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Racial/Ethnic Disparities in Self-Rated Health Status Among Adults With and Without Disabilities – United States, 2004–2006

Self-rated health status has been found to be an independent predictor of morbidity and mortality (1), and racial/ethnic disparities in self-rated health status persist among the U.S. adult population (2). Black and Hispanic adults are more likely to report their general health status as fair or poor compared with white adults (2). In addition, the prevalence of disability has been shown to be higher among blacks and American Indians/Alaska Natives (AI/ANs) (3). To estimate differences in self-rated health status by race/ethnicity and disability, CDC analyzed data from the 2004–2006 Behavioral Risk Factor Surveillance System (BRFSS) surveys. This report summarizes the results of that analysis, which indicated that the prevalence of disability among U.S. adults ranged from 11.6% among Asians to 29.9% among AI/ANs. Within each racial/ethnic population, adults with a disability were more likely to report fair or poor health than adults without a disability, with differences ranging from 16.8 percentage points among Asians to 37.9 percentage points among AI/ANs. Efforts to reduce racial/ethnic health disparities should explicitly include strategies to improve the health and well being of persons with disabilities within each racial/ethnic population.

BRFSS is a state-based, random-digit-dialed telephone survey of the noninstitutionalized, U.S. civilian population aged ≥ 18 years. In 2004, 2005, and 2006, approximately 1 million persons from all 50 states, the District of Columbia, Puerto Rico, and the U.S. Virgin Islands participated in the BRFSS survey.* Consistent with the definition of disability from *Healthy People 2010* (4), respondents were asked, “Are you limited in any way in any activities because of physical, mental, or emotional problems?” and “Do you now have any health problem that requires you to use special equipment, such as a cane, a wheelchair, a special bed, or a special telephone?”

*Hawaii did not collect data in 2004.

Participants who responded “yes” to either question were classified as having a disability. To assess self-rated health status, participants were asked, “Would you say that in general your health is excellent, very good, good, fair, or poor?”

The following racial/ethnic categories were included in this analysis: white, black, Hispanic, Asian, Native Hawaiian or Other Pacific Islander, and AI/AN.[†] Data from 2004, 2005, and 2006 were aggregated to provide sufficient power to analyze low-count racial/ethnic populations. Prevalence estimates were weighted and age adjusted to the 2000 U.S. standard population. Weighted population estimates were determined by taking the final weights for each year during 2004–2006 and dividing by three. Data were weighted to compensate for unequal probabilities of selection, to adjust for nonresponse and telephone noncoverage, to ensure that results were consistent with population data, and to make population estimates.[§]

[†] For this report, persons identified as white, black, Asian, Native Hawaiian or Other Pacific Islander, and AI/AN are all non-Hispanic. Persons identified as Hispanic might be of any race.

[§] Additional information available at <http://health.utah.gov/opha/ibishelp/brfss/issues.htm>.

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Prevalence estimates and standard errors were obtained using statistical software to account for the complex sampling design. Chi-square tests were used to compare self-rated health status between racial/ethnic populations and by disability status. Council of American Survey Research Organizations (CASRO) median response rates[†] for the 2004–2006 BRFSS surveys were 52.7% (2004), 51.1% (2005), and 51.4% (2006). The median cooperation rates^{**} for each year were 74.3% (2004), 75.1% (2005), and 74.5% (2006).

During 2004–2006, an estimated 19.9% of the total U.S. population aged ≥ 18 years (i.e., an average of 43 million persons) had a disability. The prevalence of disability was highest among AI/ANs (29.9%) and lowest among Asians (11.6%) (Table 1). Nearly 84% of the total U.S. adult population reported having good or better health, but substantial variation was observed in self-rated health status across racial/ethnic populations. Nearly 60% of white, Asian, and Native Hawaiian or Other Pacific Islander respondents (59.3%, 55.8%, and 55.4%, respectively) rated their health as very good or excellent, whereas 44.4% of black respondents reported their health to be very good or excellent. White and Asian adults had similar rates of self-rated fair or poor health (12.9% and 10.4%, respectively), whereas fair or poor health was reported more frequently among other minority populations: 21.1% among blacks, 14.8% among Native Hawaiian or Other Pacific Islanders, and 24.5% among AI/ANs. Hispanic adults rated their health status approximately equally across the three health status categories: very good or excellent (33.6%), good (35.4%), and fair or poor (31.1%).

Overall, adults with a disability were less likely to report excellent or very good health (27.2% versus 60.2%; $p < 0.01$) and more likely to report fair or poor health (40.3% versus 9.9%; $p < 0.01$), compared with adults without disability (Table 2). White adults without a disability had the highest proportion of respondents who rated their health as very good or excellent (66.9%), whereas 49.9% of black respondents without a disability reported very good or excellent health. Reports of fair or poor health among adults with a disability were most common among Hispanics and AI/ANs (55.2% and 50.5%, respectively) and least common among Asians (24.9%).

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[†] The percentage of persons who completed interviews among all eligible persons, including those who were not successfully contacted.

^{**} The percentage of persons who completed interviews among all eligible persons who were contacted.

TABLE 1. Disability and self-rated health status among U.S. adults aged ≥ 18 years, by race/ethnicity — Behavioral Risk Factor Surveillance System, United States,* 2004–2006

Race/Ethnicity†	Disability§				Excellent or very good health				Good health				Fair or poor health			
	Sample population	Weighted U.S. population¶	%**	SE††	Sample population	Weighted U.S. population	%	SE	Sample population	Weighted U.S. population	%	SE	Sample population	Weighted U.S. population	%	SE
White	195,804	32,437,544	20.3	0.1	429,877	89,109,657	59.3	0.1	225,743	42,965,935	27.8	0.1	130,116	21,053,344	12.9	0.1
Black	18,713	4,181,086	21.2	0.3	32,734	9,538,829	44.4	0.3	28,709	7,218,402	34.6	0.3	19,739	4,200,595	21.1	0.3
Hispanic	13,596	4,456,898	16.9	0.3	25,957	11,778,660	33.6	0.4	26,357	12,064,608	35.4	0.4	22,033	9,009,330	31.1	0.4
Asian	1,472	508,360	11.6	0.7	7,623	3,261,549	55.8	0.9	5,127	1,791,107	33.8	0.9	1,566	470,499	10.4	0.6
Native Hawaiian or Other Pacific Islander	351	106,044	16.6	2.1	1,043	439,397	55.4	2.7	692	231,004	29.7	2.3	300	81,042	14.8	2.2
American Indian/Alaska Native	4,385	671,346	29.9	1.0	5,652	990,624	42.7	1.0	5,131	744,749	32.8	0.9	3,981	550,738	24.5	0.8
Total§§	241,863	43,786,716	19.9	0.1	512,996	117,631,008	53.4	0.1	298,772	66,518,557	30.2	0.1	183,253	36,412,487	16.4	0.1

* Includes the District of Columbia, Puerto Rico, and the U.S. Virgin Islands. Hawaii did not collect data in 2004.

† Persons identified as white, black, Asian, Native Hawaiian or Other Pacific Islander, and American Indian/Alaska Native are all non-Hispanic. Persons identified as Hispanic might be of any race.

§ Based on a “yes” response to either of the following questions: “Are you limited in any way in any activities because of physical, mental, or emotional problems?” and “Do you now have any health problem that requires you to use special equipment, such as a cane, a wheelchair, a special bed, or a special telephone?”

¶ Weighted population estimates were determined by taking the final weights for each year during 2004–2006 and dividing by three. Data were weighted to compensate for unequal probabilities of selection, to adjust for nonresponse and telephone noncoverage, to ensure that results were consistent with population data, and to make population estimates. Additional information available at <http://health.utah.gov/oph/ibishelp/brfss/issues.htm>.

** Age adjusted to the 2000 U.S. standard population.

†† Standard error.

§§ Sample population and weighted estimates by race/ethnicity do not sum to column total because respondents who reported being multiracial or of other race were included in the total.

Editorial Note: *The Surgeon General's Call to Action to Improve the Health and Wellness of People with Disabilities* notes that good health is essential if persons with disabilities are to work, learn, and fully interact with their families and community (5). The concept of health should be the same for persons with and without disabilities (5). As in previous studies (2), the findings in this report indicated that, in 2004–2006, self ratings of fair or poor health were generally higher among black, Hispanic, Native Hawaiian or Other Pacific Islander, and AI/AN adults than among their white and Asian counterparts. Also, as in previous studies (3), the findings in this report show that a higher proportion of persons with disabilities rated their health as fair or poor compared with persons without disabilities. This analysis also determined that the difference in self-rated fair or poor health between persons with and without disabilities varied by race/ethnicity. The absolute difference between persons with and without disabilities ranged from 16.8 percentage points for Asians to 37.9 percentage points for AI/ANs. These differences are attributed, in part, to health-care and wellness promotion services being inaccessible or unavailable for certain persons with disabilities (5). Health-care delivery has been slow to reduce disparities that would enable many persons with disabilities to achieve and maintain a good level of health (5).

The findings in this report are subject to at least five limitations. First, BRFSS does not include persons living in institutions or group homes. Therefore, because persons with disabilities are likely to reside in such facilities, the results likely

underestimate the actual prevalence of adults with a disability. Second, the BRFSS questions used to define disability do not collect information on the type, severity, duration, or permanence of disability. Therefore, the definition of disability used in this analysis might have captured some persons with relatively minor or short-term disabilities (e.g., a sprained ankle). Third, because of the cross-sectional nature of the data, inferring any direction of causality between disability and fair or poor health is not possible. Fourth, BRFSS is conducted only in English and Spanish, which might preclude participation by persons who speak other languages. In addition, differences in the Spanish translation of the questionnaire might explain some of the health disparities observed in the Hispanic population (6). The Spanish language version of BRFSS uses the Spanish word “regular” for the category of “fair” health, an idiomatic difference that might alter the way the participant understands the question. Finally, racial/ethnic differences in self-rated health and disability might reflect differences in potentially confounding factors, such as education, income, and health insurance status, which are significantly associated with both race/ethnicity and disability and were not controlled for in this analysis (7,8). This is a direction for future work that CDC plans to undertake.

Despite efforts to identify and reduce health disparities among racial/ethnic populations in the United States, disproportionately high rates of disability and self-rated fair or poor health persist among certain racial/ethnic populations (9,10). Efforts to reduce health disparities among racial/ethnic popula-

TABLE 2. Disability among U.S. adults aged ≥18 years, by race/ethnicity and self-rated health status — Behavioral Risk Factor Surveillance System, United States,* 2004–2006

Race/Ethnicity† and self-rated health status	Disability§				No disability				Absolute % point difference
	Sample population	Weighted U.S. population¶	%**	SE††	Sample population	Weighted U.S. population	%	SE	
White									
Excellent or very good	45,799	8,461,103	29.7	0.3	380,061	79,708,577	66.9	0.1	37.2
Good	62,212	10,542,506	33.4	0.3	160,699	31,834,243	26.8	0.1	6.6
Fair or poor	86,670	13,289,406	36.9	0.3	41,643	7,423,348	6.3	0.1	30.6
Subtotal	194,681	32,293,015	100.0		582,403	118,966,168	100.0		
Black									
Excellent or very good	2,806	769,406	21.8	0.8	29,305	8,577,102	49.9	0.4	28.1
Good	5,229	1,244,095	31.6	0.8	22,858	5,805,151	36.4	0.4	4.8
Fair or poor	10,527	2,137,184	46.6	0.8	8,825	1,988,691	13.6	0.3	33.0
Subtotal	18,562	4,150,685	100.0		60,988	16,370,944	100.0		
Hispanic									
Excellent or very good	1,951	726,628	17.3	0.7	23,622	10,803,926	37.0	0.4	19.7
Good	3,220	1,174,570	27.5	0.9	22,670	10,525,422	37.1	0.4	9.6
Fair or poor	8,348	2,528,028	55.2	1.0	13,247	6,183,780	25.9	0.4	29.3
Subtotal	13,519	4,429,226	100.0		59,539	27,513,128	100.0		
Asian									
Excellent or very good	352	169,960	36.2	3.1	7,137	3,038,367	58.7	1.0	22.5
Good	546	197,675	38.9	3.3	4,465	1,544,372	33.3	1.0	5.6
Fair or poor	564	139,986	24.9	2.3	973	320,879	8.1	0.7	16.8
Subtotal	1,462	507,621	100.0		12,575	4,903,618	100.0		
Native Hawaiian or Other Pacific Islander									
Excellent or very good	82	24,318	22.3	4.8	944	406,170	62.4	3.1	40.1
Good	113	43,046	41.3	6.5	565	181,304	26.8	2.3	14.5
Fair or poor	151	37,196	36.5	6.6	143	41,172	10.8	2.5	25.7
Subtotal	346	104,560	100.0		1,652	628,646	100.0		
American Indian/Alaska Native									
Excellent or very good	717	135,771	22.4	2.2	4,867	838,701	51.6	1.2	29.2
Good	1,236	177,565	27.1	1.7	3,805	552,187	35.8	1.2	8.7
Fair or poor	2,403	355,375	50.5	2.0	1,522	187,744	12.6	0.7	37.9
Subtotal	4,356	668,711	100.0		10,194	1,578,632	100.0		
All racial/ethnic populations									
Excellent or very good	53,166	10,610,265	27.2	0.3	454,452	105,528,004	60.2	0.1	33.0
Good	74,723	13,800,748	32.5	0.3	219,798	51,501,505	29.9	0.1	2.6
Fair or poor	112,511	19,159,470	40.3	0.3	67,947	16,493,297	9.9	0.1	30.4
Total§§	240,400	43,570,483	100.0		742,197	173,522,806	100.0		

* Includes the District of Columbia, Puerto Rico, and the U.S. Virgin Islands. Hawaii did not collect data in 2004.

† Persons identified as white, black, Asian, Native Hawaiian or Other Pacific Islander, and American Indian/Alaska Native are all non-Hispanic. Persons identified as Hispanic might be of any race.

§ Based on a "yes" response to either of the following questions: "Are you limited in any way in any activities because of physical, mental, or emotional problems?" and "Do you now have any health problem that requires you to use special equipment, such as a cane, a wheelchair, a special bed, or a special telephone?"

¶ Weighted population estimates were determined by taking the final weights for each year during 2004–2006 and dividing by three. Data were weighted to compensate for unequal probabilities of selection, to adjust for nonresponse and telephone noncoverage, to ensure that results were consistent with population data, and to make population estimates. Additional information available at <http://health.utah.gov/oph/ibishelp/brfss/issues.htm>.

** Age adjusted to the 2000 U.S. standard population.

†† Standard error.

§§ Sample population and weighted estimates by race/ethnicity do not sum to column total because respondents who reported being multiracial or of other race were included in the total.

tions should also address the needs of adults with disabilities. Such efforts must ensure that persons with disabilities have accessible, available, and appropriate health-care and wellness promotion services (5).

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HIV Prevalence Estimates — United States, 2006

Accurate and timely data on the number of persons in the United States living with human immunodeficiency virus (HIV) infection (HIV prevalence) are needed to guide planning for disease prevention, program evaluation, and resource allocation. However, overall HIV prevalence cannot be measured directly because a proportion of persons infected with HIV have neither been diagnosed nor reported to local surveillance programs. In addition, national HIV prevalence data are incomplete because local reporting systems for confidential, name-based HIV reporting have been fully implemented only since April 2008. With the advent of highly active antiretroviral therapies that delay the progression of HIV to acquired immunodeficiency syndrome (AIDS), and of AIDS to death (1), and changes in the AIDS case definition to include an immunologic diagnosis (2), earlier back-calculation methods from the 1990s for estimating HIV prevalence based on the number of reported AIDS cases are no longer reliable. With 80% of states reporting name-based HIV diagnoses as of January 2006, an extended back-calculation method now can

be used to estimate HIV prevalence more accurately. Based on this method, CDC now estimates that 1.1 million adults and adolescents (prevalence rate: 447.8 per 100,000 population) were living with diagnosed or undiagnosed HIV infection in the United States at the end of 2006. The majority of those living with HIV were nonwhite (65.4%), and nearly half (48.1%) were men who have sex with men (MSM). The HIV prevalence rates for blacks (1,715.1 per 100,000) and Hispanics (585.3 per 100,000) were, respectively, 7.6 and 2.6 times the rate for whites (224.3 per 100,000).

An extended back-calculation method has been described in detail and was used recently to calculate the incidence of HIV infection in the United States (3). The method was used in this analysis to estimate HIV prevalence based on the number of HIV diagnoses by calendar year and disease severity (i.e., whether the person received an AIDS diagnosis in the same calendar year as the HIV diagnosis). HIV prevalence at the end of 2006 for the 50 states and District of Columbia was estimated using information from the national HIV/AIDS Reporting System for persons aged ≥ 13 years who were diagnosed with HIV during 2006 and reported to CDC by the end of June 2007. Forty states provided data on both HIV and AIDS diagnoses, whereas 10 states (California, Delaware, Hawaii, Illinois, Maryland, Massachusetts, Montana, Oregon, Rhode Island, and Vermont) and the District of Columbia provided data only for AIDS diagnoses. For the areas without name-based HIV data, statistical procedures and AIDS data were used to estimate HIV cases, based on the ratio of HIV to AIDS in states with integrated surveillance systems (4). The number of undiagnosed HIV infections was calculated by subtracting diagnosed AIDS prevalence and diagnosed HIV prevalence from the estimated overall HIV prevalence. Using an established method, data also were adjusted for reporting delays and redistribution of risk factors among persons initially reported without sufficient information to be classified into an HIV transmission category (5). HIV prevalence rates per 100,000 population were calculated for various demographic characteristics; population denominators for rate calculations were based on official postcensus estimates for 2006 (6).

Among the estimated number of persons living with HIV at the end of 2006, 46.1% (1,715.1 per 100,000 population) were black, 34.6% (224.3 per 100,000) were white, 17.5% (585.3 per 100,000) were Hispanic, 1.4% (129.6 per 100,000) were Asian/Pacific Islander, and 0.4% (231.4 per 100,000) were American Indian/Alaska Native (Table). Males accounted for 74.8% of prevalent HIV cases (685.7 per 100,000). The greatest percentage of cases was attributed to male-to-male sexual contact, accounting for 48.1% overall (and 64.3% among men). High-risk heterosexual contact, defined as heterosexual contact with a person known to have, or to be at high risk for,

TABLE. Estimated number,* percentage, and rate† of persons aged ≥13 years living with human immunodeficiency virus (HIV) infection, by selected characteristics — United States, 2006

Characteristic	HIV prevalence	(95% CI [§])	%	Rate	(95% CI)
Sex					
Male	828,000	(786,000–870,000)	74.8	685.7	(650.9–720.5)
Female	278,400	(253,400–303,400)	25.2	220.4	(200.6–240.2)
Age group (yrs)					
13–24	56,500	(45,000–68,000)	5.1	111.0	(88.4–133.6)
25–49	770,000	(730,000–810,000)	69.6	720.4	(683.0–757.9)
≥50	280,000	(255,000–305,000)	25.3	313.5	(285.5–341.4)
Race/Ethnicity					
White	382,600	(354,600–410,600)	34.6	224.3	(207.9–240.7)
Black	510,100	(478,100–542,100)	46.1	1,715.1	(1,607.5–1,822.7)
Hispanic [¶]	194,000	(175,000–213,000)	17.5	585.3	(528.0–642.6)
Asian/Pacific Islander	15,100	(12,600–17,600)	1.4	129.6	(108.2–151.1)
American Indian/Alaska Native	4,600	(3,100–6,100)	0.4	231.4	(156.0–306.9)
HIV transmission category					
Male-to-male sexual contact	532,000	(492,000–572,000)	48.1		
Injection drug use (male)	131,500	(114,500–148,500)	11.9		
Injection drug use (female)	73,100	(62,100–84,100)	6.6		
Male-to-male sexual contact and injection drug use	54,900	(44,900–64,900)	5.0		
High-risk heterosexual contact (male)**	104,000	(89,000–119,000)	9.4		
High-risk heterosexual contact (female)**	201,700	(179,700–223,700)	18.2		
Other ^{††}	9,100	(7,600–10,600)	0.8		
Total^{§§}	1,106,400	(1,056,400–1,156,400)	100	447.8	(427.5–468.0)

* Estimated numbers, from national HIV/AIDS Reporting System data, are adjusted for reporting delays and reclassification of cases reported without information regarding an HIV transmission category, but are not adjusted for underreporting. Estimates are rounded to the nearest 100.

† Per 100,000 population at the end of 2006. Rates for transmission category subgroups were not calculated because population denominators were unavailable. Rates for racial/ethnic populations do not include an adjustment for redistribution of persons of unknown race/ethnicity.

§ Confidence interval.

¶ Might be of any race.

** Heterosexual contact with a person known to have, or to be at high risk for, HIV infection.

†† Includes hemophilia, blood transfusion, perinatal exposure, and risk factors not reported or not identified.

§§ Because column totals were calculated independently of the values of the subpopulations and all values were rounded, the values might not sum to the respective column total.

HIV infection (e.g., an injection drug user) accounted for 27.6% of prevalent cases overall (12.6% of cases among men and 72.4% of cases among women). Injection drug use (IDU) accounted for 18.5% of total cases (15.9% of cases among men and 26.3% of cases among women). The remainder of cases were attributed to men who reported both male-to-male sexual contact and IDU (5.0%) or whose transmission category was classified as other (0.8%; including hemophilia, blood transfusion, perinatal exposure, and risk factors not reported or not identified). Overall, an estimated 232,700 (21.0%) persons living with HIV infection had not been diagnosed as of the end of 2006.

The HIV prevalence rate for black men (2,388.2 per 100,000 population; 95% confidence interval [CI] = 2,197.9–2,578.4) was six times the rate for white men (394.6 per 100,000; CI = 363.3–425.9) (Figure), and the rate for Hispanic men (883.4 per 100,000; CI = 784.9–982.4) was more than twice the rate for white men. The HIV prevalence rate for black women (1,122.4 per 100,000; CI = 1,002.2–1,242.5) was nearly 18 times the rate for white women (62.7 per 100,000; CI = 54.7–70.7), and the rate for Hispanic women (263.0 per 100,000; CI = 231.6–294.4) was more than four times

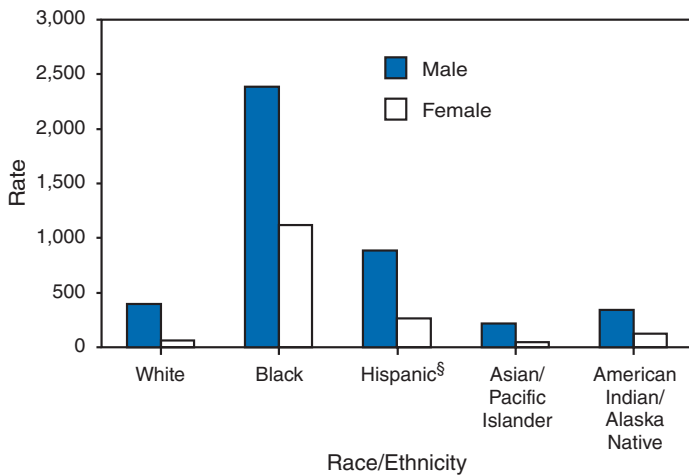
the rate for white women. The HIV prevalence rate for black women was greater than the rate for all other groups, except for black men.

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Editorial Note: Reduced mortality resulting from the use of highly active antiretroviral therapies is a major factor contributing to the number of persons in the United States living with HIV disease (1). Additionally, more than 56,000 new HIV infections are estimated to occur annually (3).

The estimate of HIV prevalence in this report is similar to an estimate for 2003 (1,039,000–1,185,000) that used the same extended back-calculation method (4). However, because of improvements in national HIV surveillance data since 2003, the two estimates cannot be compared directly. The 2006 estimate is based on a data set that 1) includes HIV diagnoses from 10 states that were not reporting in 2003 and 2) has been refined by an improved ability to identify and remove duplicate HIV case data that reflect reports by more than one state. Using the refined data set, CDC now estimates the HIV prevalence for 2003 to have been 994,000, suggesting that HIV

FIGURE. Estimated human immunodeficiency virus (HIV) prevalence rate* among persons aged ≥ 13 years, by race/ethnicity and sex — United States, 2006†



* Per 100,000 population.

† HIV prevalence at the end of 2006 for the 50 states and the District of Columbia, estimated from national HIV/AIDS Reporting System data.

§ Might be of any race.

prevalence in the United States increased by approximately 112,000 (11.3%) from 2003 to 2006. Analysis of the refined data also indicated that the percentage of HIV-positive persons who were undiagnosed decreased from approximately 25% in 2003 to 21% in 2006; an estimated 30% of this change resulted from a decrease in the number of undiagnosed persons, and 70% resulted from an increase in the total number of persons living with HIV (CDC, unpublished data, 2008).

The burden of HIV infection was disproportionate among populations. Blacks made up 12% of the adult and adolescent population in the United States in 2006 (6), but accounted for 46.1% of persons estimated to be living with HIV. Similarly, nearly half (48.1%) of the persons living with HIV were MSM, and although not precisely known, the percentage of MSM in the general population is estimated to be much lower. Data from CDC's National Survey of Family Growth indicate that, among males aged 15–44 years, 3.7% ever have had anal sex with another male, and the proportion of men who had a male sexual partner in the past 12 months was 2.9% (7).

The findings in this report are subject to at least three limitations. First, reported HIV data used in the extended back-calculation method represent only a portion of persons in the United States who were diagnosed with HIV infection; several high-morbidity areas, including California, Illinois, Maryland, and the District of Columbia, did not contribute HIV data. Availability of reported HIV data from these areas will increase accuracy of future prevalence estimates. Second, not all persons who are infected with HIV have been diagnosed and reported to the public health surveillance system, and

data must be estimated for undiagnosed persons. Finally, the data have been adjusted statistically to account for delays in reporting new cases and deaths, and cases reported without risk factor information have been redistributed among other transmission categories (5). These adjustments were based on risk redistribution assumptions from the mid-1990s that might no longer be valid, which could result in over- or under-adjustment of the data.

Previous studies have indicated that persons generally reduce their sexual risk behaviors (e.g., decrease the number of sex partners and reduce unprotected intercourse through increased condom use) after being diagnosed with HIV (8). Thus, increasing the percentage of HIV-infected persons who are diagnosed and linked with effective care and prevention services has the potential to reduce new HIV infections over time. To help achieve that, CDC has focused resources on increasing testing for HIV, particularly among populations that are disproportionately affected by HIV infection. Recent CDC activities have included publication of revised recommendations for HIV testing in health-care settings (9) and creation of a new program, the Heightened National Response to the HIV/AIDS Crisis in the African American Community (10). In 2007, as part of the President's Domestic HIV Initiative, CDC allocated funds to expand routine HIV testing, primarily among blacks. In addition to testing, expanding the number and reach of effective HIV prevention services for at-risk populations, including blacks, Hispanics, and MSM of all races, can contribute to reducing the disproportionate numbers of infections in these groups. Culturally appropriate opportunities for HIV testing, diagnosis, and access to early treatment and prevention services to reduce further HIV transmission are key to reducing new infections and ultimately decreasing HIV prevalence in the United States.

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Rabies in a Dog Imported from Iraq – New Jersey, June 2008

Rabies vaccination and stray dog control have led to successful control of canine rabies in the United States. The number of rabid dogs reported decreased from approximately 5,000 in 1950 to 79 in 2006, when the canine rabies virus variant associated with dog-to-dog rabies transmission was declared eliminated in the United States (1). On June 18, 2008, a mixed-breed dog, recently shipped from Iraq into the United States, was confirmed to have rabies by the Public Health and Environmental Laboratories of the New Jersey Department of Health and Senior Services. A total of 24 additional animals in the shipment, all potentially exposed to the rabid dog, were distributed to 16 states. This report summarizes the epidemiologic investigation by the New Jersey Department of Health and Senior Services, Bergen County Department of Health, and CDC, and the ensuing public health response. These findings underscore the need for vigilance regarding rabies (and other zoonotic diseases) during animal importation to prevent the possible reintroduction and sustained transmission of canine rabies in U.S. dog populations.

Case Report

On June 5, 2008, a shipment of 24 dogs and two cats arrived in the United States from Iraq as part of an international animal rescue operation. The goal of the operation was to reunite servicemen returning to the United States with animals they had adopted in Iraq. Upon arrival at Newark Liberty International Airport, the animals received physical examinations from volunteer licensed veterinarians. One cat became ill with neurologic signs during transport and was euthanized on arrival. The cat was tested for rabies and was negative. The remaining 24 dogs and one cat were housed for several days at the airport before distribution to their final U.S. destinations.

On June 8, one of the 24 dogs, a mixed-breed aged 11 months (dog A), became ill and was taken to a veterinarian the

next day. The dog was hospitalized with fever, diarrhea, wobbly gait, agitation, and crying. The dog's condition deteriorated, progressing to lateral recumbency with periods of agitation. On June 11, the dog was euthanized. Specimens were shipped to the Public Health and Environmental Laboratories for rabies testing, but delivery of the specimens was delayed. On June 18, the specimens were tested, and rabies was diagnosed. Specimens also were submitted to CDC, where rabies was confirmed on June 26 and typed as a rabies virus variant associated with dogs in the Middle East.

Public Health Investigation

The potentially infectious period for a dog, cat, or ferret with rabies can begin as many as 10 days before the onset of clinical signs and continue throughout the clinical course until death (2). To identify potential rabies exposure to humans or other animals while dog A was in Iraq, during transport, or at the airport shelter, an investigation was initiated by the New Jersey Department of Health and Senior Services and the Bergen County Department of Health, with participation from CDC. The dog was reportedly in the possession of a U.S. soldier in Baghdad for approximately 7 months before shipment to the United States. The dog had been kept in an indoor-outdoor run on a military base and had not been vaccinated for rabies; the owner reported no signs of illness in the dog or potential exposure to other rabid animals during the 7 months. The owner also reported no potential exposures to other persons or animals during the 2 days of potential infectivity before the dog was transferred to the animal rescue operation for shipment on May 31.

Upon arrival in the United States, none of the 24 dogs were accompanied by the valid rabies vaccination certificates required for admission by CDC animal importation regulations.* For dogs aged ≥ 3 months, a rabies vaccination must be administered at least 30 days before the date of arrival at a U.S. port. Five of the 24 dogs (not including dog A) reportedly had received a previous rabies vaccination; however, none of the information required for a valid rabies vaccination certificate was available, including vaccine manufacturer, lot numbers, or a certifying veterinarian signature. Twenty-one of the animals in the shipment, including dog A, had received a primary rabies vaccination in Iraq during May 28–31, immediately before being shipped to New Jersey. Because none of the dogs met rabies vaccination requirements for importation, in accordance with the importation regulation, a confinement agreement was issued by CDC, stating where the animals would be held for at least 30 days after vaccination. During shipment and

*42 CFR § 71.51.

upon arrival in New Jersey, all the animals were housed in separate crates; however, interviews with persons present during the animals' arrival and stay in Newark identified potential periods during which dogs, including dog A, were allowed to intermingle.

On June 10, 1 day before dog A was euthanized and 8 days before rabies was diagnosed, the remaining 23 dogs and one cat were shipped to destinations in 16 states.[†] Because none of the surviving animals had a verifiable history of vaccination at least 30 days before their potential exposure to dog A, CDC recommended immediate vaccination and a 6-month quarantine for all of them (2). State health departments in the 16 states were advised of the recommendations.

During the public health investigation, 28 persons were evaluated for potential rabies exposure; 13 were identified with potential exposure because of direct contact with possibly infectious saliva (3) and were recommended to initiate rabies postexposure prophylaxis (PEP). All 23 dogs and one cat were located by state and local health authorities within 2 weeks of the rabies diagnosis. No clinical signs consistent with rabies were reported in the animals during 20 days of follow-up. All 24 animals continue to be monitored during the 6-month quarantine period.

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Editorial Note: Rabies virus infection results in a fatal encephalomyelitis in humans and other mammals. Globally, the most common sources of human rabies are geographically distinct rabies virus variants maintained predominantly through dog-to-dog transmission (i.e., canine rabies), but sometimes with spillover[§] into other species. In the United States, occasional spillover into dogs of rabies virus variants associated with wildlife has occurred. However, since 2004, no rabies case attributable to an indigenously acquired canine rabies virus variant has been reported (1).

Canine rabies virus variants most commonly are imported via unvaccinated dogs from areas where rabies is enzootic, such as Asia, Africa, the Middle East, and parts of Latin America, where canine variants are responsible for most of the 55,000 human rabies deaths estimated worldwide each year (4). In

May 2004, an unvaccinated puppy was flown from Puerto Rico to Massachusetts as part of an animal rescue program. The day after arrival, the puppy exhibited neurologic signs, was euthanized, and was subsequently confirmed to have rabies. Six persons were recommended to receive PEP because of potential exposure. In June 2004, an unvaccinated puppy adopted by a U.S. resident in Thailand was confirmed to have rabies by the California Department of Public Health. Of 40 persons interviewed for potential rabies exposure, 12 received PEP. In March 2007, a puppy adopted by a U.S. veterinarian while volunteering in India was confirmed to have rabies by the Alaska Department of Health and Social Services. The puppy was flown in cargo to Seattle, Washington, then adopted by another veterinarian in Juneau, Alaska, where it was flown 7 days after arrival. Of 20 persons interviewed for potential rabies exposure, eight received PEP (5,6). In all three cases, the rabies virus variant was typed as a variant circulating in dogs and terrestrial wildlife in the animal's country of origin (i.e., mongoose and canine rabies virus variants enzootic in Puerto Rico, Thailand, and India, respectively).

This report reiterates the need for education of the public regarding rabies incidence in other countries and preventing rabies exposure. While traveling in areas that are endemic for rabies, travelers should not pet stray animals. In addition, travelers should not adopt stray animals without acquiring a veterinarian's health assessment and ensuring proper animal vaccination for importation. Travelers also should consider their potential for rabies exposure from animals, understand proper wound management, and promptly report animal bites to health-care providers (7). Health information for travelers is available at <http://wwwn.cdc.gov/travel/contentyellowbook.aspx>.

CDC administers federal importation regulations for dogs. These regulations allow admittance of unvaccinated dogs aged <3 months, provided the importer signs an agreement to vaccinate the dog at age 3 months and confine the animal for 30 days after the vaccination. Dogs aged ≥3 months that have not been vaccinated for rabies also must be confined until vaccinated and for 3 months after vaccination. Upon arrival in the United States, importers should declare animals to federal authorities and comply with those requirements for confinement of unvaccinated puppies.

CDC's regulations were created in the early 1950s to guide persons importing dogs or cats as their personal pets. However, recent trends in dog importations have shown an increase in the numbers of animals being imported for commercial pet trade (8). CDC is working to update current regulations and better address the importation of dogs. In July 2007, the U.S. Department of Health and Human Services posted an advance notice of proposed rulemaking to begin the process of revising

[†] California, Colorado, Connecticut, Iowa, Kentucky, Maryland, Massachusetts, Missouri, North Carolina, Ohio, Oklahoma, Pennsylvania, South Carolina, Texas, Virginia, and Washington.

[§] Transmission of a rabies virus variant to a secondary host from a primary reservoir species, usually resulting in a dead-end infection, such as human rabies acquired from a rabid dog.

CDC's animal importation regulations, including those that apply to dogs and other companion animals.[†]

U.S. animal importation regulations, rabies vaccination requirements for dogs, wildlife rabies surveillance and vaccination programs, and prophylaxis for human exposures all contribute to public health protection from rabies. Continued vigilance and partnership between federal and state agencies, as well as health professionals and pet importers, are vital to decrease the risk for reemergence of canine rabies virus in the United States.

[†] Available at <http://www.cdc.gov/ncidod/dq/anprm/index.htm>.

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Licensure of a Diphtheria and Tetanus Toxoids and Acellular Pertussis Adsorbed and Inactivated Poliovirus Vaccine and Guidance for Use as a Booster Dose

On June 24, 2008, the Food and Drug Administration licensed a combined diphtheria and tetanus toxoids and acellular pertussis adsorbed (DTaP) and inactivated poliovirus (IPV) vaccine, DTaP-IPV (Kinrix, GlaxoSmithKline Biologicals, Rixensart, Belgium). Kinrix is licensed for use as the fifth dose of the DTaP vaccine series and the fourth dose of the IPV series in children aged 4–6 years whose previous DTaP vaccine doses were DTaP (Infanrix, GlaxoSmithKline) and/or DTaP-Hepatitis B-IPV (Pediarix, GlaxoSmithKline) for the first 3 doses and DTaP (Infanrix) for the fourth dose (1,2). DTaP-IPV administered to children aged 4–6 years would reduce by one the number of injections needed to complete DTaP and IPV immunization. This report summarizes the indications for Kinrix and provides guidance from the Advisory Committee on Immunization Practices (ACIP) for its use.

ACIP reviewed data on the safety and immunogenicity of DTaP-IPV (Kinrix). On the basis of these data, expert opinion of the ACIP Combination Vaccines Workgroup, and feedback from ACIP liaison organizations including the American Academy of Pediatrics and the American Academy of Family Physicians, ACIP endorsed the licensed indications and offered the following guidance for use of DTaP-IPV. On June 26, ACIP voted to include DTaP-IPV in the federal Vaccines for Children Program.

The individual antigens (diphtheria, tetanus, and pertussis toxoids, filamentous hemagglutinin, pertactin, and poliovirus types 1, 2, and 3) contained in combined DTaP-IPV are identical to the antigens contained in GlaxoSmithKline's DTaP (Infanrix) and DTaP-Hepatitis B-IPV (Pediarix) and have been described previously (3). DTaP-IPV contains no preservatives. DTaP-IPV is administered as an intramuscular injection, preferably into the deltoid region. Two clinical trials conducted in U.S. children aged 4–6 years showed that combined DTaP-IPV and separately administered DTaP and IPV vaccines had comparable safety and reactogenicity profiles, with or without a co-administered second dose of measles, mumps, and rubella (MMR) vaccine (3,4). The immunogenicity of all antigens was similar between the treatment groups, with or without a co-administered second dose of MMR vaccine.

Indications and Guidance for Use

DTaP-IPV (Kinrix) is indicated for use as the fifth dose of DTaP and fourth dose of IPV in children aged 4–6 years

who received DTaP (Infanrix) and/or DTaP-Hepatitis B-IPV (Pediatrix) as the first 3 doses and DTaP (Infanrix) as the fourth dose (1,2). This vaccine should not be administered to children aged <4 years or ≥ 7 years; however, if DTaP-IPV (Kinrix) is inadvertently administered for an earlier dose of the DTaP and/or IPV series, the dose should be counted as valid and does not need to be repeated provided minimum interval requirements have been met (5). Data are limited on the safety and immunogenicity of interchanging DTaP vaccines from different manufacturers (6). ACIP recommends that, whenever feasible, the same manufacturer's DTaP vaccines should be used for each dose in the series; however, vaccination should not be deferred because the type of DTaP previously administered is unavailable or unknown (6).

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months (1,2). This report summarizes the indications for Pentacel and provides guidance from the Advisory Committee on Immunization Practices (ACIP) for its use.

ACIP reviewed data on the safety and immunogenicity of DTaP-IPV/Hib (Pentacel). On the basis of these data, expert opinion of the ACIP Combination Vaccines Workgroup, and feedback from ACIP liaison organizations including the American Academy of Pediatrics and the American Academy of Family Physicians, ACIP endorsed the licensed indications and offered the following guidance for use of DTaP-IPV/Hib. On June 26, ACIP voted to include DTaP-IPV/Hib in the federal Vaccines for Children Program.

Each dose of DTaP-IPV/Hib contains the same diphtheria and tetanus toxoids and pertussis antigens (inactivated pertussis toxin [PT], filamentous hemagglutinin [FHA], pertactin, and fimbriae types 2 and 3) as the FDA-licensed DTaP vaccine Daptacel (Sanofi Pasteur, Toronto, Canada) but contains an increased amount of inactivated PT and FHA (2). The poliovirus component of DTaP-IPV/Hib contains the same strains and amount of inactivated poliovirus types 1, 2, and 3 as the polio vaccine Poliovax (Sanofi Pasteur, Toronto, Canada) (2). The Hib component is identical to ActHib (*Haemophilus influenzae* type b capsular polysaccharide [polyribosyl-ribitol-phosphate {PRP}] covalently bound to tetanus toxoid) (Sanofi Pasteur, Swiftwater, Pennsylvania) (2). The DTaP-IPV component is supplied as a sterile liquid used to reconstitute a lyophilized ActHIB vaccine component. Components should not be administered separately. DTaP-IPV/Hib does not contain thimerosal.

In comparative studies, the frequency of solicited local and systemic adverse events and of serious adverse events after administration of DTaP-IPV/Hib was similar to that observed following separately administered DTaP, IPV, and Hib component vaccines (2,3). The immunologic responses after the third dose or the fourth dose of DTaP-IPV-Hib generally were comparable to those following separately administered component vaccines, and have been published (2,3). Immune responses following the first and second doses were not measured.

Indications and Guidance for Use

DTaP-IPV/Hib is licensed for use in children aged 6 weeks through 4 years. DTaP-IPV/Hib is indicated for use in infants and children at ages 2, 4, 6, and 15–18 months (1). DTaP-IPV/Hib is not licensed for use in children aged ≥ 5 years, and is not indicated for the booster dose at age 4–6 years (2). However, DTaP-IPV/Hib that is inadvertently administered to children aged ≥ 5 years should be counted as a valid dose.

For prevention of diphtheria, tetanus, and pertussis, all children are recommended to receive 4 doses of DTaP, at ages 2, 4, 6, and 15–18 months, and a booster dose at age 4–6 years.

Licensure of a Diphtheria and Tetanus Toxoids and Acellular Pertussis Adsorbed, Inactivated Poliovirus, and *Haemophilus b* Conjugate Vaccine and Guidance for Use in Infants and Children

On June 20, 2008 the Food and Drug Administration (FDA) licensed a combined diphtheria and tetanus toxoids and acellular pertussis adsorbed (DTaP), inactivated poliovirus vaccine (IPV), and *Haemophilus influenzae* type b conjugate (tetanus toxoid [TT] conjugate) vaccine, DTaP-IPV/Hib (Pentacel, Sanofi Pasteur, Swiftwater, Pennsylvania), for use as a four-dose series in infants and children at ages 2, 4, 6, and 15–18

Although an 8-week interval between doses is preferred, if an accelerated schedule is needed, a minimum interval of 4 weeks should occur between the first and second doses, and the third dose should not be administered before age 14 weeks (4). The fourth dose of DTaP-IPV/Hib may be administered as early as 12 months of age if the clinician feels an opportunity to vaccinate may be missed later and if 6 months has elapsed since the third dose of DTaP-IPV/Hib (1).

Data are limited on the safety and immunogenicity of interchanging DTaP vaccines from different manufacturers (2). ACIP recommends that, whenever feasible, the same manufacturer's DTaP product should be used for the pertussis series; however, that vaccination should not be deferred if the specific DTaP vaccine brand previously administered is unavailable or unknown (2).

For prevention of poliomyelitis, all children are recommended to receive 4 doses of IPV, at ages 2, 4, 6–18 months, and 4–6 years. DTaP-IPV/Hib may be used for 1 or more doses of the IPV series, including in children who have received 1 or more doses of another licensed IPV vaccine and who also are scheduled to receive DTaP and Hib vaccination. When an accelerated or catch-up schedule is needed, IPV doses may be administered at 4-week intervals and the fourth dose counted as valid if administered as early as age 18 weeks when the proper spacing of prior doses is maintained (1). Therefore, DTaP-IPV/Hib (Pentacel) doses administered at 2, 4, 6, and 12–18 months would provide 4 valid doses of IPV under these circumstances.

The recommended vaccination schedule for Hib-TT vaccines (e.g., Pentacel) consists of a 3-dose primary series at ages 2, 4, and 6 months, and a booster dose at age 12–15 months (1). Intervals between doses of the primary series as short as 1 month are acceptable but not optimal. Minimum intervals for the booster dose vary by age at first vaccination and have been published (5). DTaP-IPV/Hib may be administered at 12 months and counted as a valid Hib-TT dose if the minimum intervals are followed; however, the safety and efficacy of DTaP-IPV/Hib in this circumstance have not been evaluated. DTaP-IPV/Hib may be administered at separate injection sites with other vaccines administered at age 12–18 months, such as hepatitis A, hepatitis B, pneumococcal conjugate, measles, mumps, and rubella (MMR), and varicella vaccines (2).

Special Considerations

Certain American Indian/Alaska Native (AI/AN) children are at increased risk for Hib disease, particularly in the first 6 months of life (6). Furthermore, the immunologic response to different Hib conjugate vaccine preparations can vary. Compared with other Hib conjugate vaccines (e.g., Hib-TT),

administration of polyribosylribitol phosphate-meningococcal outer membrane protein (PRP-OMP)-containing Hib vaccine preparations leads to a more rapid seroconversion to protective antibody concentrations within the first 6 months of life. Although for subsequent doses, PRP-OMP and other Hib conjugate vaccines appear to have equal efficacy, failure to use PRP-OMP vaccines for the first dose has been associated with excess cases of Hib disease in AI/AN infants living in communities where Hib transmission is ongoing and exposure to colonized persons is likely (6,7). In addition, stocking of both PRP-OMP and other Hib conjugate vaccine preparations in the same clinic might lead to inadvertent administration of another vaccine for the first Hib dose. For this reason, clinics that serve predominantly AI/AN children might elect to stock and use only PRP-OMP-containing Hib vaccines (6).

Different lot numbers for the different components of DTaP-IPV/Hib are included on the DTaP-IPV vial and on the Hib powder vial. Providers should record lot numbers separately for the DTaP-IPV and Hib components.

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Notice to Readers

Get Smart About Antibiotics Week — October 6–10, 2008

October 6–10 is Get Smart About Antibiotics Week. The theme of this observance is “The power to prevent resistance is in your hands.”

Inappropriate use of antibiotics to treat upper respiratory infections (URIs) can result in unnecessary risk for adverse events and contribute to the likelihood of antibiotic resistance. Adverse events related to antibiotics (usually aller-

gies or drug intolerance) resulted in an estimated 142,500 emergency department visits annually in the United States during 2004–2006 (1). In addition, inappropriate and excessive antimicrobial use can increase a community's risk for antibiotic-resistant bacterial infections that might lead to severe or prolonged illness, hospitalization, and sometimes death. Educating clinicians and the public regarding appropriate use of antibiotics might help reduce adverse drug events, including antibiotic resistance.

As part of Get Smart About Antibiotics Week, health-care providers are urged to take the following actions to help reduce antibiotic resistance and other adverse drug events:

- Know when antibiotics are indicated, and avoid prescribing antibiotics for URIs such as pharyngitis, bronchitis, sinusitis, and the common cold, which are primarily caused by viruses.
- Instead of prescribing antibiotics for URIs, identify and validate patient concerns and recommend symptomatic therapy.

Additional information about Get Smart About Antibiotics Week is available at <http://www.cdc.gov/getsmart>.

Reference

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Notice to Readers

Epidemiology in Action: Intermediate Analytic Methods Course, January 12–15, 2009

CDC and Emory University's Rollins School of Public Health will cosponsor the course Epidemiology in Action: Intermediate Analytic Methods, January 12–15, 2009, at Emory University, Rollins School of Public Health, in Atlanta, Georgia. The course is designed for practicing public health professionals who have experience in basic applied epidemiology and would like training in additional quantitative skills related to analysis and interpretation of epidemiologic data.

The course includes a review of the fundamentals of descriptive epidemiology and biostatistics, measures of association, normal and binomial distributions, confounding, statistical tests, stratification, logistic regression models, and computer programs used in epidemiology.

The prerequisite is an introductory course in epidemiology, such as Epidemiology in Action or the International Course in Applied Epidemiology. Tuition will be charged. The application deadline is December 15, 2008, or until all slots have been filled.

Additional information and applications are available by mail (Emory University, Hubert Global Health Dept [Attn: Pia], 1518 Clifton Rd. NE, Rm. 746, Atlanta, GA 30322); telephone (404-727-3485); fax (404-727-4590); e-mail (pvaleri@sph.emory.edu), or Internet (<http://www.sph.emory.edu/epicourses>).

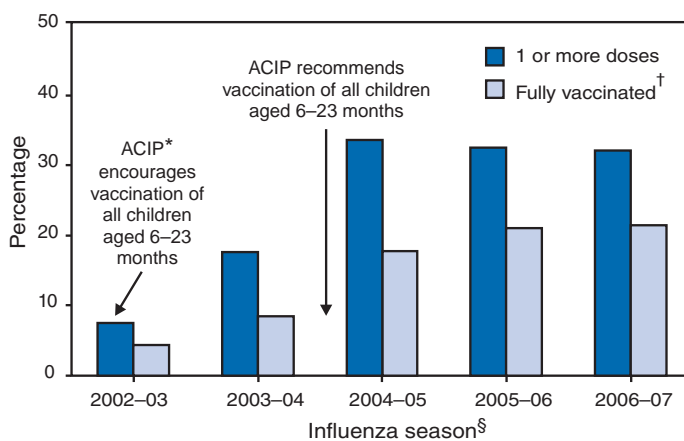
Erratum: Vol. 57, No. 36

In the report, "Subpopulation Estimates from the HIV Incidence Surveillance System — United States, 2006," an error occurred in the last sentence on page 986. The sentence should read, "Among white MSM, by age group, the largest number of new infections (4,670 [35%]) was among those aged 30–39 years (Figure)."

Erratum: Vol. 57, No. 38

In the report, "Influenza Vaccination Coverage Among Children Aged 6–23 Months — United States, 2006–07 Influenza Season," an error occurred in Figure 1 on page 1043. The corrected figure follows.

FIGURE 1. Percentage of children aged 6–23 months receiving influenza vaccination during September–December, by influenza season and vaccination status — National Immunization Survey, United States, 2002–03 to 2006–07 influenza seasons



* Advisory Committee on Immunization Practices.

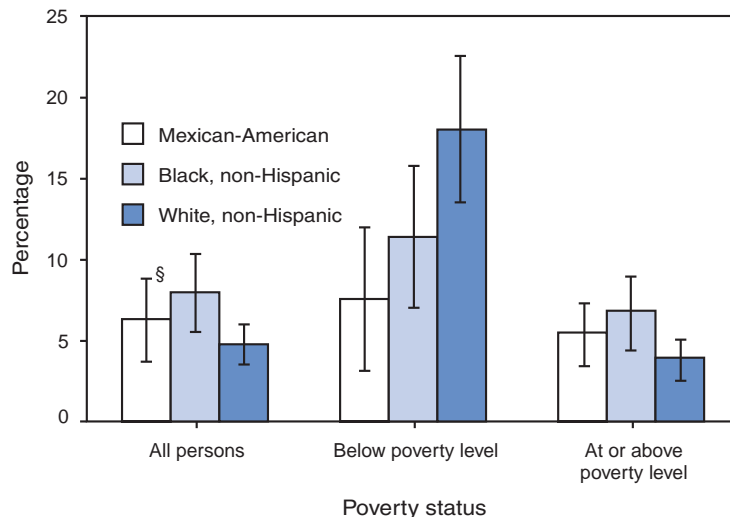
† Children were considered fully vaccinated if they had 1) received no doses of influenza vaccine before September 1 and received 2 doses from September 1 through the date of interview or January 31 (whichever was earlier), or 2) received 1 or more doses of influenza vaccine before September 1 and received 1 or more doses during September–December.

§ 2002–03 (N = 13,831); 2003–04 (N = 13,881); 2004–05 (N = 12,056); 2005–06 (N = 13,546); and 2006–07 (N = 9,710).

QuickStats

FROM THE NATIONAL CENTER FOR HEALTH STATISTICS

Percentage of Persons Aged ≥12 Years with Depression,* by Race/Ethnicity and Poverty Status† — National Health and Nutrition Examination Survey, United States, 2005–2006



* Depression was measured using the Patient Health Questionnaire (PHQ-9), a nine-item screening instrument that asks questions about the frequency of symptoms of depression during the preceding 2 weeks. Response categories “not at all,” “several days,” “more than half the days,” and “nearly every day” were given a score ranging from 0 to 3. Depression was defined as a total score of 10 or higher on the PHQ-9. This cut point has been well validated and is commonly used in clinical studies that measure depression with the PHQ-9.

† Poverty status was defined using the poverty income ratio (PIR), an index calculated by dividing the family income by a poverty threshold that is based on the size of the family. A PIR of less than 1 was used as the cut point for below the poverty level.

§ 95% confidence interval.

During 2005–2006, overall, non-Hispanic blacks had higher rates of depression (8.0%) than non-Hispanic whites (4.8%). Among persons living below the poverty level, non-Hispanic whites had higher rates of depression (18.0%) than Mexican-Americans (7.6%). Non-Hispanic blacks and non-Hispanic whites living below the poverty level had higher rates of depression than those with higher incomes, whereas rates of depression in Mexican-Americans did not vary by poverty status.

SOURCES: National Health and Nutrition Examination Survey data, 2005–2006. Available at <http://www.cdc.gov/nchs/nhanes.htm>.

Pratt LA, Brody DJ. Depression in the United States household population, 2005–2006. NCHS data brief no. 7. Hyattsville, MD: US Department of Health and Human Services, CDC, National Center for Health Statistics; 2008. Available at <http://www.cdc.gov/nchs/data/databriefs/db07.htm>.

TABLE 1. Provisional cases of infrequently reported notifiable diseases (<1,000 cases reported during the preceding year) — United States, week ending September 27, 2008 (39th week)*

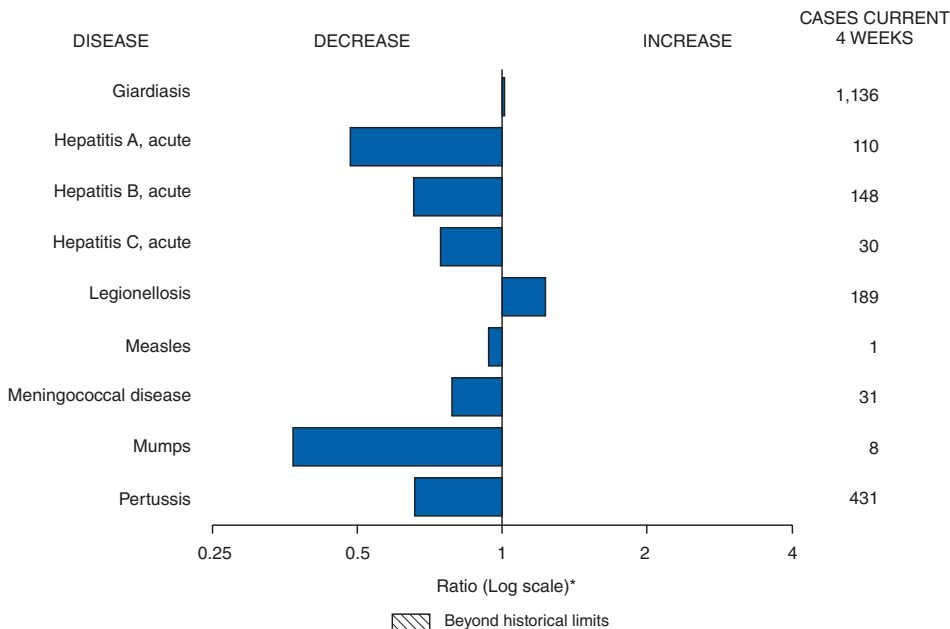
Disease	Current week	Cum 2008	5-year weekly average†	Total cases reported for previous years					States reporting cases during current week (No.)
				2007	2006	2005	2004	2003	
Anthrax	—	—	—	1	1	—	—	—	
Botulism:									
foodborne	—	6	0	32	20	19	16	20	
infant	1	69	2	85	97	85	87	76	PA (1)
other (wound & unspecified)	—	12	1	27	48	31	30	33	
Brucellosis	3	61	2	131	121	120	114	104	CA (3)
Chancroid	—	30	1	23	33	17	30	54	
Cholera	—	1	0	7	9	8	6	2	
Cyclosporiasis§	1	107	1	93	137	543	160	75	WA (1)
Diphtheria	—	—	—	—	—	—	—	1	
Domestic arboviral diseases§,¶:									
California serogroup	—	28	4	55	67	80	112	108	
eastern equine	—	2	0	4	8	21	6	14	
Powassan	—	1	—	7	1	1	1	—	
St. Louis	—	9	1	9	10	13	12	41	
western equine	—	—	—	—	—	—	—	—	
Ehrlichiosis/Anaplasmosis§,¶,¶:									
<i>Ehrlichia chaffeensis</i>	7	561	13	828	578	506	338	321	MD (1), VA (1), NC (1), GA (2), TN (2)
<i>Ehrlichia ewingii</i>	—	7	—	—	—	—	—	—	
<i>Anaplasma phagocytophilum</i>	1	224	12	834	646	786	537	362	CT (1)
undetermined	—	52	3	337	231	112	59	44	
<i>Haemophilus influenzae</i> , ††									
invasive disease (age <5 yrs):									
serotype b	1	20	0	22	29	9	19	32	MD (1)
nonserotype b	1	125	2	199	175	135	135	117	CO (1)
unknown serotype	1	142	3	180	179	217	177	227	OR (1)
Hansen disease§	—	54	2	101	66	87	105	95	
Hantavirus pulmonary syndrome§	—	11	0	32	40	26	24	26	
Hemolytic uremic syndrome, postdiarrheal§	6	147	6	292	288	221	200	178	NY (2), OK (1), CO (1), CA (2)
Hepatitis C viral, acute	8	598	17	849	766	652	720	1,102	PA (1), MI (1), NC (1), KY (2), WA (1), OR (1), CA (1)
HIV infection, pediatric (age <13 years)§§	—	—	3	—	—	380	436	504	
Influenza-associated pediatric mortality§,¶¶	—	88	0	77	43	45	—	N	
Listeriosis	9	430	21	808	884	896	753	696	NY (1), IN (1), MD (1), NC (1), FL (2), WA (1), CA (2)
Measles***	—	129	0	43	55	66	37	56	
Meningococcal disease, invasive†††:									
A, C, Y, & W-135	1	207	4	325	318	297	—	—	MD (1)
serogroup B	1	120	2	167	193	156	—	—	VA (1)
other serogroup	—	26	0	35	32	27	—	—	
unknown serogroup	10	465	9	550	651	765	—	—	NYC (2), MI (1), NE (1), DE (1), NC (1), AZ (1), WA (1), CA (2)
Mumps	3	310	16	800	6,584	314	258	231	FL (1), CA (2)
Novel influenza A virus infections	—	—	—	1	N	N	N	N	
Plague	—	1	0	7	17	8	3	1	
Poliomyelitis, paralytic	—	—	0	—	—	1	—	—	
Polio virus infection, nonparalytic§	—	—	—	—	N	N	N	N	
Psittacosis§	—	9	0	12	21	16	12	12	
Qfever§,§§§ total:	3	85	2	171	169	136	70	71	
acute	3	78	—	—	—	—	—	—	CA (3)
chronic	—	7	—	—	—	—	—	—	
Rabies, human	—	—	0	1	3	2	7	2	
Rubella¶¶¶	—	11	0	12	11	11	10	7	
Rubella, congenital syndrome	—	—	—	—	1	1	—	1	
SARS-CoV§,****	—	—	—	—	—	—	—	8	
Smallpox§	—	—	—	—	—	—	—	—	
Streptococcal toxic-shock syndrome§	1	104	1	132	125	129	132	161	NC (1)
Syphilis, congenital (age <1 yr)	—	149	8	430	349	329	353	413	
Tetanus	—	7	1	28	41	27	34	20	
Toxic-shock syndrome (staphylococcal)§	1	44	2	92	101	90	95	133	TN (1)
Trichinellosis	—	5	0	5	15	16	5	6	
Tularemia	—	79	3	137	95	154	134	129	
Typhoid fever	8	299	10	434	353	324	322	356	CT (3), VA (1), OK (1), TX (1), WA (1), CA (1)
Vancomycin-intermediate <i>Staphylococcus aureus</i> §	—	6	0	37	6	2	—	N	
Vancomycin-resistant <i>Staphylococcus aureus</i> §	—	—	—	2	1	3	1	N	
Vibriosis (noncholera <i>Vibrio</i> species infections)§	10	315	5	447	N	N	N	N	MD (2), VA (1), GA (1), FL (4), CA (2)
Yellow fever	—	—	—	—	—	—	—	—	

See Table 1 footnotes on next page.

TABLE 1. (Continued) Provisional cases of infrequently reported notifiable diseases (<1,000 cases reported during the preceding year) — United States, week ending September 27, 2008 (39th week)*

—: No reported cases. N: Not notifiable. Cum: Cumulative year-to-date counts.
 * Incidence data for reporting year 2008 are provisional, whereas data for 2003, 2004, 2005, 2006, and 2007 are finalized.
 † Calculated by summing the incidence counts for the current week, the 2 weeks preceding the current week, and the 2 weeks following the current week, for a total of 5 preceding years. Additional information is available at <http://www.cdc.gov/epo/dphsi/phs/files/5yearweeklyaverage.pdf>.
 § Not notifiable in all states. Data from states where the condition is not notifiable are excluded from this table, except in 2007 and 2008 for the domestic arboviral diseases and influenza-associated pediatric mortality, and in 2003 for SARS-CoV. Reporting exceptions are available at <http://www.cdc.gov/epo/dphsi/phs/infdis.htm>.
 ¶ Includes both neuroinvasive and nonneuroinvasive. Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Zoonotic, Vector-Borne, and Enteric Diseases (ArboNET Surveillance). Data for West Nile virus are available in Table II.
 ** The names of the reporting categories changed in 2008 as a result of revisions to the case definitions. Cases reported prior to 2008 were reported in the categories: Ehrlichiosis, human monocytic (analogous to *E. chaffeensis*); Ehrlichiosis, human granulocytic (analogous to *Anaplasma phagocytophilum*), and Ehrlichiosis, unspecified, or other agent (which included cases unable to be clearly placed in other categories, as well as possible cases of *E. ewingii*).
 †† Data for *H. influenzae* (all ages, all serotypes) are available in Table II.
 §§ Updated monthly from reports to the Division of HIV/AIDS Prevention, National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention. Implementation of HIV reporting influences the number of cases reported. Updates of pediatric HIV data have been temporarily suspended until upgrading of the national HIV/AIDS surveillance data management system is completed. Data for HIV/AIDS, when available, are displayed in Table IV, which appears quarterly.
 ¶¶ Updated weekly from reports to the Influenza Division, National Center for Immunization and Respiratory Diseases. Eighty-six cases occurring during the 2007–08 influenza season have been reported.
 *** No measles cases were reported for the current week.
 ††† Data for meningococcal disease (all serogroups) are available in Table II.
 §§§ In 2008, Q fever acute and chronic reporting categories were recognized as a result of revisions to the Q fever case definition. Prior to that time, case counts were not differentiated with respect to acute and chronic Q fever cases.
 ¶¶¶ No rubella cases were reported for the current week.
 **** Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Zoonotic, Vector-Borne, and Enteric Diseases.

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals September 27, 2008, with historical data



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

Notifiable Disease Data Team and 122 Cities Mortality Data Team
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TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 27, 2008, and September 29, 2007 (39th week)*

Reporting area	Lyme Disease					Malaria					Meningococcal disease, invasive† All serotypes				
	Current week	Previous 52 weeks		Cum 2008	Cum 2007	Current week	Previous 52 weeks		Cum 2008	Cum 2007	Current week	Previous 52 weeks		Cum 2008	Cum 2007
		Med	Max				Med	Max				Med	Max		
United States	403	382	1,375	18,140	21,640	13	22	136	726	951	12	19	53	818	828
New England	47	55	240	2,810	6,824	—	1	35	32	43	—	0	3	20	36
Connecticut	—	0	45	—	2,742	—	0	27	11	1	—	0	1	1	6
Maine§	44	2	73	468	307	—	0	1	—	6	—	0	1	4	5
Massachusetts	—	15	114	1,039	2,709	—	0	2	14	25	—	0	3	15	18
New Hampshire	—	10	124	1,036	791	—	0	1	3	8	—	0	0	—	3
Rhode Island§	—	0	30	—	161	—	0	8	—	—	—	0	1	—	1
Vermont§	3	2	38	267	114	—	0	1	4	3	—	0	1	—	3
Mid. Atlantic	255	170	968	11,442	8,852	1	5	13	172	295	2	2	6	96	105
New Jersey	—	36	182	2,212	2,623	—	0	2	—	59	—	0	2	10	14
New York (Upstate)	205	56	453	3,832	2,540	1	1	8	28	50	—	0	3	25	29
New York City	1	1	13	24	342	—	3	8	116	151	2	0	2	22	19
Pennsylvania	49	56	491	5,374	3,347	—	1	3	28	35	—	1	5	39	43
E.N. Central	7	10	71	630	1,910	—	2	7	90	101	1	3	9	128	126
Illinois	—	0	9	61	140	—	1	6	37	47	—	1	4	39	50
Indiana	—	0	8	31	42	—	0	2	5	8	—	0	4	22	20
Michigan	2	0	12	71	49	—	0	2	12	13	1	0	3	25	20
Ohio	—	0	4	29	26	—	0	3	24	19	—	1	4	32	29
Wisconsin	5	7	58	438	1,653	—	0	3	12	14	—	0	2	10	7
W.N. Central	1	5	740	745	339	1	1	9	48	28	1	2	8	75	49
Iowa	—	1	8	81	108	—	0	1	5	3	—	0	3	16	11
Kansas	—	0	1	2	8	1	0	1	6	2	—	0	1	3	4
Minnesota	—	1	731	628	206	—	0	8	21	11	—	0	7	19	14
Missouri	—	0	3	20	9	—	0	4	8	5	—	0	3	23	13
Nebraska§	1	0	2	10	5	—	0	2	8	6	1	0	2	11	2
North Dakota	—	0	9	1	3	—	0	2	—	—	—	0	1	1	2
South Dakota	—	0	1	3	—	—	0	0	—	1	—	0	1	2	3
S. Atlantic	88	54	172	2,185	3,508	4	4	13	171	202	4	3	10	126	136
Delaware	3	12	37	612	594	—	0	1	2	4	1	0	1	2	1
District of Columbia	4	2	11	126	102	—	0	1	1	2	—	0	0	—	—
Florida	7	1	8	70	21	3	1	4	41	46	—	1	3	46	53
Georgia	1	0	3	18	8	—	1	5	45	35	—	0	2	14	19
Maryland§	38	18	136	711	1,985	—	0	3	15	51	1	0	4	12	19
North Carolina	2	0	8	27	39	—	0	7	23	17	1	0	4	12	15
South Carolina§	2	0	4	18	24	—	0	2	9	5	—	0	3	19	13
Virginia§	31	12	68	569	678	1	1	7	35	41	1	0	2	18	14
West Virginia	—	0	9	34	57	—	0	0	—	1	—	0	1	3	2
E.S. Central	—	1	5	38	44	—	0	3	13	28	—	1	6	39	41
Alabama§	—	0	3	10	10	—	0	1	3	5	—	0	2	5	8
Kentucky	—	0	1	2	4	—	0	1	4	7	—	0	2	7	9
Mississippi	—	0	1	1	1	—	0	1	1	2	—	0	2	9	10
Tennessee§	—	0	3	25	29	—	0	2	5	14	—	0	3	18	14
W.S. Central	—	2	11	65	60	2	1	64	57	72	—	2	13	87	83
Arkansas§	—	0	1	2	1	—	0	1	—	—	—	0	2	7	9
Louisiana	—	0	1	1	2	—	0	1	2	14	—	0	3	19	24
Oklahoma	—	0	1	—	—	—	0	4	2	5	—	0	5	12	15
Texas§	—	2	10	62	57	2	1	60	53	53	—	1	7	49	35
Mountain	—	1	5	37	37	1	1	3	24	52	1	1	4	43	55
Arizona	—	0	1	5	2	—	0	2	11	11	1	0	2	7	11
Colorado	—	0	1	5	—	1	0	2	4	19	—	0	1	10	20
Idaho§	—	0	2	8	7	—	0	1	1	2	—	0	2	3	4
Montana§	—	0	2	4	4	—	0	0	—	3	—	0	1	4	1
Nevada§	—	0	2	9	10	—	0	3	4	2	—	0	2	6	4
New Mexico§	—	0	2	4	5	—	0	1	2	4	—	0	1	7	2
Utah	—	0	1	—	6	—	0	1	2	11	—	0	2	4	11
Wyoming§	—	0	1	2	3	—	0	0	—	—	—	0	1	2	2
Pacific	5	4	10	188	66	4	3	9	119	130	3	4	17	204	197
Alaska	—	0	2	5	5	—	0	2	4	2	—	0	2	3	1
California	3	3	8	137	56	3	2	8	88	91	2	3	17	145	145
Hawaii	N	0	0	N	N	—	0	1	2	2	—	0	2	4	8
Oregon§	1	0	5	37	4	—	0	2	4	13	—	1	3	28	25
Washington	1	0	7	9	1	1	0	3	21	22	1	0	5	24	18
American Samoa	N	0	0	N	N	—	0	0	—	—	—	0	0	—	—
C.N.M.I.	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—
Guam	—	0	0	—	—	—	0	1	1	1	—	0	0	—	—
Puerto Rico	N	0	0	N	N	—	0	1	1	3	—	0	1	3	6
U.S. Virgin Islands	N	0	0	N	N	—	0	0	—	—	—	0	0	—	—

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

U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.

* Incidence data for reporting year 2008 are provisional.

† Data for meningococcal disease, invasive caused by serogroups A, C, Y, & W-135; serogroup B; other serogroup; and unknown serogroup are available in Table I.

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

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 27, 2008, and September 29, 2007 (39th week)*

Reporting area	Streptococcal diseases, invasive, group A					<i>Streptococcus pneumoniae</i> , invasive disease, nondrug resistant† Age <5 years				
	Current week	Previous 52 weeks		Cum 2008	Cum 2007	Current week	Previous 52 weeks		Cum 2008	Cum 2007
		Med	Max				Med	Max		
United States	57	94	259	4,030	4,178	19	36	166	1,138	1,270
New England	—	6	31	299	322	—	2	14	55	96
Connecticut	—	0	26	90	95	—	0	11	—	12
Maine§	—	0	3	22	22	—	0	1	1	2
Massachusetts	—	3	8	138	161	—	1	5	39	64
New Hampshire	—	0	2	20	24	—	0	1	7	8
Rhode Island§	—	0	9	17	5	—	0	2	7	8
Vermont§	—	0	2	12	15	—	0	1	1	2
Mid. Atlantic	6	18	43	833	781	2	4	19	142	214
New Jersey	—	3	11	133	140	—	1	6	28	43
New York (Upstate)	4	6	17	276	242	2	2	14	73	75
New York City	—	3	10	150	185	—	1	12	41	96
Pennsylvania	2	6	16	274	214	N	0	0	N	N
E.N. Central	9	19	42	782	811	4	6	23	207	225
Illinois	—	5	16	206	244	—	1	6	46	56
Indiana	1	2	11	112	98	—	0	14	29	14
Michigan	3	3	10	138	169	—	1	5	54	60
Ohio	5	5	14	224	192	3	1	5	46	48
Wisconsin	—	2	10	102	108	1	1	3	32	47
W.N. Central	1	5	39	308	276	1	2	16	102	67
Iowa	—	0	0	—	—	—	0	0	—	—
Kansas	1	0	5	34	28	—	0	3	14	—
Minnesota	—	0	35	144	131	—	0	13	41	38
Missouri	—	1	10	70	74	—	1	2	28	18
Nebraska§	—	0	3	31	21	—	0	3	7	10
North Dakota	—	0	5	10	14	—	0	2	5	1
South Dakota	—	0	2	19	8	1	0	1	7	—
S. Atlantic	25	18	34	754	1,004	4	6	13	170	230
Delaware	—	0	2	6	9	—	0	0	—	—
District of Columbia	—	0	4	23	16	—	0	1	1	2
Florida	3	5	11	202	240	2	1	4	47	49
Georgia	12	4	13	188	193	—	1	5	49	51
Maryland§	2	1	6	27	171	—	0	4	5	50
North Carolina	7	2	10	117	139	N	0	0	N	N
South Carolina§	—	1	5	54	87	2	1	4	38	36
Virginia§	1	3	12	110	127	—	0	6	25	35
West Virginia	—	0	3	27	22	—	0	1	5	7
E.S. Central	5	4	9	142	171	1	2	11	71	75
Alabama§	N	0	0	N	N	N	0	0	N	N
Kentucky	1	1	3	33	32	N	0	0	N	N
Mississippi	N	0	0	N	N	—	0	3	16	5
Tennessee§	4	3	7	109	139	1	1	9	55	70
W.S. Central	4	8	85	362	251	4	5	66	198	177
Arkansas§	1	0	2	5	17	—	0	2	5	11
Louisiana	—	0	2	11	14	—	0	2	10	30
Oklahoma	1	2	19	92	58	1	1	7	52	37
Texas§	2	6	65	254	162	3	3	58	131	99
Mountain	4	11	22	433	452	3	5	12	180	173
Arizona	1	3	9	156	173	2	2	8	91	86
Colorado	3	2	8	122	113	1	1	4	51	35
Idaho§	—	0	2	11	15	—	0	1	3	2
Montana§	N	0	0	N	N	—	0	1	4	1
Nevada§	—	0	2	8	2	N	0	0	N	N
New Mexico§	—	2	8	84	76	—	0	3	15	28
Utah	—	1	5	46	68	—	0	3	15	21
Wyoming§	—	0	2	6	5	—	0	1	1	—
Pacific	3	3	10	117	110	—	0	2	13	13
Alaska	1	0	4	31	20	N	0	0	N	N
California	—	0	0	—	—	N	0	0	N	N
Hawaii	2	2	10	86	90	—	0	2	13	13
Oregon§	N	0	0	N	N	N	0	0	N	N
Washington	N	0	0	N	N	N	0	0	N	N
American Samoa	—	0	12	30	4	N	0	0	N	N
C.N.M.I.	—	—	—	—	—	—	—	—	—	—
Guam	—	0	1	—	13	—	0	0	—	—
Puerto Rico	N	0	0	N	N	N	0	0	N	N
U.S. Virgin Islands	—	0	0	—	—	N	0	0	N	N

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

U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.

* Incidence data for reporting year 2008 are provisional.

† Includes cases of invasive pneumococcal disease, in children aged <5 years, caused by *S. pneumoniae*, which is susceptible or for which susceptibility testing is not available (NNDSS event code 11717).

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

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 27, 2008, and September 29, 2007 (39th week)*

Reporting area	West Nile virus disease†														
	Varicella (chickenpox)					Neuroinvasive					Nonneuroinvasive§				
	Current week	Previous 52 weeks		Cum 2008	Cum 2007	Current week	Previous 52 weeks		Cum 2008	Cum 2007	Current week	Previous 52 weeks		Cum 2008	Cum 2007
	Med	Max				Med	Max				Med	Max			
United States	268	658	1,660	19,987	29,430	3	1	73	421	1,120	1	3	73	505	2,298
New England	14	13	68	412	1,866	—	0	2	3	4	—	0	1	2	6
Connecticut	—	0	38	—	1,081	—	0	2	3	1	—	0	1	2	2
Maine¶	—	0	26	—	238	—	0	0	—	—	—	0	0	—	—
Massachusetts	—	0	1	1	—	—	0	2	—	3	—	0	0	—	3
New Hampshire	4	6	18	197	263	—	0	0	—	—	—	0	0	—	—
Rhode Island¶	—	0	0	—	—	—	0	0	—	—	—	0	0	—	1
Vermont¶	10	6	17	214	284	—	0	0	—	—	—	0	0	—	—
Mid. Atlantic	51	56	117	1,715	3,692	—	0	6	27	19	—	0	3	9	8
New Jersey	N	0	0	N	N	—	0	1	2	1	—	0	1	2	—
New York (Upstate)	N	0	0	N	N	—	0	4	12	3	—	0	1	3	1
New York City	N	0	0	N	N	—	0	2	8	11	—	0	3	4	2
Pennsylvania	51	56	117	1,715	3,692	—	0	2	5	4	—	0	0	—	5
E.N. Central	67	163	378	4,789	8,285	—	0	11	22	99	—	0	4	12	57
Illinois	—	13	63	716	850	—	0	4	3	54	—	0	4	7	32
Indiana	—	0	222	—	—	—	0	2	2	13	—	0	0	—	10
Michigan	21	64	154	2,005	3,076	—	0	2	5	16	—	0	1	—	—
Ohio	45	55	128	1,719	3,520	—	0	3	10	11	—	0	2	2	9
Wisconsin	1	7	38	349	839	—	0	2	2	5	—	0	1	3	6
W.N. Central	6	25	145	868	1,205	—	0	6	30	238	—	0	20	120	722
Iowa	N	0	0	N	N	—	0	2	4	11	—	0	1	4	15
Kansas	6	5	36	286	447	—	0	1	4	12	—	0	3	12	26
Minnesota	—	0	0	—	—	—	0	2	3	43	—	0	6	18	56
Missouri	—	12	51	514	690	—	0	3	4	55	—	0	1	4	14
Nebraska¶	N	0	0	N	N	—	0	1	2	20	—	0	5	20	137
North Dakota	—	0	140	48	—	—	0	2	2	49	—	0	10	38	315
South Dakota	—	0	5	20	68	—	0	5	11	48	—	0	6	24	159
S. Atlantic	55	94	167	3,390	3,975	—	0	3	9	40	—	0	3	7	37
Delaware	—	1	6	39	36	—	0	0	—	1	—	0	1	1	—
District of Columbia	—	0	3	18	26	—	0	0	—	—	—	0	0	—	—
Florida	24	28	87	1,269	940	—	0	2	2	3	—	0	0	—	—
Georgia	N	0	0	N	N	—	0	1	1	22	—	0	2	1	25
Maryland¶	N	0	0	N	N	—	0	2	5	5	—	0	2	4	4
North Carolina	N	0	0	N	N	—	0	0	—	4	—	0	1	—	4
South Carolina¶	23	17	66	668	786	—	0	1	—	2	—	0	0	—	2
Virginia¶	—	21	81	847	1,317	—	0	0	—	3	—	0	1	1	2
West Virginia	8	15	66	549	870	—	0	1	1	—	—	0	0	—	—
E.S. Central	5	18	101	911	402	—	0	10	46	65	—	0	10	71	82
Alabama¶	5	18	101	901	400	—	0	5	11	15	—	0	2	4	5
Kentucky	N	0	0	N	N	—	0	1	—	3	—	0	0	—	—
Mississippi	—	0	2	10	2	—	0	6	30	43	—	0	9	61	73
Tennessee¶	N	0	0	N	N	—	0	1	5	4	—	0	2	6	4
W.S. Central	46	182	886	6,395	7,961	—	0	14	47	228	—	0	10	43	128
Arkansas¶	1	11	38	469	597	—	0	2	8	12	—	0	1	—	6
Louisiana	—	1	10	60	99	—	0	3	6	21	—	0	6	20	10
Oklahoma	N	0	0	N	N	—	0	3	3	56	—	0	3	5	42
Texas¶	45	166	852	5,866	7,265	—	0	10	30	139	—	0	6	18	70
Mountain	21	40	105	1,443	1,989	—	0	12	61	273	—	0	21	141	1,022
Arizona	—	0	0	—	—	—	0	7	33	41	—	0	10	12	38
Colorado	21	13	43	651	812	—	0	4	13	97	—	0	12	64	474
Idaho¶	N	0	0	N	N	—	0	1	2	11	—	0	7	30	116
Montana¶	—	5	27	223	301	—	0	1	—	36	—	0	2	5	165
Nevada¶	N	0	0	N	N	—	0	2	8	1	—	0	3	7	10
New Mexico¶	—	4	22	165	309	—	0	2	4	37	—	0	1	1	20
Utah	—	10	55	394	543	—	0	1	1	27	—	0	3	15	41
Wyoming¶	—	0	9	10	24	—	0	0	—	23	—	0	2	7	158
Pacific	3	1	7	64	55	3	0	31	176	154	1	0	15	100	236
Alaska	2	1	5	50	29	—	0	0	—	—	—	0	0	—	—
California	—	0	0	—	—	3	0	31	176	147	1	0	15	96	218
Hawaii	1	0	6	14	26	—	0	0	—	—	—	0	0	—	—
Oregon¶	N	0	0	N	N	—	0	0	—	7	—	0	2	4	18
Washington	N	0	0	N	N	—	0	0	—	—	—	0	0	—	—
American Samoa	N	0	0	N	N	—	0	0	—	—	—	0	0	—	—
C.N.M.I.	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—
Guam	—	2	17	55	212	—	0	0	—	—	—	0	0	—	—
Puerto Rico	—	9	20	334	586	—	0	0	—	—	—	0	0	—	—
U.S. Virgin Islands	—	0	0	—	—	—	0	0	—	—	—	0	0	—	—

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

U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.

* Incidence data for reporting year 2008 are provisional.

† Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Zoonotic, Vector-Borne, and Enteric Diseases (ArboNET Surveillance). Data for California serogroup, eastern equine, Powassan, St. Louis, and western equine diseases are available in Table I.

§ Not notifiable in all states. Data from states where the condition is not notifiable are excluded from this table, except in 2007 for the domestic arboviral diseases and influenza-associated pediatric mortality, and in 2003 for SARS-CoV. Reporting exceptions are available at <http://www.cdc.gov/epo/dphsi/phs/infdis.htm>.

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

TABLE III. Deaths in 122 U.S. cities,* week ending September 27, 2008 (39th week)

Reporting area	All causes, by age (years)						P&I† Total	Reporting area	All causes, by age (years)						P&I† Total
	All Ages	≥65	45-64	25-44	1-24	<1			All Ages	≥65	45-64	25-44	1-24	<1	
New England	439	299	93	26	9	7	40	S. Atlantic	1,204	718	315	89	47	35	67
Boston, MA	124	72	31	10	3	3	13	Atlanta, GA	128	77	31	13	2	5	2
Bridgeport, CT	34	21	10	3	—	—	4	Baltimore, MD	187	94	64	12	10	7	21
Cambridge, MA	14	12	2	—	—	—	3	Charlotte, NC	123	81	32	8	1	1	8
Fall River, MA	23	18	2	2	1	—	1	Jacksonville, FL	169	102	45	12	6	4	7
Hartford, CT	43	31	8	3	1	—	3	Miami, FL	84	57	16	6	3	2	7
Lowell, MA	21	16	4	1	—	—	2	Norfolk, VA	47	25	12	1	4	5	—
Lynn, MA	7	4	3	—	—	—	—	Richmond, VA	56	28	17	8	1	2	4
New Bedford, MA	21	15	5	—	—	1	1	Savannah, GA	57	40	12	2	3	—	1
New Haven, CT	U	U	U	U	U	U	U	St. Petersburg, FL	51	31	7	8	3	2	—
Providence, RI	46	38	7	1	—	—	1	Tampa, FL	186	117	44	14	7	4	15
Somerville, MA	1	1	—	—	—	—	—	Washington, D.C.	99	55	32	4	5	3	1
Springfield, MA	28	16	9	1	1	1	5	Wilmington, DE	17	11	3	1	2	—	1
Waterbury, CT	24	18	3	3	—	—	2	E.S. Central	870	553	206	58	19	34	43
Worcester, MA	53	37	9	2	3	2	5	Birmingham, AL	188	111	51	10	2	14	10
Mid. Atlantic	1,884	1,293	399	121	34	37	82	Chattanooga, TN	87	68	13	4	—	2	2
Albany, NY	50	36	6	3	2	3	1	Knoxville, TN	68	43	15	6	2	2	4
Allentown, PA	31	28	1	2	—	—	3	Lexington, KY	66	47	12	4	1	2	3
Buffalo, NY	78	46	26	4	1	1	5	Memphis, TN	165	94	40	16	11	4	12
Camden, NJ	23	16	3	2	—	2	—	Mobile, AL	104	63	34	3	2	2	4
Elizabeth, NJ	16	10	3	2	—	1	2	Montgomery, AL	56	42	6	6	—	2	2
Erie, PA	26	19	4	3	—	—	1	Nashville, TN	136	85	35	9	1	6	6
Jersey City, NJ	U	U	U	U	U	U	U	W.S. Central	1,511	929	383	125	41	33	76
New York City, NY	958	673	206	50	12	17	27	Austin, TX	100	64	24	8	—	4	6
Newark, NJ	40	18	13	4	1	4	3	Baton Rouge, LA	82	50	19	8	5	—	—
Paterson, NJ	16	8	5	2	1	—	2	Corpus Christi, TX	62	31	20	6	4	1	8
Philadelphia, PA	252	151	58	28	9	6	12	Dallas, TX	183	100	53	22	4	4	7
Pittsburgh, PA§	25	14	7	3	1	—	1	El Paso, TX	78	58	16	3	—	1	2
Reading, PA	20	15	4	1	—	—	—	Fort Worth, TX	128	70	38	11	3	6	4
Rochester, NY	135	99	24	8	2	2	8	Houston, TX	384	228	101	31	13	11	21
Schenectady, NY	18	14	3	1	—	—	2	Little Rock, AR	75	52	18	3	2	—	4
Scranton, PA	33	25	5	1	2	—	3	New Orleans, LA¶	U	U	U	U	U	U	U
Syracuse, NY	102	82	14	3	2	1	9	San Antonio, TX	224	137	56	19	6	6	11
Trenton, NJ	36	24	8	3	1	—	3	Shreveport, LA	86	56	19	9	2	—	12
Utica, NY	8	4	4	—	—	—	—	Tulsa, OK	109	83	19	5	2	—	1
Yonkers, NY	17	11	5	1	—	—	—	Mountain	873	551	215	68	25	14	54
E.N. Central	1,677	1,101	371	115	50	40	94	Albuquerque, NM	109	73	24	10	2	—	5
Akron, OH	54	35	10	3	2	4	1	Boise, ID	28	21	5	—	2	—	—
Canton, OH	34	28	4	1	—	1	5	Colorado Springs, CO	52	32	12	6	1	1	4
Chicago, IL	319	182	82	36	15	4	19	Denver, CO	95	50	34	7	2	2	5
Cincinnati, OH	66	39	18	4	2	3	5	Las Vegas, NV	217	143	54	11	6	3	13
Cleveland, OH	197	142	33	13	4	5	14	Ogden, UT	25	20	2	—	2	1	1
Columbus, OH	175	106	48	12	4	5	12	Phoenix, AZ	132	70	39	16	5	2	6
Dayton, OH	119	75	31	9	3	1	8	Pueblo, CO	29	19	7	3	—	—	2
Detroit, MI	122	68	31	12	5	6	6	Salt Lake City, UT	86	56	16	8	3	3	10
Evansville, IN	53	39	12	—	2	—	2	Tucson, AZ	100	67	22	7	2	2	8
Fort Wayne, IN	60	40	14	3	1	2	2	Pacific	1,545	1,050	330	91	45	29	113
Gary, IN	18	9	4	1	2	2	—	Berkeley, CA	14	9	2	2	—	1	—
Grand Rapids, MI	59	40	14	2	—	3	4	Fresno, CA	114	82	24	4	4	—	6
Indianapolis, IN	U	U	U	U	U	U	U	Glendale, CA	26	21	5	—	—	—	2
Lansing, MI	50	35	12	1	1	1	2	Honolulu, HI	77	63	8	3	2	1	11
Milwaukee, WI	70	42	20	7	1	—	1	Long Beach, CA	64	45	12	2	2	3	6
Peoria, IL	40	29	8	1	1	1	3	Los Angeles, CA	212	130	50	21	5	6	21
Rockford, IL	64	49	10	4	1	—	3	Pasadena, CA	19	14	3	1	—	1	1
South Bend, IN	53	40	7	2	4	—	3	Portland, OR	103	67	21	12	2	1	2
Toledo, OH	80	64	8	4	2	2	4	Sacramento, CA	157	107	39	4	4	3	12
Youngstown, OH	44	39	5	—	—	—	—	San Diego, CA	192	131	39	13	5	4	11
W.N. Central	676	434	157	44	24	17	48	San Francisco, CA	94	60	21	6	4	3	8
Des Moines, IA	87	59	24	2	2	—	4	San Jose, CA	150	110	31	3	5	1	13
Duluth, MN	26	19	6	—	—	1	1	Santa Cruz, CA	31	20	8	1	2	—	1
Kansas City, KS	21	14	5	—	2	—	1	Seattle, WA	121	77	32	8	3	1	5
Kansas City, MO	85	49	19	6	6	5	7	Spokane, WA	69	42	15	6	4	2	8
Lincoln, NE	37	29	4	3	—	1	2	Tacoma, WA	102	72	20	5	3	2	6
Minneapolis, MN	84	48	21	8	2	5	3	Total**	10,679	6,928	2,469	737	294	246	617
Omaha, NE	100	69	23	6	1	1	18								
St. Louis, MO	97	53	28	9	5	2	2								
St. Paul, MN	51	35	7	6	2	1	6								
Wichita, KS	88	59	20	4	4	1	4								

U: Unavailable. —:No reported cases.

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

† Pneumonia and influenza.

§ Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks.

¶ Because of Hurricane Katrina, weekly reporting of deaths has been temporarily disrupted.

** Total includes unknown ages.

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