; 67:623–6.
- Lenhart SW, Schafer MP, Singal M, et al. Histoplasmosis: protecting workers at risk. US Department of Health and Human Services, CDC, National Institute for Occupational Safety and Health; 1997. Publication No. 97-146.

West Nile Virus Activity — United States, October 27– November 2, 2004

During October 27–November 2, a total of 10 cases of human West Nile virus (WNV) illness were reported from eight states (Arizona, Georgia, Iowa, Michigan, New Mexico, Ohio, Oklahoma, and Pennsylvania).

During 2004, 40 states and the District of Columbia (DC) have reported 2,241 cases of human WNV illness to CDC through ArboNET (Figure and Table). Of these, 710 (32%) cases were reported in California, 381 (17%) in Arizona, and 276 (12%) in Colorado. A total of 1,295 (59%) of the 2,211 cases for which such data were available occurred in males; the median age of patients was 52 years (range: 1 month–

FIGURE. Areas reporting West Nile virus (WNV) activity — United States, 2004*



* As of 3 a.m., Mountain Standard Time, November 2, 2004.

TABLE.	Number	of human	cases	of	West	Nile	virus	(WNV)
illness,	by area -	- United S	tates, 2	200	4*			

	Neuro- invasive	West Nile	Other clinical/	Total reported	
Area	disease [†]	fever§	unspecified [¶]	to CDC**	Deaths
Alabama	13	0	0	13	0
Arizona	128	70	183	381	10
Arkansas	12	9	1	22	0
California	143	248	319	710	20
Colorado	39	237	0	276	3
Connecticut	0	1	0	1	0
District of Colum	nbia 1	0	0	1	0
Florida	29	8	0	37	2
Georgia	11	6	0	17	0
Idaho	0	0	2	2	0
Illinois	28	27	1	56	2
Indiana	5	0	2	7	1
Iowa	11	7	4	22	2
Kansas	18	25	0	43	2
Kentucky	1	6	0	7	0
Louisiana	68	17	0	85	7
Maryland	6	5	1	12	0
Michigan	10	1	0	11	0
Minnesota	13	20	0	33	2
Mississippi	23	5	2	30	3
Missouri	25	9	2	36	1
Montana	2	3	1	6	0
Nebraska	4	26	0	30	0
Nevada	25	19	0	44	0
New Jersey	1	0	0	1	0
New Mexico	29	49	4	82	4
New York	3	3	0	6	0
North Carolina	3	0	0	3	0
North Dakota	2	18	0	20	1
Ohio	11	1	0	12	2
Oklahoma	10	6	0	16	1
Oregon	0	1	0	1	0
Pennsylvania	8	3	1	12	2
South Carolina	0	1	0	1	0
South Dakota	6	45	0	51	1
Tennessee	9	1	0	10	0
Texas	83	26	0	109	8
Utah	6	5	0	11	0
Virginia	4	0	1	5	1
Wisconsin	4	6	0	10	1
Wyoming	2	5	2	9	0
Total	796	919	526	2.241	76

* As of November 2, 2004.

⁺ Cases with neurologic manifestations (i.e., West Nile meningitis, West Nile encephalitis, and West Nile myelitis).

§ Cases with no evidence of neuroinvasion.

Illnesses for which sufficient clinical information was not provided.

** Total number of human cases of WNV illness reported to ArboNet by state and local health departments.

99 years). Date of illness onset ranged from April 23 to October 21; a total of 76 cases were fatal.

A total of 199 presumptive West Nile viremic blood donors (PVDs) have been reported to ArboNET in 2004. Of these, 73 (37%) were reported in California; 38 (19%) in Arizona; 16 in Texas; 15 in New Mexico; seven in Colorado; six each in Louisiana and Oklahoma; five in Nevada; four in Georgia and Iowa; three each in Florida, Michigan, and South Dakota; two each in Minnesota, Mississippi, Missouri, and Wisconsin; and one each in Delaware, Kentucky, Nebraska, New Jersey, New York, North Dakota, Oregon, and Pennsylvania. Of the 199 PVDs, three persons aged 35, 69, and 77 years subsequently had neuroinvasive illness, and 46 persons (median age: 52 years; range: 17–73 years) subsequently had West Nile fever.

In addition, 5,441 dead corvids and 1,328 other dead birds with WNV infection have been reported from 45 states and New York City during 2004. WNV infections have been reported in horses in 37 states; one bat in Wisconsin; nine dogs in Nevada, New Mexico, and Wisconsin; six squirrels in Arizona and Wyoming; and 14 unidentified animal species in nine states (Arizona, Idaho, Illinois, Iowa, Kentucky, Missouri, Nevada, New York, and South Carolina). WNV seroconversions have been reported in 1,345 sentinel chicken flocks in 13 states (Alabama, Arizona, Arkansas, California, Delaware, Florida, Iowa, Louisiana, Nebraska, Nevada, Pennsylvania, South Dakota, and Utah) and in 25 wild hatchling birds in Missouri and Ohio. Four seropositive sentinel horses were reported in Minnesota and Puerto Rico. A total of 7,558 WNV-positive mosquito pools have been reported in 38 states, DC, and New York City.

Additional information about national WNV activity is available from CDC at http://www.cdc.gov/ncidod/dvbid/ westnile/index.htm and at http://westnilemaps.usgs.gov.

Notice to Readers

National Epilepsy Awareness Month — November 2004

November is National Epilepsy Awareness Month. Epilepsy affects approximately 2.5 million persons in the United States and is characterized by unprovoked seizures. Persons with epilepsy often face physical and cognitive side effects of both seizures and treatment, social stigma, lost productivity, and decreased quality of life. The impact on children is especially burdensome as they transition from childhood to adulthood. Outside the medical community, epilepsy is a poorly understood condition, even among families and friends of affected persons.

To improve social acceptance and understanding of this disorder and to increase support for those with it, the Epilepsy Foundation (EF), in partnership with CDC, is enhancing its Entitled to Respect campaign. During November, the campaign will expand last year's focus on educating black youth, an underserved segment of the community, by providing outreach to black women of childbearing age and by building new partnerships and improving services to affected children, adults, and families in the black community. Additional information about epilepsy and the campaign is available from EF, telephone 800-332-1000 or at http://www.epilepsy foundation.org.

Notice to Readers

Annual Health Information and Technology Week

November 7–13, 2004, is Annual Health Information and Technology Week. During this week, approximately 44,000

students and professionals will celebrate Health Information Management (HIM) professions, which include qualityassurance managers to information-security officers in settings from managed care organizations to home health agencies in both the private and public sectors. The American Health Information Management Association represents the community of professionals engaged in HIM. This year's theme, Health Information: Powered by Professionals, reflects the work performed by HIM professions in obtaining a more secure and electronic health information environment.



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CASES CURRENT DISEASE DECREASE INCREASE 4 WEEKS Hepatitis A, acute 242 Hepatitis B, acute 334 Hepatitis C, acute 42 94 Legionellosis Measles, total 1 Meningococcal disease 47 9 Mumps 898 Pertussis 0 Rubella 2 0.03125 0.0625 0.125 0.25 0.5 4 1

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals October 30, 2004, with historical data

Ratio (Log scale)[†]

Beyond historical limits

* No rubella cases were reported for the current 4-week period yielding a ratio for week 43 of zero (0). † 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.

IABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending October 30,	30, 2004 (43rd week)^
---	-----------------------

	Cum. 2004	Cum. 2003		Cum. 2004	Cum. 2003
Anthrax	-	-	HIV infection, pediatric [†]	126	169
Botulism:	-	-	Influenza-associated pediatric mortality**	-	NA
foodborne	11	11	Measles, total	23**	51 ^{§§}
infant	60	56	Mumps	162	177
other (wound & unspecified)	9	26	Plague	1	1
Brucellosis [†]	85	82	Poliomyelitis, paralytic	-	-
Chancroid	30	50	Psittacosis [†]	9	11
Cholera	4	1	Q fever [†]	60	56
Cyclosporiasis [†]	203	61	Rabies, human	3	2
Diphtheria	-	1	Rubella	10	7
Ehrlichiosis:	-	-	Rubella, congenital syndrome	-	1
human granulocytic (HGE) [†]	259	273	SARS-associated coronavirus disease [†] **	-	8
human monocytic (HME) [†]	237	221	Smallpox [†] ^{¶¶}	-	NA
human, other and unspecified	27	38	Staphylococcus aureus:	-	-
Encephalitis/Meningitis:	-	-	Vancomycin-intermediate (VISA)† 1	-	NA
California serogroup viral [†] §	74	106	Vancomycin-resistant (VRSA)† 🖤	1	NA
eastern equine ^{†§}	3	13	Streptococcal toxic-shock syndrome [†]	89	136
Powassan [†] §	-	-	Tetanus	14	16
St. Louis ^{†§}	7	40	Toxic-shock syndrome	106	103
western equine ^{†§}	-	-	Trichinosis	4	1
Hansen disease (leprosy) [†]	69	68	Tularemia [†]	74	74
Hantavirus pulmonary syndrome [†]	17	18	Yellow fever	-	-
Hemolytic uremic syndrome, postdiarrheal [†]	119	141			

-: No reported cases.

* Incidence data for reporting years 2003 and 2004 are provisional and cumulative (year-to-date).

^T Not notifiable in all states.

[§] Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Infectious Diseases (ArboNet Surveillance).

¹ 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 26, 2004.

++ Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases.

⁺⁺ Of 23 cases reported, 10 were indigenous, and 13 were imported from another country.

§§ Of 51 cases reported, 31 were indigenous, and 20 were imported from another country.

^{¶¶} Not previously notifiable.

(Hord Week)	AIE	os	Chlan	nvdia†	Coccidiodomycosis Cryptosporidiosis		oridiosis	Encephalitis/Meningitis West Nile [§]		
Poporting area	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
	31 120	35.017	722.358	710 785	4 924	3 078	2 770	2 853	794	2 831
NEW ENGLAND Maine N.H. Vt. Mass. R.I.	981 15 37 14 343 109	1,201 49 34 15 517 82	24,825 1,719 1,450 852 11,052 2,790	22,757 1,640 1,300 888 9,008 2,419	4,324 N - -	S,070	154 18 29 22 54 4	164 18 18 29 72 12		2,031 29 - 2 - 12 5
Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	463 6,925 724 3,949 1,140 1,112	504 8,345 745 4,488 1,291 1,821	6,962 86,773 18,142 26,412 12,475 29,744	7,502 88,195 16,378 28,552 13,061 30,204	N - - - - N	N - - - - N	27 441 151 85 25 180	15 361 107 104 14 136	- 12 1 2 1 8	10 220 - 56 21 143
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	2,742 525 300 1,290 493 134	3,208 641 431 1,472 511 153	123,808 30,828 14,935 32,970 31,119 13,956	130,323 35,970 14,184 39,621 26,148 14,400	15 N N - 15	7 N - 7	795 200 80 69 137 309	857 128 77 86 116 450	57 11 5 28 9 4	150 84 15 30 14 7
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr.** Kans.	641 152 50 277 14 8 41 99	632 123 67 305 3 8 42 84	44,669 8,123 5,293 17,427 1,229 2,136 4,260 6,201	41,201 8,878 4,185 15,036 1,292 2,126 3,842 5,842	5 N 3 N - 2 N	2 N 1 N - 1 N	335 115 73 56 10 33 23 25	502 129 110 41 11 36 21 154	79 13 11 25 2 6 4 18	695 48 80 39 94 151 194 89
S. ATLANTIC Del. Md. D.C. Va. W.Va. N.C. S.C.** Ga. Fla	9,492 121 1,252 621 513 67 482 535 1,327 4,574	9,954 191 1,271 811 789 71 887 664 1,502 3 768	143,838 2,436 15,744 2,732 18,554 2,314 24,286 17,050 26,394 34 328	133,512 2,483 13,399 2,602 15,874 2,154 21,168 12,002 29,277 34,553	N - - N N - - - - - - - - - - - - - - -	5 N 5 - N N - N	465 15 12 55 5 70 15 178 115	313 4 20 11 38 4 43 8 99 86	54 - 6 1 - 3 - 11 29	182 12 48 3 19 1 16 24 57
E.S. CENTRAL Ky. Tenn.** Ala. Miss.	1,528 187 617 360 364	1,630 141 700 389 400	46,039 4,728 18,220 9,382 13,709	45,897 6,727 16,905 11,964 10,301	4 N N - 4	1 N N 1	109 39 29 20 21	115 21 37 47 10	46 1 9 13 23	89 11 21 25 32
W.S. CENTRAL Ark. La. Okla. Tex.**	3,581 174 719 154 2,534	3,463 163 519 177 2,604	88,435 5,964 18,539 9,116 54,816	86,866 6,539 16,280 9,448 54,599	2 1 1 N N	- N N	66 14 3 20 29	97 17 4 13 63	172 12 68 9 83	593 23 89 56 425
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	1,178 6 15 16 257 152 437 53 242	1,272 11 21 5 327 96 535 52 225	39,926 1,946 2,277 876 9,779 4,333 13,330 3,002 4,383	40,164 1,590 2,057 809 10,736 6,160 11,056 3,083 4,673	3,174 N 2 N 20 3,067 33 52	1,939 N N 1 N 9 1,889 8 32	143 34 24 3 48 11 17 4 2	119 18 26 4 32 10 5 17 7	231 2 39 29 128 6 25	871 75 - 92 621 74 7 2
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	4,052 313 239 3,357 39 104	5,312 365 202 4,642 15 88	124,045 14,555 6,974 95,226 3,056 4,234	121,870 13,531 6,117 94,610 3,141 4,471	1,724 N - 1,724 -	1,124 N - 1,124 -	262 36 30 194 - 2	325 43 36 245 1 -	143 - 143 -	2 - - 2 -
Guam P.R. V.I. Amer. Samoa C.N.M.I.	2 595 10 U 2	5 851 29 U U	2,701 272 U 32	517 2,182 348 U U	N U	- N - U U	N U	- N - U U	- - - U	- - - U U

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending October 30, 2004, and October 25, 2003 (43rd Week)*

N: Not notifiable.

N: Not notifiable. U: Unavailable. -: No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands. * Incidence data for reporting years 2003 and 2004 are provisional and cumulative (year-to-date). † Chlamydia refers to genital infections caused by *C. trachomatis.* § Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Infectious Diseases (ArboNet Surveillance). ¶ 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 26, 2004.

** Contains data reported through National Electronic Disease Surveillance System (NEDSS).

MMWR

(4510 WEEK)											
		Escher	<i>ichia coli</i> , Ente	rohemorrhagio							
	015	7.117	Shiga tox	in positive,	Shiga toxi	n positive,	Cier	liecie	Conorrhoo		
	Cum		Cum	Cum	Cum	groupea Cum	Cum	Cum	Gond	Cum	
Reporting area	2004	2003	2004	2003	2004	2003	2004	2003	2004	2003	
UNITED STATES	2,009	2,147	219	204	142	135	14,660	15,642	256,824	271,927	
NEW ENGLAND	135	132	44	36	17	12	1,452	1,295	5,790	5,946	
Maine	10	10	-	1	-	-	112	155	184	162	
Vt.	10	15	-	-	-	-	147	106	73	71	
Mass.	56	57	13	8	17	12	620	657	2,610	2,357	
R.I. Conn	9 31	1	1 25	- 24	-	-	107 429	90 255	700 2 116	800 2 455	
	225	217	46	24	29	22	3 052	3 100	27.042	22,400	
Upstate N.Y.	106	79	33	10	13	17	1.089	848	5.917	6.371	
N.Y. City	32	7	-	-	-	-	799	1,011	8,421	11,191	
N.J. Pa	35	29 102	4	2	5	-	312 852	419	4,881	6,676 9,647	
	200	102	9	20	10	17	2 012	0.710	52,409	5,047	
Ohio	85	483	35 10	29 15	24 18	17	2,013	751	52,498 15.922	58,303 18.979	
Ind.	51	70	-	-	-	-		-	5,487	5,515	
III. Mich	49	111	1	2	1	-	338	794	14,383	17,845	
Wis.	100	130	17	12	5	-	381	526	3,582	4,642	
W.N. CENTRAL	432	390	27	46	16	20	1.670	1.675	14,181	14.398	
Minn.	106	119	15	20	1	1	626	599	2,479	2,508	
lowa	117	90	-	-		-	249	231	938	1,035	
N Dak	67 14	73 10	- 11	13	6	1	420	429	7,470	7,191	
S. Dak.	31	25	-	4	-	-	50	69	239	182	
Nebr.	60 27	43	1	5	-	-	117	124	861	1,284	
	57	30	-	-	2	10	100	191	2,107	2,123	
S. ATLANTIC Del	148	126	37 N	38 N	46 N	37 N	2,352	2,236	65,219 742	66,495 951	
Md.	20	12	4	3	3	1	100	97	6,755	6,390	
D.C.	1	1	-	-	-	-	57	41	2,061	2,052	
va. W Va	35	33 4	16	- 11	-	-	449	287	7,406	7,404	
N.C.	-	-	-	-	32	29	N	N	12,778	12,123	
S.C.	7	2	-	-	-	-	51	123	8,343	7,052	
Ga. Fla.	59	25 41	6	5 19	11	7	933	888	14,751	14,504	
E S CENTRAL	77	74	4	2	9	6	325	328	20.065	22 991	
Ky.	24	24	2	2	6	6	N	N	2,156	3,002	
Tenn.	31	32	2	-	3	-	159	150	6,887	7,013	
Miss.	7	14	-	-	-	-	- 100	- 178	5,743	7,002 5.314	
WS CENTRAL	65	78	2	4	2	4	265	252	34 267	35,950	
Ark.	14	9	1	-	-	-	103	128	2,995	3,473	
La.	3	3	-	-	-	-	37	10	8,710	9,382	
Tex.	31	25 41	- 1	4	2	4	125 N	N	3,879 18.683	3,893	
MOUNTAIN	211	269	23	24	-	6	1 272	1.322	8 660	8 621	
Mont.	16	16	-	-	-	-	68	94	58	87	
Idaho	46	68	15	15	-	-	163	167	79	61	
VVyo. Colo	8 44	61	1	-	-	-	21 437	20 383	54 2 168	36 2 384	
N. Mex.	9	10	2	4	-	-	59	43	603	987	
Ariz.	21	29	N	N	N	N	143	203	3,233	3,045	
Nev.	40 21	23	2	- 1	-	-	102	292 120	1.998	1.700	
PACIFIC	346	378	1	1	_	_	2 259	2 713	28 201	25 338	
Wash.	127	97	-	1	-	-	317	311	2,240	2,246	
Oreg.	60	.95	1	3	-	-	372	356	1,042	825	
Calif. Alaska	148	175	-	-	-	-	1,431	1,899	23,468	20,823	
Hawaii	10	4 7	-	-	-	-	66	70	1,002	994	
Guam	N	N	-	-	-	-	-	2	-	55	
P.R.	-	1	-	-	-	-	103	234	202	234	
V.I. Amor Samea	-	-	-	-	-	-	-	-	80	77	
C.N.M.I.	-	U	-	U	-	U	-	U	3	U	

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending October 30, 2004, and October 25, 2003 (43rd Week)*

MMWR

		Henatitis								
	All	ages		(viral, acute), by type						
	All se	rotypes	Serot	ype b	Non-sei	otype b	Unknowr	n serotype		A
Reporting area	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003
UNITED STATES	1.507	1.546	12	24	89	95	145	168	4.522	5.717
NEW ENGLAND	128	115	1	2	5	5	3	3	849 11	268 11
N.H. Vt	16	12 8	-	1	2	-	- 1	-	20	15
Mass.	51	54	1	1	-	5	2	1	728	151
Conn.	40	31	-	-	3	-	-	-	61	73
MID. ATLANTIC	310 100	328 119	-	2	4 4	3	33 5	41 8	530 86	1,087 109
N.Y. City	62	57	-	-	-	-	12	11	207	383
N.J. Pa.	63 85	60 92	-	-	-	-	3 13	9 13	104 133	182 413
E.N. CENTRAL	226	260	-	3	6	4	35 15	46 11	456	533
Ind.	41	41	-	-	4	-	1	5	88	54
III. Miab	50	94	-	-	-	-	11	20	158	161
Wis.	31	42	-	-	-	-	2	9	41	43
W.N. CENTRAL Minn.	88 40	96 40	2 1	2	3 3	7 7	10 1	12 2	148 32	147 37
lowa	1	-	1	-	-	-	-	-	43	25
N. Dak.	20	2	-	-	-	-	-	-	1	47
S. Dak. Nebr.	- 8	1	-	-	-	-	- 1	-	3 10	- 12
Kans.	7	15	-	-	-	-	2	1	22	25
S. ATLANTIC Del.	380	339	1	2	21	14	29	19	903 5	1,461 8
Md.	51	79	-	1	4	6	-	1	94	151
Va.	35	45	-	-	-	-	-	5	115	33 85
W.Va. N C	15 52	14 36	-	-	1	-	3 1	- 2	6 94	13 81
S.C.	4	6	-	-	-	-	-	2	24	35
Ga. Fla.	124 99	62 96	-	- 1	10	5	22	6 3	316 242	696 359
E.S. CENTRAL	59	71	1	1	-	3	8	8	140	242
ry. Tenn.	38	42	-	-	-	2 1	6	5	29 80	176
Ala. Miss	13	21 2	1	1	-	-	2	3	8 23	23 15
W.S. CENTRAL	62	70	1	2	7	10	2	4	308	548
Ark.	3	6	-	-	-	1	1	-	56	27
Okla.	47	20 41	-	-	7	7	-	-	19	17
Tex.	1	3	1	2	-	-	-	-	189	465
Mont.	167	- 139	4 -	6	- 25	- 22	- 18	- 16	388	402
Idaho	5	4	-	-	-	-	2	1	19	13
Colo.	41	32	-	-	-	-	5	6	46	59
N. Mex.	34	16	1	-	7	4	5	1	19	19
Ariz. Utah	61 13	64 12	- 2	6	12	9	2	4	235	223
Nev.	12	10	1	-	3	4	1	-	12	45
PACIFIC	87	128	2	4	18	27	7	19	800	1,029
Oreg.	39	33	-	-	-	-	3	2	59	49
Calif.	33	55	-	4	18	20	1	9	662	906
Alaska Hawaii	4 8	18 11	-	-	-	-	1 1	5	5 21	8 12
Guam	-	-	-	-	-	-	-	-	-	2
V.I.	-	-	-	-	-	-	-	-	∠ I -	-
Amer. Samoa C.N.M.I.	U	U U	U -	U U	U	U U	U	U U	U	U U

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending October 30, 2004, and October 25, 2003 (43rd Week)*

 C.N.M.I.
 U
 U

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

 * Incidence data for reporting years 2003 and 2004 are provisional and cumulative (year-to-date).

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Reporting areaCuiUNITED STATES5,22NEW ENGLAND30	B n. Cum. 4 5,816 1 304 2 1 3 16 5 4	Cum. 2004 714 10	C C 2003 874	Legio Cum. 2004	nellosis Cum.	Lister Cum.	iosis Cum.	Lyme d Cum.	isease Cum
Reporting areaCur 200UNITED STATES5,22NEW ENGLAND300	n. Cum. 2003 4 5,816 1 304 2 1 3 16 5 4	Cum. 2004 714 10	Cum. 2003 874	Cum. 2004	Cum.	Cum.	Cum.	Cum.	Cum
UNITED STATES 5,22 NEW ENGLAND 30	4 5,816 1 304 2 1 3 16 5 4	714	874	2004	1 2003	2004	2003	2004	2003
NEW ENGLAND 30	1 304 2 1 3 16 5 4	10		1,506	1,766	520	568	14,726	17,441
Maine N.H. 3 Vt. Mass. 17 R.I.	1 191 5 12	5 4 -	7 - 7 -	52 10 5 8 14	99 2 8 5 50 13	33 7 3 2 5 1	43 6 3 1 17	2,260 53 179 45 774 183	3,309 137 147 40 1,441 466 4020
Conn. a MID. ATLANTIC 99 Upstate N.Y. 7 N.Y. City 9 N.J. 58 Pa. 24	5 80 9 629 9 76 1 163 2 153 7 237	126 14 - 112	- 104 13 - - 91	426 88 42 76 220	523 129 61 76 257	125 41 17 20 47	116 29 21 22 44	9,802 3,338 - 2,635 3,829	1,078 11,635 3,858 187 2,665 4,925
E.N. CENTRAL 46 Ohio 10 Ind. 3 III. 7 Mich. 22 Wis. 2	3 433 4 116 8 33 1 52 7 191 3 41	99 6 7 12 74	127 7 8 18 89 5	405 194 66 20 118 7	362 186 25 41 93 17	84 37 16 5 23 3	74 22 6 19 18 9	790 59 16 1 28 686	862 62 20 67 6 707
W.N. CENTRAL 26 Minn. 4 Iowa 1 Mo. 15 N. Dak. 5 S. Dak. 3 Nebr. 3 Kans. 1	0 269 4 31 3 10 4 185 4 2 - 2 1 23 4 16	42 17 25 - -	191 8 1 180 - 2 -	43 7 5 21 2 4 1 3	61 3 9 31 1 2 5 10	14 5 1 5 - 1 2	15 4 - 6 - - 4 1	493 393 42 47 - - 7 4	333 223 48 55 - 1 2 4
S. ATLANTIC 1,63 Del. 2 Md. 13 D.C. 1 Va. 22 W.Va. 3 N.C. 13 S.C. 6 Ga. 57 Fla. 41	$\begin{array}{cccc} 0 & 1,677 \\ 8 & 8 \\ 5 & 105 \\ 9 & 10 \\ 4 & 150 \\ 4 & 27 \\ 8 & 132 \\ 5 & 143 \\ 2 & 568 \\ 5 & 534 \end{array}$	143 - 14 3 16 21 11 6 17 55	131 7 7 3 11 24 13 66	323 12 67 8 42 8 29 3 39 115	450 24 114 82 16 36 7 32 125	96 N 14 - 3 21 3 16 23	113 N 23 1 9 6 16 4 28 26	1,192 137 690 8 149 22 105 12 13 56	1,055 185 619 8 81 20 91 8 10 33
E.S. CENTRAL 38 Ky. 5 Tenn. 17 Ala. 6 Miss. 8	0 385 9 57 3 169 2 80 6 79	86 23 35 4 24	68 13 15 5 35	82 35 33 11 3	93 37 32 19 5	21 4 10 5 2	27 7 8 10 2	44 15 17 3 9	56 13 15 8 20
W.S. CENTRAL 25 Ark. 6 La. 5 Okla. 4 Tex. 8	1 928 5 71 3 108 7 50 6 699	105 2 60 3 40	143 3 94 2 44	56 - 4 5 47	63 2 1 7 53	26 2 3 - 21	46 1 3 3 39	34 8 4 - 22	88 - 6 - 82
MOUNTAIN 39 Mont. 1 Idaho 1 Wyo. Colo. Colo. 4 N.Mex. 1 Ariz. 20 Utah 4 Nev. 6	3 483 2 14 0 7 7 28 8 68 2 32 8 220 1 41 5 73	41 2 8 7 5 4	44 2 1 - 10 - 7 - 24	69 2 7 5 17 4 11 19 4	56 4 3 2 9 2 10 20 6	24 1 - 12 - 3 8	31 2 9 2 10 2	30 6 3 1 6 11	14 - 3 2 - 1 3 2 3
PACIFIC 54 Wash. 4 Oreg. 9 Calif. 38 Alaska 1 Hawaii 1	7 708 5 63 8 94 0 526 4 4 0 21	62 19 14 25 -	59 17 12 28 -	50 10 N 40	59 8 N 51 -	97 9 5 79 - 4	103 7 4 87 -	81 13 30 36 2 N	89 3 14 69 3 N
Guam P.R. 4 V.I. Amer. Samoa	- 9 6 103 	- - - U	5 - - U	- 1 - U	- - - U	- - - U	- - - U	N U	N

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending October 30, 2004, and October 25, 2003

(4510 Week)	Ма	laria	Mening	jococcal	Perti	ussis	Rabies	animal	Rocky Mountain spotted fever		
Poporting area	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	
UNITED STATES	1,039	1,102	1,059	1,371	11,871	7,329	4,727	5,893	1,228	746	
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	66 6 5 4 33 4 14	57 2 6 2 27 2 18	57 9 5 2 32 2 7	65 6 3 40 2 11	1,336 2 69 62 1,160 31 12	1,116 12 82 60 892 16 54	563 39 25 33 244 32 190	516 61 23 30 181 61 160	17 - - 14 1 2	7 - - 7 -	
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	245 40 112 52 41	300 46 163 57 34	129 29 23 31 46	165 40 38 22 65	2,351 1,628 128 190 405	863 389 120 134 220	493 453 11 - 29	782 362 6 62 352	77 3 19 27 28	39 - 13 16 10	
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	91 27 14 22 18 10	92 17 2 40 23 10	149 60 23 12 43 11	217 52 39 62 39 25	2,554 485 175 319 247 1,328	776 229 55 68 99 325	143 69 10 46 16 2	156 50 25 23 44 14	27 15 5 2 5	19 8 1 5 5	
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. S. Dak. Nebr. Kans.	60 25 4 17 3 1 3 7	42 20 5 1 2 - 9	78 22 15 18 2 2 4 15	110 25 23 43 1 1 6 11	1,553 302 132 251 693 30 40 105	377 141 117 69 6 3 8 33	435 81 95 51 53 10 53 92	579 31 95 39 50 120 92 152	105 1 88 4 12	59 1 2 47 - 5 3 1	
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	284 6 64 11 43 1 18 9 54 78	273 2 63 13 33 4 20 4 60 74	196 4 10 4 17 5 27 11 21 97	235 8 24 5 23 5 30 21 27 92	561 8 104 3 170 18 72 42 32 112	536 7 74 2 87 16 109 113 29 99	1,664 9 253 410 56 519 125 290 2	2,288 43 305 449 77 692 205 329 188	627 4 62 30 4 427 17 64 19	442 1 95 1 28 5 207 32 64 9	
E.S. CENTRAL Ky. Tenn. Ala. Miss.	27 4 7 11 5	27 8 5 7 7	53 9 15 14 15	73 16 19 20 18	234 57 135 28 14	138 44 63 18 13	123 20 36 56 11	186 33 97 55 1	169 2 89 44 34	115 1 62 21 31	
W.S. CENTRAL Ark. La. Okla. Tex.	90 7 4 7 72	113 4 4 4 101	97 15 33 9 40	152 13 37 14 88	653 63 10 33 547	640 43 10 72 515	953 45 - 95 813	1,015 25 2 175 813	176 98 5 71 2	56 - 42 14	
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	40 1 13 3 11 7 5	37 - 1 21 2 7 4 1	57 3 7 13 7 12 5 7	70 4 6 2 20 8 21 1 8	1,246 46 35 28 623 126 194 156 38	807 5 69 124 282 62 118 113 34	196 25 7 6 42 4 101 8 3	165 20 15 6 38 5 62 14 5	25 3 4 2 2 2 8	8 1 2 2 - - 1 -	
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	136 16 16 100 1 3	161 23 9 122 1 6	243 29 52 153 3 6	284 29 50 186 7 12	1,383 613 340 399 10 21	2,076 633 402 981 50 10	157 6 143 8	206 - 6 191 9 -	5 - 3 2 -	1 - 1 -	
Guam P.R. V.I. Amer. Samoa C.N.M.I.	- - - U	1 2 - U U	5 - U	9 - U U	6 - U	1 2 - U U	52 - - -	- 65 - U U	N U	- N - U U	

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending October 30, 2004, and October 25, 2003

MMWR

(43rd Week)^							Strei	ptococcus pne	<i>umoniae</i> . inv	asive	
	Salmor	ellosis	Shige	llosis	Streptococc	al disease, group A	Drug res all a	sistant, ges	Age <	Age <5 years	
Reporting area	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	
UNITED STATES	32,961	35,911	9,802	19,348	3,814	4,761	1,797	1,658	567	572	
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	1,743 79 123 54 989 107 391	1,802 114 127 63 1,049 108 341	248 4 8 2 156 18 60	284 6 7 7 191 13 60	156 8 17 8 106 17	408 24 28 19 181 11 145	26 2 7 N 17	85 - 6 N 10 69	60 3 N 3 47 7 U	8 - 4 N 4 U	
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	4,550 1,027 1,016 734 1,773	4,154 970 1,147 696 1,341	955 376 308 185 86	1,990 393 343 319 935	610 206 83 141 180	827 309 125 157 236	111 47 U - 64	107 56 U 51	94 64 U 6 24	83 62 U 2 19	
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	4,094 1,103 504 1,073 748 666	4,819 1,165 459 1,711 669 815	871 148 186 251 150 136	1,584 262 128 861 220 113	747 199 86 159 260 43	1,115 263 107 283 318 144	402 280 122 - N N	365 236 129 - N N	136 67 33 - N 36	252 79 25 101 N 47	
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	2,030 522 385 519 37 112 130 325	2,105 465 324 785 30 101 144 256	356 61 131 3 10 22 68	668 89 63 319 6 16 84 91	265 130 N 54 11 16 14 40	294 141 N 66 15 21 24 27	16 N 11 5 N	15 - N 11 3 1 - N	85 55 N 12 3 - 6 9	62 43 N 3 5 - 5 6	
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fia.	9,353 81 682 52 1,047 189 1,377 765 1,710 3,450	8,958 91 717 35 896 114 1,125 642 1,711 3,627	2,306 6 121 32 141 6 293 275 596 836	5,800 161 527 66 383 - 837 413 1,050 2,363	849 3 138 9 65 22 115 37 262 198	784 6 193 8 91 31 93 38 156 168	946 4 5 N 94 N 69 276 498	890 1 18 - N 61 N 124 201 485	46 N 33 3 N 10 U N N N	18 N 7 N 11 U N N N N	
E.S. CENTRAL Ky. Tenn. Ala. Miss.	2,180 294 520 624 742	2,511 344 646 631 890	666 60 323 237 46	819 114 268 275 162	186 54 132 -	169 41 128 -	119 26 92 - 1	120 16 104 -	5 N N 5	N N N	
W.S. CENTRAL Ark. La. Okla. Tex.	2,811 480 601 352 1,378	5,312 701 771 409 3,431	2,308 67 231 399 1,611	4,978 97 414 722 3,745	221 16 2 56 147	239 6 1 74 158	51 8 43 N N	65 20 45 N N	103 8 24 36 35	88 7 17 44 20	
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	2,036 176 135 48 482 229 614 209 143	1,859 93 154 71 424 228 547 186 156	698 4 13 5 140 108 338 43 43	1,034 2 28 6 274 212 410 42 60	442 - 8 131 70 184 38 3	393 1 18 2 112 96 132 30 2	33 N 10 5 N 16 2	7 N 6 - N 1 -	38 N 35 N 3	61 N 45 11 N 5	
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	4,164 488 361 2,945 53 317	4,391 493 367 3,291 61 179	1,394 96 59 1,191 5 43	2,191 147 201 1,796 8 39	338 53 N 183 - 102	532 56 N 366 - 110	93 - N - 93	4 - N - 4		N N N N	
Guam P.R. V.I. Amer. Samoa C.N.M.I.	225 - U 3	40 548 - U U	8 - U -	33 25 - U U	N U	- N - U U	N U U	N U U	N U	N U U	

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending October 30, 2004, and October 25, 2003

		Svp	hilis						Varicella (Chickenpox)		
	Primary	& secondary	Cong	enital	Tube	rculosis	Typho	oid fever			
Reporting area	Cum.	Cum. 2003	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	Cum. 2004	Cum. 2003	
UNITED STATES	6,081	5,778	277	362	8,583	10,375	239	308	14,685	13,494	
NEW ENGLAND Maine N.H.	158 2 4	174 7 16	5 - 3	1 - -	291 13	354 19 11	19 - -	26 - 2	607 180	2,686 755	
Vt. Mass. R.I. Conn.	- 100 21 31	1 111 19 20	- - 1 1	- - - 1	- 187 29 62	9 185 43 87	- 13 1 5	- 15 2 7	427 - - -	595 143 5 1,188	
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	778 83 464 126 105	712 33 407 142 130	39 4 12 22 1	56 9 30 17	1,689 216 852 343 278	1,826 233 939 360 294	54 9 18 13 14	72 12 34 21 5	76 - - 76	33	
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	672 181 46 266 152 27	758 172 38 316 217 15	50 1 8 12 29	63 3 12 19 28 1	954 163 109 418 193 71	945 167 110 451 165 52	17 5 - 10 2	32 2 4 16 10	4,648 1,119 - 3,137 392	4,543 1,024 - 2,784 735	
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. S. Dak. Nebr. Kans	128 15 5 81 - 5 22	128 40 8 49 2 2 5 5 22	5 1 - 2 - - - 2	4 - 4 - -	354 147 29 85 3 8 27 55	380 156 26 97 - 16 16 69	9 5 - 2 - 2	6 2 1 - - 1	130 N 5 82 43	48 N 48	
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Ela	1,581 8 290 67 89 2 161 99 270 595	1,520 6 259 43 69 2 128 84 403 526	43 1 7 1 3 - 10 7 1	74 12 1 16 12 13 20	1,565 195 66 223 15 233 151 11 671	2,080 23 201 219 19 261 142 427 788	40 - 11 - 7 - 6 - 6 10	44 - 9 - 14 - 7 - 5 9	1,904 4 22 487 1,137 N 254	1,815 26 1 26 478 1,064 N 220	
E.S. CENTRAL Ky. Tenn. Ala. Miss.	336 40 109 142 45	264 31 112 99 22	18 1 8 7 2	11 1 2 6 2	444 94 164 153 33	567 96 193 185 93	7 3 4	5 - 2 3 -	- - -	- - -	
W.S. CENTRAL Ark. La. Okla. Tex.	1,010 35 237 24 714	769 42 137 56 534	43 - 2 41	63 2 1 1 59	842 94 - 135 613	1,520 74 - 119 1,327	19 - 1 18	30 - 1 29	5,247 - 46 - 5,201	3,880 - 16 - 3,864	
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	299 - 18 3 36 46 157 7 32	267 10 29 54 158 6 10	45 2 - 1 42 -	30 2 3 7 18	391 4 4 85 18 175 33 68	366 5 8 4 83 39 175 30 22	6 - - 1 - 2 1 2	6 - - 3 - 2 -	2,073 35 1,599 83 356	489 - 45 - 3 - 441	
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	1,119 110 24 978 1 6	1,186 64 39 1,076 1 6	29 - - 28 - 1	60 - 58 - 2	2,053 191 71 1,665 32 94	2,337 198 90 1,910 47 92	68 6 2 54 - 6	87 3 4 79 - 1	- - - - -	- - - -	
Guam P.R. V.I. Amer. Samoa C.N.M.I.	112 4 U 2	1 169 1 U U	- 5 - U -	- 13 - U U	- 62 - U 10	48 95 - U U	- - - U -	- - U U	230 U	121 506 - U U	

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending October 30, 2004, and October 25, 2003 (43rd Week)*

TABLE III. Deaths in 122 U.S. cities,* week ending October 30, 2004 (43rd Week)

		All o	causes, b	y age (ye	ears)				All causes, by age (years)						
Reporting Area	All Ages	<u>></u> 65	45-64	25–44	1–24	<1	P&I [†] Total	Reporting Area	All Ages	<u>></u> 65	45-64	25–44	1–24	<1	P&l⁺ Total
NEW ENGLAND	527	373	95	38	14	7	52	S. ATLANTIC	1,166	690	291	118	43	24	54
Boston, Mass.	155	101	27	14	8	5	17	Atlanta, Ga.	133	65	41	13	11	3	2
Bridgeport, Conn.	23	16	/	-	-	-	2	Baltimore, Md.	155	73	50	20	9	3	13
Cambridge, Mass.	23	19	2	2	-	-	0	Charlotte, N.C.	102	67	19	12	3	4	1
Hartford Conn	60	10	12	2 4	-	-	2 1	Miami Fla	140	92	37	12	11	4	11
Lowell Mass	25	18	6	1	-	-	1	Norfolk Va	49	35	10	2	2	-	5
Lvnn. Mass.	15	10	5	-	-	-	-	Richmond, Va.	58	33	19	5	1	-	3
New Bedford, Mass.	20	18	1	1	-	-	2	Savannah, Ga.	60	40	13	6	1	-	4
New Haven, Conn.	U	U	U	U	U	U	U	St. Petersburg, Fla.	72	51	11	6	3	1	2
Providence, R.I.	52	35	8	7	2	-	6	Tampa, Fla.	175	103	47	20	4	1	9
Somerville, Mass.	5	3	1	-	1	-	-	Washington, D.C.	200	121	40	25	6	8	4
Springfield, Mass.	38	23	(4	2	2	3	Wilmington, Del.	14	10	4	-	-	-	-
Worcester Mass	24 65	19 50	3 1/	2	-	-	3	E.S. CENTRAL	890	581	216	40	26	27	55
	05	50	14	1	-	-	9	Birmingham, Ala.	182	130	34	8	5	5	18
MID. ATLANTIC	2,060	1,428	421	130	36	44	100	Chattanooga, Tenn.	82	60	12	2	2	6	8
Albany, N.Y.	52	35	12	5	-	-	2	Knoxville, Tenn.	100	70	20	4	4	2	1
Allentown, Pa.	29	24	4	-	-	1	1	Lexington, Ky.	63 205	36	21	3	2	1	4
Camden N I	26	40	9	2 1	-	6	4	Mobile Ala	205	135	24	5	2	1	14
Flizabeth N.I	14	9	2	2	1	-	-	Montgomery Ala	57	32	15	5	2	3	4
Erie. Pa.	39	30	7	2	-	-	-	Nashville, Tenn.	124	70	39	7	4	4	6
Jersey City, N.J.	37	27	4	5	1	-	-		4 077	000	200	407	10	40	50
New York City, N.Y.	1,087	760	232	60	16	19	43	W.S. CENTRAL	1,377	899	306	107	49	16	56
Newark, N.J.	52	30	14	7	-	1	1	Baton Rouge La	75	50	10	0		-	4
Paterson, N.J.	11	7	3	1	-	-	-	Corpus Christi Tex	46	29	9	1	5	2	5
Philadelphia, Pa.	303	176	80	25	10	11	20	Dallas, Tex.	194	115	51	16	10	2	11
Pittsburgh, Pa. ^s	25	16	4	4	-	1	-	El Paso, Tex.	90	65	16	6	2	1	-
Reading, Pa.	20	18	16	1	-	-	2	Ft. Worth, Tex.	127	85	27	9	3	3	4
Schenectady NY	24	20	3	1	-	-	2	Houston, Tex.	310	189	77	30	12	2	13
Scranton Pa	18	15	2	1	-	-	1	Little Rock, Ark.	87	60	13	9	4	1	-
Syracuse, N.Y.	74	55	12	3	4	-	12	New Orleans, La.	47	24	16	7	-	-	-
Trenton, N.J.	22	18	2	2	-	-	-	San Antonio, Tex.	206	137	41	18	8	2	14
Utica, N.Y.	20	13	6	1	-	-	-	Tulsa Okla	125	5Z 03	13	1	2	∠ 1	1
Yonkers, N.Y.	16	14	1	1	-	-	1		120	33	20		2		-
E.N. CENTRAL	2,071	1,414	466	110	41	39	149	MOUNTAIN	754	496	159	57	27	15	40
Akron, Ohio	61	45	13	3	-	-	16	Albuquerque, N.M.	132	92	25	10	3	2	11
Canton, Ohio	33	22	9	1	-	1	4	Colo Springs Colo	30 65	20	16	2 5	1	∠ 1	3
Chicago, III.	362	237	81	24	8	11	21	Denver Colo	103	60	24	7	- 8	4	3
Cincinnati, Ohio	83	56	20	3	2	2	5	Las Vegas, Nev.	235	150	58	19	4	4	5
Cleveland, Ohio	237	170	53	9	4	1	14	Ogden, Utah	31	25	4	2	-	-	3
Davton Ohio	197	139	42	9	3	0	17	Phoenix, Ariz.	U	U	U	U	U	U	U
Dayton, Onio	160	96	20 54	12	3	1	9 10	Pueblo, Colo.	37	28	8	1	-	-	2
Evansville Ind	47	35	6	4	2	-	-	Salt Lake City, Utah	113	74	19	11	7	2	10
Fort Wayne, Ind.	40	28	10	-	2	-	5	Tucson, Ariz.	U	U	U	U	U	U	U
Gary, Ind.	19	6	7	5	1	-	-	PACIFIC	1,766	1,212	376	101	40	36	151
Grand Rapids, Mich.	56	45	8	2	1	-	5	Berkeley, Calif.	15	9	4	2	-	-	3
Indianapolis, Ind.	197	131	41	12	6	7	10	Fresno, Calif.	170	121	34	7	4	4	15
Lansing, Mich.	47	33	12	2	-	-	2	Glendale, Calif.	6	6	-	-	-	-	1
Milwaukee, Wis.	114	76	27	5	2	4	9	Honolulu, Hawaii	80	63	13	1	1	2	3
Peona, III.	50 57	30		3	-	-	2 0	Long Beach, Calli.	00 270	270	70	25 25	2	0	0 27
South Bend Ind	47	44 35	0 6	3	2	-	0 4	Pasadena Calif	15	12	1	25	1	• -	37
Toledo Ohio	88	63	21	4	-	-	3	Portland Oreg	126	84	26	8	2	5	5
Youngstown. Ohio	65	48	11	3	2	1	2	Sacramento, Calif.	135	96	27	5	5	2	11
	000	405	440	00	40		50	San Diego, Calif.	146	102	25	11	6	2	19
W.N. CENTRAL	636	405	110	29	18	14	59	San Francisco, Calif.	104	70	25	7	1	1	17
Duluth Minn	40	42	3	-	2		7	San Jose, Calif.	201	126	56	12	5	2	14
Kansas City Kans	20	10	6	3	2 1	-	2	Santa Cruz, Calif.	27	18	6	2	-	1	4
Kansas City, Mo.	89	56	21	3	4	5	5	Seattle, Wash.	143	93	35	10	3	2	5
Lincoln, Nebr.	35	28	7	-	-	-	5	Spokane, Wash.	57	32	17	3	1	4	9
Minneapolis, Minn.	77	55	13	5	2	2	6	lacoma, Wash.	97	69	21	2	3	2	1
Omaha, Nebr.	74	65	6	-	2	1	4	TOTAL	11,247¶	7,558	2,440	730	294	222	716
St. Louis, Mo.	103	63	24	8	3	5	10								
St. Paul, Minn.	49	41	7	-	1	-	4								
Wichita, Kans.	84	62	13	9	-	-	10								

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

¹ Total includes unknown ages.

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