

Weekly

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National Action Week for the Bone and Joint Decade — October 12–20, 2006

National Action Week for the Bone and Joint Decade, a global, multidisciplinary initiative promoting the care of persons with bone and joint disorders, is being observed October 12–20. This initiative focuses on improving quality of life and advancing the understanding and treatment of musculoskeletal conditions through research, prevention, and education. CDC, the National Institutes of Health, the World Health Organization, and the United Nations are among the governmental and nongovernmental organizations supporting this initiative. In 2002, the United States officially proclaimed the years 2002–2011 as the National Bone and Joint Decade.

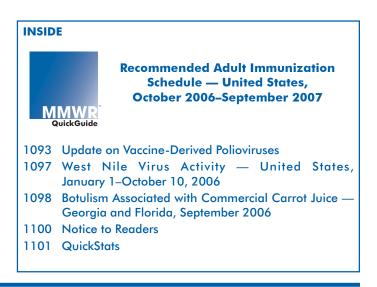
Bone and joint disorders are the leading causes of disability in the United States (1) and impose substantial burdens on the health-care system and society (2). The United States Bone and Joint Decade organization is committed to raising awareness of the growing burden of musculoskeletal conditions and promoting their prevention, advancing research, and improving diagnosis and treatment. This week, two education programs will be inaugurated by the U.S. group: Straighten Up America and Joint Health and Arthritis. Additional information regarding the United States Bone and Joint Decade is available at http://www.usbjd.org and regarding the national action week is available at http://www.usbjd.org/rd/?naw. Information about activities worldwide is available at http://www.boneandjointdecade.org.

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Prevalence of Doctor-Diagnosed Arthritis and Arthritis-Attributable Activity Limitation — United States, 2003–2005

Arthritis is highly prevalent among U.S. adults, the leading cause of disability (1), and associated with substantial activity limitation, work disability, reduced quality of life, and high health-care costs (2-4). As the population ages, arthritis is expected to affect an estimated 67 million adults in the United States by 2030 (5). This report updates estimates of the national prevalence of doctor-diagnosed arthritis and arthritisattributable activity limitation in the adult U.S. population, using data from the National Health Interview Survey (NHIS) for 2003-2005. The findings indicated that an estimated 21.6% of the adult U.S. population (46.4 million persons) had doctor-diagnosed arthritis, and 8.3% (17.4 million) had arthritis-attributable activity limitations. Public and private health agencies should promote measures to increase the availability of evidence-based arthritis prevention and management interventions.



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NHIS is an annual, household-based survey of a representative sample of the U.S. civilian, noninstitutionalized population, using in-person interviews. This study used the sample adult core component of the NHIS survey, which collects information on adults aged ≥ 18 years residing in selected households. In 2003, 2004, and 2005, the sample sizes were 30,852, 31,326, and 31,428, respectively, for the adult core component, and the final response rates were 74.2%, 72.5%, and 69.0%, respectively. Respondents were defined as having doctor-diagnosed arthritis if they answered "yes" to the question, "Have you ever been told by a doctor or other health professional that you have some form of arthritis, rheumatoid arthritis, gout, lupus, or fibromyalgia?" Those who answered "yes" were asked, "Are you limited in any way in any of your usual activities because of arthritis or joint symptoms?" Persons responding "yes" to both questions were defined as having an arthritis-attributable activity limitation.

For this study, prevalence estimates are presented overall and by sex, age group, race/ethnicity, education level, body mass index (BMI)* category, and physical activity level. Physical activity level of respondents was determined from six questions that asked about frequency and duration of participation in leisure-time activities of moderate and vigorous intensity; those reporting no participation in such activities were classified as inactive, and all others as active. Estimates were calculated by using combined data from 2003-2005 and applying an annual average weighting; 95% confidence intervals (CIs) were calculated using sample design factors and statistical software to account for the multistage probability sample. To facilitate comparisons between demographic subgroups, estimates were age adjusted to the standard 2000 U.S. population (6). All differences noted in this report are statistically significant (p<0.05) with nonoverlapping 95% CIs.

In unadjusted analyses for 2003–2005 (Table), the prevalence of doctor-diagnosed arthritis among adults was estimated at 21.6%, or 46.4 million persons. Prevalence was higher among women (25.4%) compared with men (17.6%), older age groups (50% for persons aged \geq 65 years and 29.3% for persons aged 45–64 years) compared with younger age groups (7.9% for persons aged 18–44 years), and non-Hispanic whites (24.3%) compared with non-Hispanic blacks (19.2%) and Hispanics (11.4%). Prevalence also was higher among those who were obese (31.6%) or overweight (21.7%) compared with those who were normal weight or underweight (16.3%)

^{*} BMI was calculated using self-reported weight and height as follows: weight (kg) / height (m²). Categories were defined as follows: underweight/normal weight, ≤24.9; overweight, 25.0–29.9; and obese, ≥30.0.

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TABLE. Unadjusted and age-adjusted* estimates of the prevalence of doctor-diagnosed arthritis and arthritis-attributable activity limitations[†] among adults aged ≥18 years, by selected characteristics — National Health Interview Survey, United States, 2003–2005

			Adı		Proportion with arthritis-attributable								
	Do	ctor-diagn					ole activity	/limitation	activity limitation among those				
	((46.4 millio	n persor	ıs)		(17.4 milli	on persoi	with doctor-diagnosed arthritis					
	Una	djusted	Age	adjusted	Unadjusted		Age	adjusted	Unadjusted		Age adjusted		
Characteristic	% (95% Cl§)	%	(95% CI)	%	(95% CI)	%	(95% CI)	%	(95% CI)	%	(95% CI)	
Sex													
Men	17.6	(<u>+</u> 0.5)	18.1	(<u>+</u> 0.5)	6.4	(<u>+</u> 0.3)	6.6	(<u>+</u> 0.4)	36.6	(<u>+</u> 1.4)	35.2	(<u>+</u> 1.6)	
Women	25.4	(<u>+</u> 0.6)	24.4	(<u>+</u> 0.5)	10.0	(<u>+</u> 0.3)	9.7	(<u>+</u> 0.3)	40.1	(<u>+</u> 1.0)	37.5	(<u>+</u> 1.3)	
Age (yrs)													
18–44	7.9	(<u>+</u> 0.3)		_	2.7	(<u>+</u> 0.2)			33.9	(<u>+</u> 1.8)			
45–64	29.3	(<u>+</u> 0.7)		_	11.3	(<u>+</u> 0.5)			38.6	(<u>+</u> 1.4)			
<u>≥</u> 65	50.0	(<u>+</u> 0.9)		_	20.5	(<u>+</u> 0.7)			41.5	(<u>+</u> 1.3)			
Race/Ethnicity													
White, non-Hispanic	24.3	(<u>+</u> 0.5)	22.6	(<u>+</u> 0.4)	8.9	(<u>+</u> 0.3)	8.4	(<u>+</u> 0.3)	37.4	(<u>+</u> 0.9)	35.0	(<u>+</u> 1.2)	
Black, non-Hispanic	19.2	(<u>+</u> 0.9)	21.4	(<u>+</u> 0.9)	8.8	(<u>+</u> 0.6)	10.0	(<u>+</u> 0.6)	45.7	(<u>+</u> 2.4)	43.4	(<u>+</u> 3.1)	
Hispanic	11.4	(<u>+</u> 0.6)	16.5	(<u>+</u> 0.8)	5.0	(<u>+</u> 0.4)	7.3	(<u>+</u> 0.6)	43.8	(<u>+</u> 2.7)	42.3	(<u>+</u> 3.2)	
Other non-Hispanic	14.7	(<u>+</u> 1.3)	17.3	(<u>+</u> 0.5)	5.7	(<u>+</u> 0.8)	7.0	(±1.0)	41.5	(<u>+</u> 4.6)	40.0	(<u>+</u> 5.3)	
Education													
Did not graduate from high school	27.0	(<u>+</u> 1.0)	23.2	(<u>+</u> 0.8)	13.6	(<u>+</u> 0.6)	11.7	(<u>+</u> 0.6)	50.6	(<u>+</u> 1.6)	49.3	(<u>+</u> 3.0)	
High school graduate or more	20.8	(<u>+</u> 0.4)	21.2	(<u>+</u> 0.4)	7.4	(<u>+</u> 0.2)	7.7	(<u>+</u> 0.2)	36.1	(<u>+</u> 0.9)	34.5	(<u>+</u> 1.1)	
Body mass index (BMI [¶])													
Underweight/Normal weight	16.3	(<u>+</u> 0.5)	17.4	(<u>+</u> 0.5)	5.5	(<u>+</u> 0.3)	5.9	(<u>+</u> 0.3)	34.3	(<u>+</u> 1.4)	32.4	(<u>+</u> 1.8)	
Overweight	21.7	(<u>+</u> 0.6)	20.5	(<u>+</u> 0.5)	7.5	(<u>+</u> 0.3)	7.1	(<u>+</u> 0.3)	35.0	(<u>+</u> 1.3)	33.4	(<u>+</u> 1.8)	
Obese	31.6	(<u>+</u> 0.8)	29.3	(<u>+</u> 0.7)	14.4	(<u>+</u> 0.6)	13.8	(<u>+</u> 0.6)	46.4	(<u>+</u> 1.6)	43.2	(<u>+</u> 1.9)	
Physical activity level													
Inactive	25.0	(<u>+</u> 0.6)	22.3	(<u>+</u> 0.5)	13.2	(<u>+</u> 0.5)	11.7	(<u>+</u> 0.4)	52.6	(<u>+</u> 1.3)	49.8	(<u>+</u> 2.1)	
Active	19.5	(<u>+</u> 0.5)	20.8	(<u>+</u> 0.5)	6.1	(<u>+</u> 0.3)	6.6	(<u>+</u> 0.3)	31.3	(<u>+</u> 1.0)	29.9	(<u>+</u> 1.2)	
Total	21.6	(± 0.4)	21.5	(<u>+</u> 0.4)	8.3	(<u>+</u> 0.2)	8.3	(<u>+</u> 0.2)	38.8	(<u>+</u> 0.8)	36.6	(<u>+</u> 1.0)	

* Adjusted to the projected 2000 population aged \geq 18 years by three age groups: 18–44 years, 45–64 years, and \geq 65 years.

Adjusted to the projected 2000 population aged 210 years by tince age groups, no 41 years, no 41 years, and 200 years and 200 population aged 210 years by tince age groups. To 41 years, no 41 years, and 200 years and 200 population aged 210 years by tince age groups. To 41 years, no 41 years, and 200 years and 200 population aged 210 years by tince age groups. To 41 years, no 41 years, and 200 years are provided at the professional that you have some form of arthritis, rheumatoid arthritis, gout, lupus, or fibromyalgia? Those who answered "yes" were asked, "Are you limited in any way in any of the professional that you have some form of arthritis, rheumatoid arthritis, gout, lupus, or fibromyalgia? Those who answered "yes" were asked, "Are you limited in any way in any of your usual activities because of arthritis or joint symptoms?" Persons responding "yes" to both questions were defined as having an arthritis-attributable s activity limitation.

Confidence interval

¹BMI = weight (kg) / height (m²). Underweight/normal weight, \leq 24.9; overweight, 25.0–29.9; and obese, \geq 30.0.

and among those who were physically inactive (25.0%) compared with those who were physically active (19.5%). After adjustment for age, all of these differences (except among age groups) were slightly attenuated but remained significant, with the exception of differences between non-Hispanics whites (22.6%) and non-Hispanic blacks (21.4%).

Unadjusted analyses for arthritis-attributable activity limitation among adults indicated an estimated overall prevalence of 8.3%, or 17.4 million persons, with differences among groups that were similar to those for doctor-diagnosed arthritis prevalence. The exception was a similar prevalence for non-Hispanic blacks (8.8%) and non-Hispanic whites (8.9%). Age-adjusted analyses identified differences among groups that were similar to the unadjusted figures, except that prevalence among non-Hispanic blacks (10.0%) significantly exceeded that for non-Hispanic whites (8.4%).

In unadjusted analyses of all adults reporting arthritis, 38.8% reported arthritis-attributable activity limitation (Table). Proportions were significantly higher among women (40.1%) compared with men (36.6%) and among non-Hispanic blacks (45.7%) and Hispanics (43.8%) compared with non-Hispanic whites (37.4%). Persons with arthritis and activity limitations also were more likely to have less than a high school education (50.6% versus 36.1%) or to be obese (46.4% versus 34.3%) underweight/normal weight) or physically inactive (52.6%) versus 31.3%). Age-adjusted analyses eliminated the significant difference between men and women, but did not otherwise change the results.

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Editorial Note: The findings in this report indicate that 21.6% (46.4 million) of U.S. adults reported doctor-diagnosed arthritis, and 8.3% (17.4 million) reported arthritisattributable activity limitation during 2003–2005. This represents an increase from 2002, when an estimated 20.8% (42.7 million) reported doctor-diagnosed arthritis and 7.8% (16.0 million) reported arthritis-attributable activity limitation (2). The increase in both the prevalence of doctor-diagnosed arthritis and arthritis-attributable activity limitation is consistent with future projections, largely based on the aging of the population (5). However, in 2003, the NHIS transitioned to a weighting structure based on the 2000 U.S. Census population; therefore, interpretation of this increased prevalence should be made with caution. Additional years of data are needed to determine whether these growth trends will be lasting.

Disparities exist with regard to arthritis and activity limitations. Women, older adults, persons with little education, or those who are obese, overweight, or physically inactive are more likely affected. In unadjusted analyses, doctor-diagnosed arthritis was less prevalent among non-Hispanic blacks and Hispanics than among non-Hisipanic whites; however, both groups reported greater proportions of persons with arthritisattributable activity limitation.

In contrast to previous estimates of arthritis prevalence based on 1 year of data, prevalences for a 3-year period were used to reduce the year-to-year fluctuation that can result from smaller sample sizes from a single year. This approach might provide more reliable estimates, especially for smaller groups such as certain racial/ethnic populations and older adults.

The findings in this report are subject to at least three limitations. First, doctor-diagnosed arthritis was self-reported and not confirmed by a health-care professional, although selfreport of arthritis has been determined valid for surveillance purposes (7). Second, the cross-sectional study design does permit determining the temporal sequence of arthritis onset and selected characteristics (e.g., obesity or physical inactivity). However, other studies have identified excess body weight as a risk factor for incident osteoarthritis, the most common type of arthritis, and physical activity has been determined to prevent or delay onset of functional limitation and disability among adults with osteoarthritis (8). Finally, certain factors that might contribute to differences in arthritis prevalence (e.g., history of joint injury or comorbid conditions such as cardiovascular disease, diabetes, or depression) were not analyzed.

Population-based national surveillance of arthritis prevalence and associated effects such as arthritis-attributable activity limitation are important to identify groups at greatest risk, target interventions, and measure progress toward achieving national health objectives (9). Currently, the CDC Arthritis Program is focusing on expanding the availability of evidence-based physical-activity and self-management interventions proven to reduce pain and improve function among adults with arthritis. Such interventions include those related to safe physical activity for persons with arthritis (e.g., Arthritis Foundation's Exercise Program, Arthritis Foundation's Aquatics Program, and EnhanceFitness) and self-management education (e.g., Arthritis Foundation's Self-Help Course and the Chronic Disease Self-Management Program). In addition, the CDC Arthritis Program is working with 35 state health department programs and various local chapters of the Arthritis Foundation to disseminate a health communications campaign designed to promote greater physical activity among adults with arthritis. The campaign, "Physical Activity. The Arthritis Pain Reliever," was developed to target an audience of low-income men and women aged \geq 45 years with arthritis. A similar campaign targeted to Spanish-speaking adults, "Buenos Dias Artritis," is being developed and tested.[†] Further research is needed to investigate possible underlying reasons for the differences among groups in arthritis prevalence and activity limitation and to develop more targeted solutions to improve the quality of life for all adults with arthritis, particularly among those most affected.

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[†]Additional information on arthritis programs is available at http://www.cdc.gov/ arthritis.

Update on Vaccine-Derived Polioviruses

In 1988, the World Health Assembly resolved to eradicate polio worldwide. The Global Polio Eradication Initiative (PEI) of the World Health Organization (WHO) has led to a decline in global polio incidence, from an estimated 350,000 cases in 1988 to fewer than 2,000 reported cases in 2005, and polio remains endemic to only four countries (Afghanistan, India, Nigeria, and Pakistan) (1). However, two additional obstacles to global eradication involve vaccine-derived polioviruses (VDPVs). Polio outbreaks continue to be associated with circulating vaccine-derived polioviruses (cVDPVs) in areas with low oral poliovirus vaccine (OPV) coverage. In addition, long-term excretion of neurovirulent immunodeficiency-associated vaccine-derived polioviruses (iVDPVs) can lead to poliovirus spread to contacts. Overcoming these obstacles is challenging. High rates of OPV coverage will prevent all poliovirus spread, including spread of VDPVs, but will not prevent establishment of prolonged VDPV infections in certain persons with B-cell immunodeficiencies (i.e., having defects in antibody production). Inevitable gaps in vaccination coverage will give rise to cVDPVs as long as OPV use continues. This report updates a previous report on VDPVs and describes the potential implications of VDPVs in the final stages of global polio eradication (2). The findings underscore the critical need to strengthen strategies to prevent emergence of VDPVs and to stop all OPV use once wild polioviruses (WPVs) are eradicated (2-5).

Biologic Properties of VDPVs

The critical biologic properties of VDPVs are their capacity to cause paralytic polio in humans and their potential or demonstrated capacity for sustained circulation. VDPVs have lost key attenuating mutations and resemble WPVs biologically (2). All known cVDPVs (except those from China) (Table 1), but no iVDPVs, are recombinants with nonstructural protein sequences derived from species C enteroviruses, a property associated with poliovirus circulation (2). Most VDPVs are antigenic variants of the Sabin strains, but antigenic evolution appears to be faster in iVDPVs than in cVDPVs. Unlike cVDPV isolates, iVDPV isolates commonly contain mixed VDPV populations. These biologic distinctions (and the differing conditions favoring iVDPV and cVDPV emergence) have helped in recognition of the likely origins of many ambiguous VDPVs (aVDPVs) (2).

Categories of VDPVs

VDPVs differ from the majority of vaccine-related isolates by having genetic properties consistent with prolonged replication or transmission. Because poliovirus genomes evolve at a rate of approximately 1% per year, vaccine-related isolates

TABLE 1. Outbreaks of circulating vaccine-derived polioviruses (cVDPVs) — worldwide, 1988–2006

Location	Years	Reported no. of polio cases	Serotype	No. of clinical isolates (% VP1 divergence from Sabin strain)	Recombination with species C enteroviruses*	Estimated duration of circulation	Routine vaccination coverage with 3 doses of oral polio vaccine (OPV)
Egypt [†]	1988–1993	30	2	30 (4.0–7.0)	Yes	10.0 yrs	Reported high
Haiti [§]	2000-2001	8	1	8 (1.9–2.6)	Yes	2.5 yrs	<30% nationwide
Dominican Republic [§]	2000-2001	13	1	13 (1.9–2.6)	Yes	>0.5 yr	<30% around most cases
Philippines	2001	3	1	4 (3.1–3.5)	Yes	2.5 yrs	OPV shortage previous 2 yrs
Madagascar [¶] **	2002	4	2	6 (2.5-3.0)	Yes	2.5 yrs	<50% nationwide
China ^{†† §§}	2004	2	1	4 (1.0–1.2)	No	1.0 yr	<50% around cases
Madagascar ^{§§}	2005	3	2	3 (1.1–1.8)	Yes	1.0 yr	<50% nationwide
Indonesia ^{§§} ¶¶	2005	46	1	46 (1.1–3.0)	Yes	2.0 yrs	<40% in Madura
Cambodia ^{§§}	2005–2006	3	3	3 (1.9–2.4)	Yes	>1.0 yr	<50% around cases

* All cVDPV isolates except those from China were vaccine/nonvaccine recombinants.

[†] Inferred retrospectively from sequence studies of stored isolates. Not investigated in the field.

§ Common outbreak. In 2000, cVDPV spread from Haiti to the Dominican Republic.

[¶] In 2001, an unrelated type 2 ambiguous VDPV isolate (1% VP1 divergence) was obtained from a patient with acute flaccid paralysis in a separate community in Madagascar.

** VDPVs were isolated from four polio patients (March–April 2002) and from two healthy children (from a stool survey of 316 healthy children conducted in June 2002 in the outbreak area).

^{††} Localized outbreak in Guizhou province.

§§ New cases reported since publication of previous report (2).

^{¶¶} Localized outbreak on Madura Island off coast of Java.

that differ from the corresponding OPV strain by more than 1% of nucleotide positions (usually determined by sequencing the genomic region encoding the major viral surface protein, VP1) are estimated to have replicated for at least 1 year after administration of an OPV dose, substantially longer than the normal period of vaccine virus replication of 4-6 weeks. Poliovirus isolates are divided into three categories, identified by the extent of VP1 nucleotide sequence divergence from the corresponding Sabin OPV strain: 1) OPV-like viruses (<1% divergent), 2) VDPVs (1%-15% divergent), and 3) WPVs (>15% divergent) (2). VDPVs are further divided into 1) iVDPVs isolated from persons with primary immunodeficiencies who have prolonged VDPV infections after exposure to OPV, 2) cVDPVs that emerge in communities with inadequate OPV coverage, and 3) aVDPVs, which are clinical isolates from persons with no known immunodeficiency and environmental isolates whose ultimate source has not been identified (2).

iVDPVs

A small proportion of immunodeficient persons exposed to OPV have excreted iVDPV over prolonged periods (>6 months). WHO maintains an iVDPV registry; since the introduction of OPV in 1961-1962, only 30 persons excreting iVDPVs have been identified. Persons with primary B-cell immunodeficiencies, but not persons with T-cell immunodeficiencies (e.g., from human immunodeficiency virus infection), are at risk for iVDPV infections (6). Approximately 70% of iVDPV infections have spontaneously ceased within 3 years of exposure to OPV, or the patients have died from complications of their immunodeficiency. Five persons excreted virus for 3-8 years, and in three persons, the duration of excretion exceeded 9 years (Table 2). Eighteen (60%) documented iVDPV infections were associated with type 2 poliovirus infection, eight (27%) with type 1, one (3%) with type 3, and three (9%) with mixed infections (Table 2, Figure). The first reports of iVDPVs came from high-income countries (e.g., the United States, countries of Western Europe, and Japan) but recent reports of iVDPVs include middleincome countries (Table 2). No iVDPVs have been reported from low-income countries, where survival rates for persons with B-cell immunodeficiencies are low (7). Exposure usually is from receipt of OPV, but three of the known iVDPV infections occurred in unimmunized persons (Table 2). Strategies for resolving iVDPV infections are needed, both because of the risk for paralytic disease to infected persons and the risk for transmission to the wider community. No antiviral drug that has been shown to resolve iVDPV infections is currently available. However, new antiviral drugs broadly effective against VDPVs are under development (8).

cVDPVs

VDPVs do not circulate when high vaccination coverage leads to high population immunity. However, low vaccination coverage increases the proportion of nonimmune persons in a population; this increases the potential for VDPVs to circulate. Under circumstances of low vaccination coverage, cVDPVs have produced several localized polio outbreaks. Eight independent outbreaks (i.e., two or more polio cases) in eight countries have been associated with cVDPVs (Table 1, Figure). The largest documented outbreak (46 polio cases) occurred on the Indonesian island of Madura. Genetic studies on stored isolates suggest that a type 2 cVDPV circulated endemically in Egypt for 10 years (approximately from 1983 to 1993) and probably caused more polio cases than were reported (2). Outbreaks of cVDPVs have been associated with all three poliovirus serotypes. Two independent type 2 cVDPV outbreaks occurred in Madagascar in 2002 and 2005 (2), possibly signaling a higher potential for the emergence of type 2 cVDPVs.

aVDPVs

aVDPVs are VDPV isolates that cannot be clearly assigned to either of the other two well-defined categories. They have been isolated from paralyzed persons with no evidence of additional paralyzed VDPV-infected persons among household or community contacts. Highly divergent (>12% VP1 nucleotide divergence) aVDPVs also have been isolated from sewage in Estonia, Israel, and Slovakia. The sewage isolates have similar genetic and antigenic properties as iVDPVs, but measures to identify the infected persons have been unsuccessful. In 1966, aVDPVs were found in Belarus after local suspension of OPV use; in 1999, they were found in Russia among children in orphanages (2). A growing number of aVDPVs having VP1 sequence divergence slightly above 1% have been found by the Global Polio Laboratory Network.

Limited person-to-person transmission for certain aVDPVs has occurred. In 2005, a type 3 aVDPV was isolated from one polio patient and seven nonparalyzed contacts in Madagascar. Similarly, a type 1 VDPV was isolated from one patient and seven contacts in Romania in 2002, a type 2 VDPV was isolated from one patient and two contacts in Laos in 2004 (2), a type 1 VDPV was isolated from an unimmunized severe combined immunodeficiency (SCID) patient and four community members in rural Minnesota in 2005 (9), and a type 1 VDPV was isolated from one patient and six contacts in Myanmar in 2006. Other aVDPVs with genetic properties resembling those of cVDPVs were found in Peru in 1983, in Pakistan in 2000, and in Nigeria in 2002 and 2006 (2). TABLE 2. Selected characteristics of persons excreting immunodeficiency-associated vaccine-derived polioviruses (iVDPVs) — worldwide, 1962–2006

Location	Year detected	Immunodeficiency	Paralysis	Serotype	Maximum % VP1 divergence from Sabin e strain	Estimated interval between last oral polio vaccine (OPV) dose and detection of iVDPV infection (yrs)*	Outcome	Estimated duration of virus excretion (yrs)
United Kingdom	1962	Hypogammaglobulinemia	No	1	Unknown	0†	Died	2.7
United Kingdom	1962	Hypogammaglobulinemia	No	3	2.3	0†	Died	1.8
Japan	1977	X-linked agammaglobulinemia	Yes	2	Unknown	1.5	Died	3.4
United States	1980	Agammaglobulinemia	Yes	2	Unknown	1	Died	1§
United States	1981	Common variable immunodeficiency	Yes	1	10.0	7.1	Died	7.6
United States	1986	X-linked agammaglobulinemia	Yes	2	2.0	0.4	Survived	0.4
United States [¶]	1986	Common variable immunodeficiency	Yes	1	5.4	4.7	Survived	9.6
	1992	_	_	2	11.8	_	_	
United Kingdom	1987	Common variable immunodeficiency	No	2	4.1	4	Survived	3.6
United States	1989	Agammaglobulinemia	Yes	1	1.1	0.3	NA**	Unknown
Germany	1990	Common variable immunodeficiency	Yes	1	8.3	4	Survived	9.5
United States	1990	Severe combined immunodeficiency	Yes	2	1.9	0.5	Died	0.8
United States	1991	Common variable immunodeficiency	Yes	2	1.4	0.4	Survived	0.6
Iran	1995	Antibody deficiency	Yes	2	2.2	Unimmunized	Died	1.5
United Kingdom	1995	Common variable immunodeficiency	No	2	12.9	15.7	Survived	20
United States	1995	Severe combined immunodeficiency	Yes	2	2.1	0.3	Died	3.7
Argentina	1998	X-linked agammaglobulinemia	Yes	1	2.8	Unimmunized	Survived	2
Germany	2000	Antibody deficiency	Yes	1	3.5	NA	Survived	1.5
Taiwan	2001	Common variable immunodeficiency	Yes	1	3.5	1.6	Survived	3
United Kingdom United Kingdom	2002 2002	Common variable immunodeficiency Immunodeficiency-centromeric	No	2	3.3	NA	Survived	3.3
		instability-facial abnormalities syndrom		2	2.5	NA	Survived	2.5
Kazakhstan	2002	Hypogammaglobulinemia	Yes	2	2.3	NA	Died	2
Kuwait	2002	Major histocompatibility complex class					.	
-		molecule deficiency	No	2	2.0	0.9	Died	0.4
Peru	2003	Agammaglobulinemia	Yes	2	1.2	0.6	Survived	<1.0
Thailand	2003	Hypogammaglobulinemia	Yes	2	2.2	0.3	NA	<0.5
China ^{††}	2005	X-linked agammaglobulinemia	Yes	2+3	2.7	0.6	NA	NA
Iran ^{††}	2005	Common variable immunodeficiency	Yes	2	1.4	0.7	Died	0.7
Morocco ^{††}	2005	Severe combined immunodeficiency	Yes	2	2.5	1	Died	1
Syria ^{††}	2005	Hypogammaglobulinemia	Yes	2	1.3	<0.1	Survived	<0.1
United States ^{††}	2005	Severe combined immunodeficiency	No	1	2.3	Unimmunized	Survived	<0.5 ^{§§}
Tunisia ^{††}	2006	Severe combined immunodeficiency	No	2	2.0	NA	Survived	0.1

* Several estimates are approximate because of no follow-up sampling, long sampling intervals, or uncertain date of associated OPV exposure. Because criteria for estimates varied in different studies, certain estimates were rounded off to the nearest integer.

[†] Immunodeficient children were administered OPV, and virus excretion was monitored.

§ Neural isolate obtained at autopsy, approximately 4.3 years after last OPV dose.

[¶] Two different iVDPVs were isolated from the same patient (a type 1 in 1986 and a type 2 in 1992).

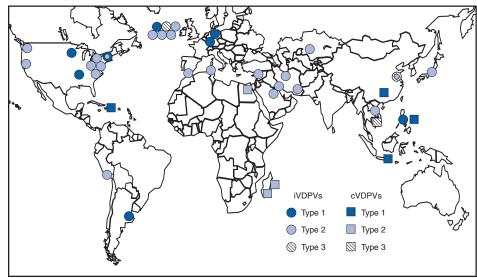
** Information not available.

⁺⁺ New cases reported since publication of previous report (2).

§§ Excretion stopped after bone marrow transplant.

Risk Factors for VDPV Emergence

The key factors favoring cVDPV emergence and spread are the same as for WPV circulation: low OPV coverage, poor sanitation, high population densities, and (usually) tropical conditions. In all but the remaining polio-endemic areas, immunity to polio is no longer acquired from natural infection; immunization is the only current means to prevent the spread of emerging VDPVs or imported WPVs (*3*). Although OPV is not recommended for immunodeficient patients, it is often inadvertently administered because certain primary immunodeficiencies (e.g., common variable immunodeficiency [CVID]) develop later in life. Certain persons with CVID who excrete iVDPVs had onset of polio several years after the implicated OPV dose was administered, and three have demonstrated no signs of paralysis. Survival of patients with primary immunodeficiencies can be extended FIGURE. Locations of persons excreting immunodeficiency-associated vaccinederived polioviruses (iVDPVs), 1962–2006, and polio outbreaks associated with circulating vaccine-derived polioviruses (cVDPVs), 1988–2006



in upper- and middle-income countries by intravenous immunoglobulin therapy; however, for patients in low-income countries, such therapy often is too expensive and difficult to obtain (7).

Global VDPV Surveillance

Since the cVDPV outbreak in Haiti and the Dominican Republic in 2000-2001 (Figure, Table 1), all polioviruses isolated in the WHO Global Poliovirus Laboratory Network from patients with acute flaccid paralysis have been characterized by one molecular method, to identify polioviruses by their genetic properties (usually using the polymerase chain reaction), and one antigenic method, to detect antigenic differences from the OPV strains (using either an enzyme-linked immunosorbent assay [ELISA] or panels of specific neutralizing monoclonal antibodies) (10). Isolates found to be genetically related to an OPV strain but with antigenic differences are possible VDPVs. VP1 sequencing is routinely performed on all possible VDPV and WPV isolates. Approximately 12,000 isolates from all WHO regions have been routinely screened for VDPVs since 2001 (10). Temporal or geographic clustering of vaccine-related isolates of the same serotype has prompted the detection and investigation of cVDPV outbreaks in eight countries (Table 1).

Reported by: WHO Global Poliovirus Laboratory Network. Immunization, Vaccines and Biologicals Dept, WHO, Geneva, Switzerland. Div of Viral Diseases and Global Immunization Div, National Center for Immunization and Respiratory Diseases (proposed), CDC.

Editorial Note: VDPVs will continue to emerge as long as OPV is used. Intensified surveillance has indicated that cVDPVs can emerge repeatedly under conditions of low OPV coverage (e.g., Madagascar). VDPVs also can be found in developed countries with no paralytic cases (e.g., Estonia, Israel, and Slovakia) and can circulate in isolated pockets of unimmunized persons in countries with overall high rates of vaccination coverage (e.g., China and the United States). Although iVDPVs can emerge in middle-income developing countries, cVDPVs have not been found in some areas of high biologic risk, such as in northern India, presumably because of the current high rates of OPV coverage.

Occurrences of VDPVs, including cVDPV-related outbreaks, are rare events, and all recent outbreaks of cVDPVs have been rapidly interrupted using OPV campaigns. The recent increase in the detection of VDPVs is probably primarily attributable to intensified surveillance and improved laboratory methods. Enhanced surveillance for VDPVs has allowed for better understanding of the risks associated with the different types of VDPVs. Areas with continued use of OPV but lacking optimal coverage (e.g., Indonesia in 2005) are at increased risk for cVDPV emergence. The importance of detecting aVDPVs with limited VP1 divergence is not clear; the presence of aVDPVs in certain settings might not have any public health consequences, whereas aVDPVs found elsewhere might signal conditions favoring the emergence of a cVDPV.

Under certain circumstances, OPV viruses regain both neurovirulence and the capacity to circulate and cause outbreaks and therefore are of concern to the PEI. After global eradication of WPVs, the continued use of OPV would continually generate cVDPVs and could eventually pose a challenge to the goal of stopping all poliovirus infections in the human population. The increasing risk of cVDPV emergence in countries with widening immunity gaps and the ongoing risks for vaccine-associated paralytic polio and iVDPVs have prompted an evaluation of the feasibility of orderly cessation of OPV use as soon as possible in the posteradication era (4)while population immunity and surveillance sensitivity are still high (6). Continued development and implementation of a comprehensive strategy to minimize the risks for VDPV emergence in the posteradication era presents a challenge to the PEI and to the public health and scientific communities.

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West Nile Virus Activity — United States, January 1–October 10, 2006

This report summarizes West Nile virus (WNV) surveillance data reported to CDC through ArboNET as of 3 a.m. Mountain Daylight Time, October 10, 2006. A total of 41 states and the District of Columbia had reported 3,135 cases of human WNV illness to CDC (Table, Figure). A total of 1,717 (55%) cases for which such data were available occurred in males; median age of patients was 50 years (range: 3 months–99 years). Dates of illness onset ranged from January 6 to September 25; a total of 97 cases were fatal.

A total of 260 presumptive West Nile viremic blood donors (PVDs) have been reported to ArboNET during 2006. Of these, 40 were reported from Nebraska; 27 from Texas; 24 from Utah; 21 from Colorado; 15 from California; 14 from Louisiana; 11 each from North Dakota and South Dakota; 10 each from Iowa and Wisconsin; nine each from Arizona, Mississippi, and Oklahoma; eight from Kansas; six from Idaho; five each from Minnesota and Virginia; four each from Kentucky and Missouri; three each from Illinois, Montana, and Nevada; two from Michigan; and one each from Arkansas, Maryland, New York, Ohio, Oregon, Pennsylvania, and

TABLE. Number of human cases of West Nile virus (WNV)
illness, by state — United States, 2006*

inness, by s		West	Other	Total	
	Neuroinvasive	Nile	clinical/	reported	
State	disease [†]	fever§	unspecified ¹	to CDC**	Deaths
Alabama	4	0	1	5	0
Arizona	15	14	16	45	3
Arkansas	21	5	0	26	0
California	65	164	13	242	3
Colorado	54	219	0	273	3
Connecticut	6	2	0	8	1
District of Co	lumbia 0	1	0	1	0
Florida	3	0	0	3	0
Georgia	2	4	1	7	1
Idaho	94	542	6	642	10
Illinois	111	55	23	189	9
Indiana	11	5	12	28	0
lowa	17	12	0	29	0
Kansas	14	10	0	24	3
Kentucky	5	1	0	6	1
Louisiana	66	49	0	115	0
Maryland	2	1	1	4	0
Massachuse	tts 2	1	0	3	0
Michigan	29	2	6	37	3
Minnesota	29	34	0	63	3
Mississippi	72	79	0	151	6
Missouri	41	9	1	51	2
Montana	10	19	1	30	0
Nebraska	33	123	0	156	1
Nevada	34	73	14	121	1
New Jersey	2	2	1	5	0
New Mexico	1	2	0	3	0
New York	7	3	1	11	2
North Dakota	a 20	115	0	135	1
Ohio	27	7	0	34	3
Oklahoma	21	12	1	34	5
Oregon	4	42	8	54	0
Pennsylvania	a 7	1	0	8	2
South Dakota	a 37	71	0	108	3
Tennessee	7	1	0	8	1
Texas	175	81	0	256	23
Utah	48	88	0	136	4
Virginia	0	0	2	2	0
Washington	0	2	0	2	0
West Virginia	a 1	0	0	1	0
Wisconsin	10	8	0	18	1
Wyoming	14	36	11	61	2
Total	1,121	1,895	119	3,135	97
	her 10,0000	· -		, -	

* As of October 10, 2006.

[†] 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.

Wyoming. Of the 260 PVDs, three persons (median age: 73 years [range: 26–74 years]) subsequently had neuroinvasive illness, one person aged 41 years had other illness, and 54 persons (median age: 46 years [range: 17–70 years]) had West Nile fever.

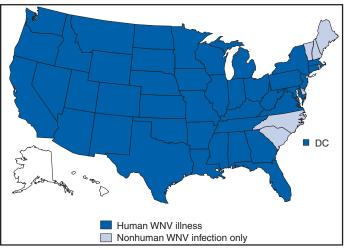


FIGURE. Areas reporting West Nile virus (WNV) activity — United States, 2006*

* As of October 10, 2006.

In addition, 2,746 dead corvids and 636 other dead birds with WNV infection have been reported in 41 states and New York City during 2006. WNV infections have been reported in horses in 34 states, in one squirrel in Kansas, and in two unidentified animal species in North Carolina and Wyoming. WNV seroconversions have been reported in 682 sentinel chicken flocks in 12 states (Arizona, Arkansas, California, Florida, Iowa, Montana, Nevada, North Carolina, North Dakota, Pennsylvania, Utah, and Virginia). A total of 10,157 WNV-positive mosquito pools have been reported from 38 states, the District of Columbia, 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.

Botulism Associated with Commercial Carrot Juice — Georgia and Florida, September 2006

On October 6, this report was posted as an MMWR Dispatch on the MMWR website (http://www.cdc.gov/mmwr).

On September 8, 2006, the Georgia Division of Public Health (GDPH) and CDC were notified of three suspected cases of foodborne botulism in Washington County, Georgia. On September 25, the Florida Department of Health and CDC were notified of an additional suspected case in Tampa, Florida. This report describes the joint investigation and control measures undertaken by state and local health departments, CDC, and the Food and Drug Administration (FDA).

On September 8, the three patients from Washington County, Georgia, went to a local hospital with cranial nerve palsies and progressive descending flaccid paralysis resulting in respiratory failure; the patients had shared meals on September 7. On the evening of September 8, physicians suspected foodborne botulism, notified the state health department, and collected clinical specimens for testing at CDC. On the same evening, CDC provided clinical consultation and dispatched botulinum antitoxin, which was administered to each of the patients the following morning. After receiving antitoxin, the patients had no progression of neurologic symptoms, but they remain hospitalized and on ventilators.

On September 9, the Washington County Health Department, Richmond County Health Department, and GDPH launched an investigation. The three patients had consumed several food items during their two meals together on September 7, including juice from a single 1-liter bottle of Bolthouse Farms carrot juice. The bottle had a "best if used by" date of September 18, 2006. Clinical specimens and leftover food and juice were collected and sent to CDC for testing. On September 13, botulinum toxin type A was identified in the serum and stool of all three patients. On September 15, leftover carrot juice recovered from the home of one of the patients also tested positive for botulinum toxin type A.

During September 8-15, FDA, the Georgia Department of Agriculture, the Georgia Hospital Association, and public health officials in all 50 states were notified of the outbreak and the implicated product as information became available. After these notifications, no additional cases of botulism in Georgia were reported to the state and local health departments or to CDC. During this time, FDA launched an investigation of the Bolthouse Farms, Inc., manufacturing plant in Bakersfield, California. FDA and CDC tested other bottles of the implicated brand of carrot juice, including bottles from different lots, and all were negative for botulinum toxin. Because botulinum toxin was found only in the bottle of carrot juice consumed by the three patients, a lapse in refrigeration of the carrot-juice bottle during transport or storage was suspected, which would have allowed for growth of Clostridium botulinum and subsequent production of botulinum toxin. Based on the CDC test results, on September 17, FDA issued a consumer advisory on the importance of keeping carrot juice refrigerated. However, information obtained from patient interviews regarding storage and transport of the carrot juice did not confirm mishandling by the patients.

On September 25, officials at the Florida Department of Health, the Hillsborough County Health Department, and

MMWR

CDC were notified that a patient had been hospitalized in Tampa, Florida, on September 16, with respiratory failure and descending paralysis. On September 28, botulinum toxin type A was identified in the patient's serum. Circulating toxin persisted more than 10 days after illness onset in this completely paralyzed patient, indicating ingestion of a massive toxin dose. Accordingly, the patient was treated with antitoxin, which prevents binding of circulating botulinum toxin to nerve endings. The patient remains hospitalized, paralyzed, and on a ventilator. The Hillsborough County Health Department collected an open, 450-milliliter bottle of Bolthouse Farms carrot juice, which had been found by a family member in the hotel room where the patient had been staying during the month before being hospitalized. The hotel room had no refrigerator. The bottle, which had a "best if used by" date of September 19, 2006, had a different lot number than the bottle associated with the Georgia cases. On September 29, botulinum toxin was identified in carrot juice from the bottle found in the patient's hotel room; the toxin was subsequently identified as botulinum toxin type A. The Hillsborough County Health Department and CDC notified FDA, public health officials in all 50 states, and infection-control practitioners in Hillsborough County about the botulism case and implicated product. The manufacturer provided FDA with bottles of carrot juice from the same lot as the bottle found in the patient's room. FDA tested juice from all of these bottles, and it was negative for botulinum toxin.

C. botulinum spores are found in the environment and can be present naturally in carrot juice and other foods that have not undergone the retort canning process, which involves high temperatures and high pressure. Anaerobic conditions, low acidity (pH>4.6), low salt and sugar concentrations, and temperatures >39°F (>4°C) promote germination of *C. botulinum* spores and botulinum toxin production. Carrot juice has low acidity, with a natural pH of approximately 6.0; therefore, in the absence of another inhibitor, refrigeration at temperatures <40°F (<4°C) is necessary to prevent germination of *C. botulinum* spores and production of botulinum toxin. Inhibiting *C. botulinum* growth in other ways, such as through acidification, can retard its growth in juice that is not properly refrigerated.

Acidification has been used as a solution to previous foodborne botulism outbreaks. In 1985, 36 patients in the United States and Canada were identified with botulism after eating at a restaurant in Vancouver, British Columbia. A casecontrol study implicated commercially produced, chopped garlic in soybean oil stored at room temperature as the source of the outbreak (1). In 1989, a second outbreak of botulism associated with chopped garlic in oil occurred when three patients in New York were identified with botulism after consuming a meal containing unrefrigerated, commercially produced, chopped garlic in virgin olive oil (2). After these outbreaks, FDA rules were altered to require that garlic-in-oil products contain an acidifying agent such as phosphoric or citric acid.

The carrot juice consumed by these four patients was manufactured by Bolthouse Farms, Inc., and distributed in all 50 states, Mexico, Canada, and Hong Kong with the labels "Bolthouse Farms 100% Carrot Juice," "Earthbound Farm Organic Carrot Juice," and "President's Choice Organics 100% Pure Carrot Juice." Investigations of these cases by state and local health departments and investigations of the manufacturer by FDA are ongoing. On September 29, GDPH and the Georgia Department of Agriculture recommended that Georgia residents not purchase or consume Bolthouse Farms carrot juice. The same day, the FDA warned consumers not to drink Bolthouse Farms carrot juice with "best if used by" dates of November 11, 2006 or earlier (i.e., all bottles produced before the date the warning was issued), and Bolthouse Farms issued a voluntary recall of these products. Additional information regarding the recall is available from the Bolthouse Farms website at http://www.bolthouse.com/ bolthouserecallFAQ.pdf or from FDA (telephone, 888-723-3366).

Suspected botulism cases should be reported immediately to local or state public health officials, who then should call the 24-hour CDC Emergency Operations Center at 770-488-7100; the center will immediately connect them with an on-call botulism specialist. Health-care providers and public health officials are encouraged to inquire specifically about consumption of carrot juice as part of the food history of suspect botulism cases. Additional information on botulism is available at http:// www.cdc.gov/ncidod/dbmd/diseaseinfo/botulism_g.htm.

Reported by: C Shuler, DVM, C Drenzek, DVM, S Lance, DVM, PhD, G Gonzalez, MD, J Miller, MSPH, M Tobin-D'Angelo, MD, J Gabel, DVM, C Burnett, MPH, Georgia Div of Public Health. D Atrubin, MPH, Florida Dept of Health. J Sobel, MD, P Juliao, PhD, S Maslanka, PhD, Div of Foodborne, Bacterial, and Mycotic Diseases, National Center for Zoonotic, Vector-Borne, and Enteric Diseases; P Wiersma, MD, A Sheth, MD, EIS officers, CDC.

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Notice to Readers

Availability of Provisional AIDS and HIV/AIDS Data in MMWR Table IV and Pediatric HIV Surveillance Data in MMWR Table I

CDC is upgrading the national HIV/AIDS surveillance data management system. Because of this transition, CDC will not update AIDS or HIV/AIDS surveillance data for display in quarterly *MMWR* Table IV for the last two quarters of 2006. In addition, CDC will not provide monthly updates of HIV infection data for persons aged <13 years in *MMWR* Table I for the remainder of this year. Explanatory footnotes will be included with Tables I and IV during the period when no updates are available.

Erratum: Vol. 55, No. 35

In the *MMWR* report, "Update: Delayed-Onset *Pseudomonas fluorescens* Bloodstream Infections After Exposure to Contaminated Heparin Flush — Michigan and South Dakota, 2005–2006," the last line of the first column on page 961 should read, "The patients all had indwelling central venous catheters and received treatment during **October 2004**– **February 2005** at clinics known to have used the contaminated flush."

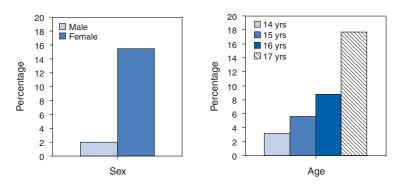
Erratum: Vol. 55, No. 39

In the *MMWR* report, "Childhood Influenza Vaccination Coverage — United States, 2004–05 Influenza Season," on page 1061, an error occurred in the fifth sentence of the first paragraph. The sentence should read, "Others recommended to receive influenza vaccination include children aged **5–18** years who have certain high-risk medical conditions, are on chronic aspirin therapy, or who are household contacts of persons at high risk for influenza complications (1)."

QuickStats

FROM THE NATIONAL CENTER FOR HEALTH STATISTICS

Percentage of Teens Aged 14–17 Years Who Used Indoor Tanning Devices During the Preceding 12 Months, by Sex and Age — United States, 2005*



* Data are based on household interviews of a sample of the civilian, noninstitutionalized population.

The World Health Organization recommends that no person aged <18 years use a tanning bed because of the associated increased risk for skin cancer. In addition, CDC recommends that school programs to prevent skin cancer advise students to avoid using sunlamps and tanning beds. Nonetheless, in 2005, 8.7% of teens aged 14–17 years used indoor tanning devices. Girls aged 14–17 years were seven times more likely to use these devices than boys in the same age group. The use of indoor tanning devices increased with age from 14 to 17 years.

SOURCES: National Health Interview Survey, 2005. Available at http://www.cdc.gov/nchs/nhis.htm.

World Health Organization. The World Health Organization recommends that no person under 18 should use a sunbed. Available at http://www.who.int/mediacentre/news/notes/2005/np07/en/index.html.

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TABLE I. Provisional cases of infrequently reported notifiable diseases (<1,000 cases reported during the preceding year) — United States, week ending October 7, 2006 (40th Week)*

	Current	Cum	5-year weekly	Total o	cases rer	orted for	previou	s vears	
Disease	week	2006	average [†]	2005	2004	2003	2002	2001	States reporting cases during current week (No.)
Anthrax		1	0				2	23	
Botulism:			0				2	20	
foodborne	1	8	0	19	16	20	28	39	FL (1)
infant	1	64	2	90	87	76	69	97	UT (1)
other (wound & unspecified)	_	42	1	33	30	33	21	19	01(1)
Brucellosis	_	78	2	122	114	104	125	136	
Chancroid	_	23	0	17	30	54	67	38	
Cholera	_	6	0	8	5	2	2	3	
Cyclosporiasis§	_	91	1	734	171	75	156	147	
Diphtheria	_	_	_	_	_	1	1	2	
Domestic arboviral diseases ^{§,¶} :									
California serogroup	_	33	5	80	112	108	164	128	
eastern equine	_	6	0	21	6	14	10	9	
Powassan	_	1		1	1	_	1	N	
St. Louis	_	3	1	13	12	41	28	79	
western equine	_	_	_			_	_	_	
Ehrlichiosis [§] :									
human granulocytic	3	282	8	790	537	362	511	261	NY (1), PA (1), AL (1)
human monocytic	4	281	8	522	338	321	216	142	NY (1), MO (2), TN (1)
human (other & unspecified)	1	130	1	122	59	44	23	6	MO (1)
laemophilus influenzae,**									
invasive disease (age <5 yrs):									
serotype b	_	8	0	9	19	32	34	_	
nonserotype b	2	67	2	135	135	117	144	_	CT (1), OK (1)
unknown serotype	1	152	2	217	177	227	153	_	ID (1)
lansen disease§	3	55	1	88	105	95	96	79	NH (1), FL (2)
lantavirus pulmonary syndrome§	1	25	0	29	24	26	19	8	AZ (1)
lemolytic uremic syndrome, postdiarrheal§	7	183	5	221	200	178	216	202	OH (1), MN (2), NC (2), AL (1), UT (1)
lepatitis C viral, acute	11	594	32	771	713	1,102	1,835	3,976	OH (2), MI (3), MO (1), MD (1), OK (1), TX (1),
									WA (1), OR (1)
HV infection, pediatric (age <13 yrs)§,††	_	52	4	380	436	504	420	543	
nfluenza-associated pediatric mortality ^{§,§§,111}	_	40	0	45		N	N	N	
isteriosis	7***	498	19	892	753	696	665	613	NY (2), PA (1), OH (1), FL (1), AL (1), WA (1)
leasles		43	0	66	37	56	44	116	
Aeningococcal disease, ^{†††} invasive:		170	0	007					14/4 (4)
A, C, Y, & W-135	1	170	3	297	_	_	_	_	WA (1)
serogroup B	_	108	2	157	_		_	_	
other serogroup	9	14	0 5	27 314	258	231	270	266	
/umps	9	5,791	5 0	314					NY (1), PA (1), MO (1), NC (6)
Plague	I	12	0		3	1	2	2	UT (1)
Poliomyelitis, paralytic Psittacosis [§]	_	17	0	1 19	12	12	18	25	
2 fever [§]	4	120	1	139	70	71	61	25 26	MO (2), TN (1), TX (1)
Rabies, human	4	120	0	2	70	2	3	20	MO(2), TN(T), TA(T)
Rubella	1	8	0	11	10	2	18	23	NY (1)
Rubella, congenital syndrome		1		1		1	10	23	
SARS-CoV ^{§,§§}	_		_		_	8	N	N	
Smallpox [§]	_	_	_	_	_	_			
Streptococcal toxic-shock syndromes		78	1	129	132	161	118	77	
Streptococcus pneumoniae,§		70		123	102	101	110		
invasive disease (age <5 yrs)	15	787	11	1,257	1,162	845	513	498	NY (3), MI (2), MN (2), NE (2), OK (1), TX (2), CO (3)
Syphilis, congenital (age <1 yr)	5	205	8	361	353	413	412	490	NY (2), IL (1), NC (1), AZ (1)
etanus	_	17	0	27	34	20	25	37	
oxic-shock syndrome (other than streptococ	cal)§ —	72	2	96	95	133	109	127	
richinellosis		11	0	19	5	6	109	22	
ularemia [§]	_	68	3	154	134	129	90	129	
vphoid fever	_	220	9	324	322	356	321	368	
	reus§ —	220	-	2	522	330 N	N	308 N	
ancomycin-intermediate Stanhylococcus au				~					
/ancomycin-intermediate Staphylococcus au /ancomycin-resistant Staphylococcus aureus		_	0	3	1	N	N	N	

-: No reported cases. N: Not notifiable. Cum: Cumulative year-to-date counts.

Incidence data for reporting year 2006 is provisional, whereas data for 2001, 2002, 2003, 2004, and 2005 are finalized.

Calculated by summing the incidence counts for the current week, the two weeks preceding the current week, and the two 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. t

8 Not notifiable in all states.

Not notifiable in all states. Includes both neuroinvasive and non-neuroinvasive. Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Zoonotic, Vector-Borne, and Enteric Diseases (proposed) (ArboNET Surveillance). 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 (proposed). Implementation of HIV reporting influences the number of cases reported. Pediatric HIV data will not be updated monthly for the remainder of this year due to upgrading of the national HIV/ AIDS surveillance data management system. Data for HIV/AIDS are available in Table IV quarterly. ††

§§ Updated weekly from reports to the Influenza Division, National Center for Immunization and Respiratory Diseases (proposed).

11 Cumulative totals for 2005 and 2006 do not include reports from states where influenza-associated pediatric mortality is not a notifiable condition.

*** No measles cases were reported for the current week.

ttt Data for meningococcal disease (all serogroups and unknown serogroups) are available in Table II.



Recommended Adult Immunization Schedule — United States, October 2006–September 2007

Weekly

October 13, 2006 / Vol. 55 / No. 40

The Advisory Committee on Immunization Practices (ACIP) annually reviews the recommended Adult Immunization Schedule to ensure that the schedule reflects current recommendations for the licensed vaccines. In June 2006, ACIP approved the Adult Immunization Schedule for October 2006–September 2007. This schedule has also been approved by the American Academy of Family Physicians and the American College of Obstetricians and Gynecologists.

Changes in the Schedule for October 2006–September 2007

The 2006–2007 schedule differs from the previous schedule as follows:

- The broken red line has been deleted on the age-based schedule (Figure 1). Vaccination of persons with specific risk factors is now shown only with purple bars.
- Human papillomavirus (HPV) vaccine has been added to the age-based schedule, with a yellow bar indicating that the vaccine is recommended for women ≤26 years.
- Tetanus, diphtheria, and acellular pertussis (Tdap) vaccine has been added to the age-based schedule, with a hatched yellow bar indicating that Tdap is a one-time, 1-dose recommendation for persons ≤64 years.
- The purple bar for varicella vaccine has been shortened in anticipation of the recommendation for the use of zoster vaccine in persons aged ≥60 years.
- A new column has been added to the medical/other indications schedule (Figure 2) to clarify indications for hepatitis A and B vaccines. The indications "chronic liver disease" and "recipients of clotting factor concentrates" have been removed from the previous schedule's third and fifth columns, respectively, and combined into a new column. The column has a yellow bar for hepatitis A and B vaccines, clarifying that these vaccines are recommended for all persons with these medical indications.
- HPV vaccine has been added to the medical/other indications schedule, with a yellow bar to indicate the vaccine

Suggested citation: Centers for Disease Control and Prevention. Recommended Adult Immunization Schedule—United States, October 2006–September 2007. MMWR 2006;55:Q1–Q4. is recommended for women aged ≤ 26 years with all indications except pregnancy.

- Tdap was added to the medical/other indications schedule, with a hatched yellow bar to indicate that Tdap is a one-time, 1-dose recommendation for all indications except pregnancy.
- The tetanus and diphtheria footnote (#1) has been reworded to reflect ACIP recommendations for use of Tdap.
- A footnote (#2) has been added to reflect ACIP recommendations for HPV vaccination for all women aged ≤26 years.
- The measles, mumps, and rubella (MMR) footnote (#3) has been reworded to reflect ACIP recommendations to administer a second dose of mumps vaccine to adults in certain age groups and with certain risk factors.
- The varicella footnote (#4) has been reworded in accordance with ACIP recommendations for administering a routine second dose for all adults without evidence of immunity. The footnote also has been revised to reflect the new definition of immunity to varicella.
- The influenza footnote (#5) has been revised to reflect recent ACIP recommendations to vaccinate close contacts of children aged 0–59 months rather than 0–23 months (1).
- The hepatitis B footnote (#9) has been revised to reflect recommendations to vaccinate any adult seeking protection from hepatitis B virus infection and vaccinate adults in specific settings (e.g., sexually transmitted disease clinics) (2).

The Adult Immunization Schedule is available in English and Spanish at http://www.cdc.gov/nip/recs/adultschedule.htm. General information about adult vaccinations, including recommendations concerning vaccination of person with HIV and other immunosuppressive conditions, is available from state and local health departments and at http:// www.cdc.gov/nip. Vaccine information statements are available at http://www.cdc.gov/nip/publications/vis. ACIP statements for each recommended vaccine and provisional vaccine recommendations can be viewed, downloaded, and printed at http://www.cdc.gov/nip/publications/acip-list.htm. Instructions for reporting adverse events to the Vaccine Adverse Event Reporting System are available at http://www.vaers.hhs.gov or by telephone, 800-822-7967.

References

- 1. CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 2006;55(No. RR-10).
- 2. CDC. A comprehensive immunization strategy to eliminate transmission of hepatitis B virus infection in the United States: recommendations of the Advisory Committee on Immunization Practices (ACIP). Part II: immunization of adults. MMWR. In press 2006.

The Recommended Adult Immunization Schedule has been approved by the Advisory Committee on Immunization Practices, the American College of Obstetricians and Gynecologists, and the American Academy of Family Physicians. The standard *MMWR* footnote format has been modified for publication of this schedule.

		Age group (yrs)	
Vaccine	19–49	50–64	≥65
Tetanus, diphtheria, pertussis (Td/Tdap) ¹ *	Substitute 1 do	1-dose Td booster every 10 yrs se of Tdap for Td	
Human papillomavirus (HPV) ^{2*}	3 doses (females)		
Measles, mumps, rubella (MMR) ³ *	1 or 2 doses	1 d	ose
Varicella ⁴ *	2 doses (0, 4–8 wks)	2 doses (0, 4–8 wks)	
Influenza ⁵ *	1 dose annually	1 dose a	annually
Pneumococcal (polysaccharide) ^{6,7}	1–2 c	loses	1 dose
Hepatitis A ⁸ *	2	doses (0, 6–12 mos, or 0, 6–18 mo	s)
Hepatitis B ⁹ *		3 doses (0, 1–2, 4–6 mos)	P
Meningococcal ¹⁰		1 or more doses	
	hs in this category who meet the age requirements and e.g., lack documentation of vaccination or have no evide		me other risk factor is present (e.g., on the cupational, lifestyle, or other indications)

FIGURE 1. Recommended adult immunization schedule, by vaccine and age group — United States, October 2006–September 2007

* Covered by the Vaccine Injury Compensation Program.

NOTE: These recommendations must be read along with the footnotes, which can be found on pages Q2–Q4 of this schedule.

Approved by the Advisory Committee on Immunization Practices, the American College of Obstetricians and Gynecologists, and the American Academy of Family Physicians

1. Tetanus, diphtheria, and acellular pertussis (Td/Tdap) vaccination. Adults with uncertain histories of a complete primary vaccination series with diphtheria and tetanus toxoid-containing vaccines should begin or complete a primary vaccination series. A primary series for adults is 3 doses; administer the first 2 doses at least 4 weeks apart and the third dose 6-12 months after the second. Administer a booster dose to adults who have completed a primary series and if the last vaccination was received >10 years previously. Tdap or tetanus and diphtheria (Td) vaccine may be used; Tdap should replace a single dose of Td for adults aged <65 years who have not previously received a dose of Tdap (either in the primary series, as a booster, or for wound management). Only one of two Tdap products (Adacel [Sanofi pasteur, Swiftwater, Pennsylvania]) is licensed for use in adults. If the person is pregnant and received the last Td vaccination \geq 10 years previously, administer Td during the second or third trimester; if the person received the last Td vaccination in <10 years, administer Tdap during the immediate postpartum period. A one-time administration of 1-dose of Tdap with an interval as short as 2 years from a previous Td vaccination is recommended for postpartum women, close contacts of infants aged <12 months, and all health-care workers with direct patient contact. In certain situations, Td can be deferred during pregnancy and Tdap substituted in the immediate postpartum period, or Tdap can be given instead of Td to a pregnant woman after an informed discussion with the woman (see http://www.cdc.gov/nip/ publications/acip-list.htm). Consult the ACIP statement for recommendations for administering Td as prophylaxis in wound management (http://www.cdc.gov/mmwr/preview/mmwrhtml/ 00041645.htm).

2. Human papillomavirus (HPV) vaccination. HPV vaccination is recommended for all women aged ≤26 years who have not completed the vaccine series. Ideally, vaccine should be administered before potential exposure to HPV through sexual activity; however, women

who are sexually active should still be vaccinated. Sexually active women who have not been infected with any of the HPV vaccine types receive the full benefit of the vaccination. Vaccination is less beneficial for women who have already been infected with one or more of the four HPV vaccine types. A complete series consists of 3 doses. The second dose should be administered 2 months after the first dose: the third dose should be administered 6 months after the first dose. Vaccination is not recommended during pregnancy. If a woman is found to be pregnant after initiating the vaccination series, the remainder of the 3-dose regimen should be delayed until after completion of the pregnancy. 3. Measles, mumps, rubella (MMR) vaccination. Measles component: adults born before 1957 can be considered immune to measles. Adults born during or after 1957 should receive >1 dose of MMR unless they have a medical contraindication, documentation of ≥ 1 dose, history of measles based on health-care provider diagnosis, or laboratory evidence of immunity. A second dose of MMR is recommended for adults who 1) have been recently exposed to measles or in an outbreak setting; 2) have been previously vaccinated with killed measles vaccine; 3) have been vaccinated with an unknown type of measles vaccine during 1963-1967; 4) are students in postsecondary educational

during 1963–1967; 4) are students in postsecondary educational institutions; 5) work in a health-care facility; or 6) plan to travel internationally. Withhold MMR or other measles-containing vaccines from HIV-infected persons with severe immunosuppression. *Mumps component:* adults born before 1957 can generally be considered immune to mumps. Adults born during or after 1957 should receive 1 dose of MMR unless they have a medical contraindication, history of mumps based on health-care provider diagnosis, or laboratory evidence of immunity. A second dose of MMR is recommended for adults who 1) are in an age group that is affected during a mumps outbreak; 2) are students in postsecondary educational institutions; 3) work in a health-care workers born before 1957 who do not have other evidence of

FIGURE 2. Recommended adult immunization schedule, by vaccine and medical and other indications — United States, October 2006–September 2007

				Indica	ation			
Vaccine	Pregnancy	Congenital immuno- deficiency, leukemia, ¹¹ lymphoma, generalized malignancy, cerebrospinal fluid leaks, therapy with alkylating agents, anti- metabolites, radiation, or high-dose, long-term corticosteroids	Diabetes, heart disease, chronic pulmonary disease, chronic alcoholism	Asplenia ¹¹ (including elective splenectomy and terminal complement component deficiencies)	Chronic liver disease, recipients of clotting factor concentrates	Kidney failure, end-stage renal disease, recipients of hemodialysis	Human immuno- deficiency virus (HIV) infection ^{3,11}	Health-care workers
Tetanus, diphtheria, pertussis (Td/Tdap) ^{1*}			1-0	dose Td boost	er every 10 y			
Human papillomavirus (HPV) ² *			3 dose			6 yrs (0, 2, 6 n	nos)	
Measles, mumps, rubella (MMR) ³ *					1	or 2 doses		
Varicella ⁴ *				2 doses (0, 4–8 wks)			2 doses
Influenza ⁵ *		1 dose annually		1 dose annually		1 dose a	innually	
Pneumococcal (polysaccharide) ^{6,7}	1–2 doses			1–2 d	oses			1–2 doses
Hepatitis A ^{8*}	2 do	l ses (0, 6–12 mo l	s, or 0, 6–18	3 mos)	2 dos	es (0, 6–12 mo	os, or 0, 6–18	mos)
Hepatitis B ^{9*}		3 doses (0, 1–	2, 4–6 mos)			3 doses (0, 1-	-2, 4–6 mos)	
Meningococcal ¹⁰		1 dose		1 dose		1 dc)Se	
For all persons in this categ of immunity (e.g., lack docu						risk factor is present al, lifestyle, or other in		Contraindicated

* Covered by the Vaccine Injury Compensation Program.

NOTE: These recommendations must be read along with the footnotes, which can be found on pages Q2–Q4 of this schedule.

mumps immunity, consider giving 1 dose on a routine basis and strongly consider giving a second dose during an outbreak. *Rubella component:* administer 1 dose of MMR vaccine to women whose rubella vaccination history is unreliable or who lack laboratory evidence of immunity. For women of childbearing age, regardless of birth year, routinely determine rubella immunity and counsel women regarding congenital rubella syndrome. Do not vaccinate women who are pregnant or who might become pregnant within 4 weeks of receiving vaccine. Women who do not have evidence of immunity should receive MMR vaccine upon completion or termination of pregnancy and before discharge from the health-care facility.

4. Varicella vaccination. All adults without evidence of immunity to varicella should receive 2 doses of varicella vaccine. Special consideration should be given to those who 1) have close contact with persons at high risk for severe disease (e.g., health-care workers and family contacts of immunocompromised persons) or 2) are at high risk for exposure or transmission (e.g., teachers of young children; child care employees; residents and staff members of institutional settings, including correctional institutions; college students; military personnel; adolescents and adults

living in households with children; non-pregnant women of childbearing age; and international travelers). Evidence of immunity to varicella in adults includes any of the following: 1) documentation of 2 doses of varicella vaccine at least 4 weeks apart; 2) U.S.-born before 1980 (although for health-care workers and pregnant women, birth before 1980 should not be considered evidence of immunity); 3) history of varicella based on diagnosis or verification of varicella by a health-care provider (for a patient reporting a history of or presenting with an atypical case, a mild case, or both, health-care providers should seek either an epidemiologic link with a typical varicella case or evidence of laboratory confirmation, if it was performed at the time of acute disease); 4) history of herpes zoster based on health-care provider diagnosis; or 5) laboratory evidence of immunity or laboratory confirmation of disease. Do not vaccinate women who are pregnant or might become pregnant within 4 weeks of receiving the vaccine. Assess pregnant women for evidence of varicella immunity. Women who do not have evidence of immunity should receive dose 1 of varicella vaccine upon completion or termination of pregnancy and before discharge from the health-care facility. Dose 2 should be administered 4-8 weeks after dose 1.

5. Influenza vaccination. Medical indications: chronic disorders of the cardiovascular or pulmonary systems, including asthma; chronic metabolic diseases, including diabetes mellitus, renal dysfunction, hemoglobinopathies, or immunosuppression (including immunosuppression caused by medications or HIV); any condition that compromises respiratory function or the handling of respiratory secretions or that can increase the risk of aspiration (e.g., cognitive dysfunction, spinal cord injury, or seizure disorder or other neuromuscular disorder); and pregnancy during the influenza season. No data exist on the risk for severe or complicated influenza disease among persons with asplenia; however, influenza is a risk factor for secondary bacterial infections that can cause severe disease among persons with asplenia. Occupational indications: health-care workers and employees of long-term-care and assisted living facilities. Other indications: residents of nursing homes and other long-term-care and assisted living facilities; persons likely to transmit influenza to persons at high risk (i.e., in-home household contacts and caregivers of children aged 0-59 months, or persons of all ages with high-risk conditions); and anyone who would like to be vaccinated. Healthy, nonpregnant persons aged 5-49 years without high-risk medical conditions who are not contacts of severely immunocompromised persons in special care units can receive either intranasally administered influenza vaccine (FluMist[®]) or inactivated vaccine. Other persons should receive the inactivated vaccine.

6. Pneumococcal polysaccharide vaccination. *Medical indications:* chronic disorders of the pulmonary system (excluding asthma); cardiovascular diseases; diabetes mellitus; chronic liver diseases, including liver disease as a result of alcohol abuse (e.g., cirrhosis); chronic renal failure or nephrotic syndrome; functional or anatomic asplenia (e.g., sickle cell disease or splenectomy [if elective splenectomy is planned, vaccinate at least 2 weeks before surgery]); immunosuppressive conditions (e.g., congenital immunodeficiency, HIV infection [vaccinate as close to diagnosis as possible when CD4 cell counts are highest], leukemia, lymphoma, multiple myeloma, Hodgkin disease, generalized malignancy, or organ or bone marrow transplantation); chemotherapy with alkylating agents, antimetabolites, or high-dose, long-term corticosteroids; and cochlear implants. *Other indications:* Alaska Natives and certain American Indian populations and residents of nursing homes or other long-term–care facilities.

7. Revaccination with pneumococcal polysaccharide vaccine. Onetime revaccination after 5 years for persons with chronic renal failure or nephrotic syndrome; functional or anatomic asplenia (e.g., sickle cell disease or splenectomy); immunosuppressive conditions (e.g., congenital immunodeficiency, HIV infection, leukemia, lymphoma, multiple myeloma, Hodgkin disease, generalized malignancy, or organ or bone marrow transplantation); or chemotherapy with alkylating agents, antimetabolites, or high-dose, long-term corticosteroids. For persons aged \geq 65 years, one-time revaccination if they were vaccinated \geq 5 years previously and were aged <65 years at the time of primary vaccination.

8. Hepatitis A vaccination. *Medical indications:* persons with chronic liver disease and persons who receive clotting factor concentrates. *Behavioral indications:* men who have sex with men and persons who use illegal drugs. *Occupational indications:* persons working with hepatitis A virus (HAV)–infected primates or with HAV in a research laboratory setting. *Other indications:* persons traveling to or working in countries that have high or intermediate endemicity of hepatitis A (a

list of countries is available at http://www.cdc.gov/travel/diseases.htm) and any person who would like to obtain immunity. Current vaccines should be administered in a 2-dose schedule at either 0 and 6–12 months, or 0 and 6–18 months. If the combined hepatitis A and hepatitis B vaccine is used, administer 3 doses at 0, 1, and 6 months.

9. Hepatitis B vaccination. Medical indications: persons with endstage renal disease, including patients receiving hemodialysis; persons seeking evaluation or treatment for a sexually transmitted disease (STD); persons with HIV infection; persons with chronic liver disease; and persons who receive clotting factor concentrates. Occupational indications: health-care workers and public-safety workers who are exposed to blood or other potentially infectious body fluids. Behavioral indications: sexually active persons who are not in a long-term, mutually monogamous relationship (i.e., persons with >1 sex partner during the previous 6 months); current or recent injection-drug users; and men who have sex with men. Other indications: household contacts and sex partners of persons with chronic hepatitis B virus (HBV) infection; clients and staff members of institutions for persons with developmental disabilities; all clients of STD clinics; international travelers to countries with high or intermediate prevalence of chronic HBV infection (a list of countries is available at http://www.cdc.gov/travel/diseases.htm); and any adult seeking protection from HBV infection. Settings where hepatitis B vaccination is recommended for all adults: STD treatment facilities; HIV testing and treatment facilities; facilities providing drug-abuse treatment and prevention services; health-care settings providing services for injection-drug users or men who have sex with men; correctional facilities; end-stage renal disease programs and facilities for chronic hemodialysis patients; and institutions and nonresidential daycare facilities for persons with developmental disabilities. Special formulation indications: for adult patients receiving hemodialysis and other immunocompromised adults, 1 dose of 40 µg/mL (Recombivax HB[®]) or 2 doses of 20 µg/mL (Engerix-B^w).

10. Meningococcal vaccination. Medical indications: adults with anatomic or functional asplenia, or terminal complement component deficiencies. Other indications: first-year college students living in dormitories; microbiologists who are routinely exposed to isolates of Neisseria meningitidis; military recruits; and persons who travel to or live in countries in which meningococcal disease is hyperendemic or epidemic (e.g., the "meningitis belt" of sub-Saharan Africa during the dry season [December-June]), particularly if their contact with local populations will be prolonged. Vaccination is required by the government of Saudi Arabia for all travelers to Mecca during the annual Hajj. Meningococcal conjugate vaccine is preferred for adults with any of the preceding indications who are aged \leq 55 years, although meningococcal polysaccharide vaccine (MPSV4) is an acceptable alternative. Revaccination after 5 years might be indicated for adults previously vaccinated with MPSV4 who remain at high risk for infection (e.g., persons residing in areas in which disease is epidemic)

11. Selected conditions for which *Haemophilus influenzae* type b (Hib) vaccine may be used. Hib conjugate vaccines are licensed for children aged 6 weeks–71 months. No efficacy data are available on which to base a recommendation concerning use of Hib vaccine for older children and adults with the chronic conditions associated with an increased risk for Hib disease. However, studies suggest good immunogenicity in patients who have sickle cell disease, leukemia, or HIV infection or who have had splenectomies; administering vaccine to these patients is not contraindicated.

Additional information about the vaccines in this schedule and contraindications for vaccination is also available at http://www.cdc.gov/nip or from the CDC-INFO Contact Center at 800-CDC-INFO (800-232-4636) in English and Spanish, 24 hours a day, 7 days a week.

This schedule indicates the recommended age groups and medical indications for routine administration of currently licensed vaccines for persons aged \geq 19 years, as of October 1, 2006. Licensed combination vaccines may be used whenever any components of the combination are indicated and when the vaccine's other components are not contraindicated. For detailed recommendations on all vaccines, including those used primarily for travelers or that are issued during the year, consult the manufacturers' package inserts and the complete statements from the Advisory Committee on Immunization Practices (http://www.cdc.gov/nip/publications/acip-list.htm).

Report all clinically significant postvaccination reactions to the Vaccine Adverse Event Reporting System (VAERS). Reporting forms and instructions on filing a VAERS report are available at http://www.vaers.hhs.gov or by telephone, 800-822-7967.

Information on how to file a Vaccine Injury Compensation Program claim is available at http://www.hrsa.gov/vaccinecompensation or by telephone, 800-338-2382. To file a claim for vaccine injury, contact the U.S. Court of Federal Claims, 717 Madison Place, N.W., Washington, D.C. 20005; telephone, 202-357-6400.

			Chlamyd	lia†			Coccidioidomycosis						Cryptosporidiosis				
	0		vious	0	0	0		ious	0	0	0		vious	0	0		
Reporting area	Current week	Med	veeks Max	Cum 2006	Cum 2005	Current week	Med	eeks Max	Cum 2006	Cum 2005	Current week	Med	veeks Max	Cum 2006	Cum 2005		
United States	10,833	18,961	35,170	723,079	736,812	10	149	1,643	6,216	3,222	111	69	594	3,594	5,738		
New England	1,068	623	1,550	24,919	24,817	_	0	0	_	_	8	4	29	235	285		
Connecticut Maine [§]	400 51	167 43	1,214 74	7,196 1,725	7,275 1,714	N N	0 0	0 0	N N	N N	_	0 0	26 4	26 32	64 25		
Massachusetts	543	284	442	11,392	11,086	—	0	0	—	—	_	1	14	88	129		
New Hampshire Rhode Island	51	37 60	65 100	1,511 2,244	1,424 2,572	_	0 0	0 0	_	_	4	1 0	4 6	36 11	30 7		
Vermont [§]	23	19	43	851	746	Ν	Ő	Ő	Ν	Ν	4	Ő	5	42	30		
Mid. Atlantic New Jersev	1,475 132	2,380 375	3,696 497	91,650 14,080	90,744 14,850	N	0 0	0 0	N	N	3	11 0	444 3	403 9	2,286		
New York (Upstate)	444	499	1,727	18,506	17,969	N	0	0	N	N	2	3	441	130	52 1,900		
New York City Pennsylvania	439 460	731 739	1,570 1,074	29,148 29,916	29,441 28,484	N N	0 0	0 0	N N	N N	1	1 5	9 13	50 214	118 216		
E.N. Central	1,836	3,123	12,578	121,131	123,466	_	1	3	37	9	18	16	109	878	1,364		
Illinois	647	964	1,691	39,135	38,714		0	0	_	_	_	2	8	72	141		
Indiana Michigan	320 684	393 645	510 9,888	15,042 26,632	15,526 20,545	N	0 0	0 3	N 33	N 9	5 2	1 2	18 7	68 102	59 89		
Ohio	41	686	1,433	25,115	33,067		0	1	4	N	10	5	76	285	650		
Wisconsin W.N. Central	144 488	399 1,152	531 1,456	15,207 44,315	15,614 45,432	N	0 0	12	N 1	4	1 14	5 11	52 73	351 645	425 516		
Iowa	—	154	225	5,730	5,484	Ν	0	0	Ν	Ν	_	1	28	151	113		
Kansas Minnesota	_	154 230	269 346	5,443 8,222	5,665 9,493	N	0 0	0 12	N	N 3		1 2	7 22	58 155	32 103		
Missouri	332	441	612	17,497	17,469		0	1	1	1	1	2	18	145	221		
Nebraska [§] North Dakota	71 21	95 33	176 58	4,109 1,273	3,960 1,230	N N	0 0	1 0	N N	N N	_2	1 0	16 4	72 8	19 1		
South Dakota	64	51	116	2,041	2,131	Ν	0	0	Ν	Ν	—	1	7	56	27		
S. Atlantic	3,117 63	3,454 68	4,927 92	138,603	137,469 2,572	N	0 0	1 0	3	1 N	46	14 0	63 3	753	546 3		
Delaware District of Columbia	23	52	103	2,714 1,829	2,9572		0	0	<u>N</u>		_	0	3	11 12	9		
Florida Georgia	840 1	942 635	1,154 2,142	37,497 22,464	33,379 24,134	N	0 0	0 0	N	N	30 4	6 3	32 11	356 154	243 113		
Maryland§	307	331	486	13,456	14,176	_	0	1	3	1	_	0	3	15	26		
North Carolina South Carolina [§]	1,017 349	562 306	1,772 1,306	25,920 13,895	24,902 14,846	N N	0 0	0 0	N N	N N	8 4	0 1	11 13	79 81	69 18		
Virginia§	495	423	840	18,390	18,458	N	0	0	N	N	_	1	6	38	53		
West Virginia E.S. Central	22	57	226	2,438	2,045	N	0	0 0	N	N		0 3	3	7	12		
Alabama [§]	552	1,419 391	1,947 756	55,669 15,314	53,492 11,982	N	0	0	N	N	11 10	1	20 7	140 61	173 21		
Kentucky Mississippi	4	160 382	402 802	6,427 14,283	6,858 16,599	N	0 0	0 0	N	N	1	1 0	19 3	31 14	117 2		
Tennessee§	548	495	599	19,645	18,053	Ν	0	0	Ν	Ν	—	1	5	34	33		
W.S. Central	615	2,151	3,605	82,185	84,961	_	0	1	1	_	4	4	26	194	185		
Arkansas Louisiana	181 225	158 265	333 761	6,270 11,278	6,692 12,966	_	0 0	0 1	1	N	1	0 0	2 7	18 41	4 72		
Oklahoma Texas [§]	209	228 1,392	2,159 1,774	9,209 55,428	8,735 56,568	N N	0 0	0 0	N N	N N	3	1 2	4 20	32 103	36 73		
Mountain	1,300	1,027	1,839	38,608	48,496	10	114	452	4,330	2,100	5	2	38	280	111		
Arizona	881	354	642	14,119	16,579	10	111	448	4,256	2,020	_	0	2	19	9		
Colorado Idaho§	62 191	156 50	482 159	4,512 2,236	11,670 1,998	N N	0 0	0 0	N N	N N	2 1	1 0	7 5	58 25	38 13		
Montana Nevedo [§]	100	42	195	1,825	1,775	N	0 0	0 4	N	N	—	0	26	104 4	16		
Nevada [§] New Mexico [§]	166	77 172	432 339	3,732 7,422	5,642 6,504	_	0	4	21 13	48 16	_	0	1 4	4 16	11 10		
Utah Wyoming	_	94 27	170 55	3,731 1,031	3,457 871	_	1 0	3 2	38 2	13 3	2	0 0	3 11	16 38	11 3		
Pacific	382	3,315	5,079	125,999	127,935	_	43	1,179	1,844	1,108	2	2	52	66	272		
Alaska	57	85	152	3,247	3,263	—	0	0	_	· _	—	0	1	4	3		
California Hawaii	_	2,570 103	4,231 135	98,604 3,948	99,331 4,264	N	43 0	1,179 0	1,844 N	1,108 N	_	0 0	14 1	4	156 1		
Oregon [§] Washington	325	174 350	315 604	6,624 13,576	6,836 14,241	N N	0 0	0 0	N N	N N	2	1 0	6 38	58	60 52		
American Samoa	325 U	0	46	13,576 U	14,241 U	U	0	0	U	U IN	 U	0	0	 U	52 U		
C.N.M.I.	Ŭ	0	0	Ŭ	Ū	Ŭ	0	0	U	Ŭ	Ŭ	0	0	Ŭ	Ŭ		
Guam Puerto Rico	_	18 75	37 161	2,945	629 3,187	N	0 0	0 0	N	N	N	0 0	0 0	N	N		
U.S. Virgin Islands	_	5	16	178	196	_	0	0	_	_	_	0	0	_	_		

Med: Median.

Max: Maximum.

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending October 7, 2006, and October 8, 2005

Cum: Cumulative year-to-date counts.

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-or * Incidence data for reporting year 2006 is provisional. * Chlamydia refers to genital infections caused by *Chlamydia trachomatis*. § Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

(40th Week)*	Giardiasis						G	ionorrhe	a		Hae		<i>is influen</i> es, all se	<i>izae</i> , inva rotypes	sive
	Current		/ious /eeks	Cum	Cum	Current		vious veeks	Cum	Cum	Current		vious veeks	Cum	Cum
Reporting area	week	Med	Max	2006	2005	week	Med	Max	2006	2005	week	Med	Max	2006	2005
United States	297	322	1,029	12,510	14,626	3,548	6,499	14,136	250,416	252,809	14	38	142	1,550	1,777
New England Connecticut	23 8	24 0	75 37	967 222	1,331 280	145 82	107 41	288 241	4,211 1,698	4,476 1,908	1 1	2 0	19 9	127 38	136 40
Maine [†]	6	2	13	133	168	2	2	6	98	110	_	0	4	17	8
Massachusetts New Hampshire	1	10 0	25 9	357 24	602 50	58 1	46 4	86 9	1,855 149	1,936 127	_	1 0	7 2	52 7	68 7
Rhode Island Vermont [†]		0 3	25 10	92 139	86 145	2	8 1	19 4	360 51	350 45	_	0 0	7 2	4 9	7 6
Mid. Atlantic	39	61	254	2,239	2,646	448	636	1,014	24,324	26,068	2	7	30	299	333
New Jersey New York (Upstate)	27	9 24	13 227	297 911	351 909	88 107	102 123	143 455	3,767 4,787	4,399 5,240	2	1 2	4 27	45 103	67 97
New York City	2 10	10 15	31 29	438 593	708 678	98 155	175 215	357 394	7,198	7,873 8,556	_	1 3	4	36 115	61 108
Pennsylvania E.N. Central	28	49	29 86	1,846	2,612	654	1,285	7,047	8,572 49,286	50,104	2	5	14	219	308
Illinois Indiana	N	9	21 0	317 N	615 N	207 137	377 161	709 237	14,969	15,252 6,259	2	1	6 11	47 66	104 54
Michigan	3	14	22	502	631	262	252	5,880	6,713 11,097	8,512	—	Ö	3	18	19
Ohio Wisconsin	25	16 10	32 40	624 403	605 761	15 33	329 134	648 172	11,390 5,117	15,637 4,444	_	2 0	6 4	65 23	94 37
W.N. Central	19	29	260	1,423	1,614	173	364	436	14,024	14,447	2	2	15	112	88
lowa Kansas	1	5 3	15 11	223 148	215 158	_	33 44	46 124	1,199 1,519	1,219 2,010	_	0 0	1 3	1 14	9
Minnesota Missouri	 15	2 10	238 32	477 420	667 365	155	62 190	105 251	2,113 7,753	2,660 7,309	2	0 0	9 6	56 30	37 29
Nebraska [†]	3	2	8	86	103	11	23	56	1,062	899	_	0	2	7	12
North Dakota South Dakota	_	0 1	7 7	12 57	12 94	1 6	3 6	7 15	87 291	79 271	_	0 0	3 0	4	1
S. Atlantic	80	49	95	1,920	2,118	1,187	1,491	2,334	61,369	60,107	1	10	26	409	420
Delaware District of Columbia	_	1 1	4 5	33 52	44 41	27 24	27 34	44 61	1,132 1,238	663 1,633	_	0 0	1 1	1 4	7
Florida Georgia	40 29	18 10	39 44	821 411	743 565	381 4	437 300	553 1,014	17,853 10,865	15,359 11,261	1	3 2	9 12	133 80	103 90
Maryland† North Carolina	8 N	4 0	11 0	158 N	161 N	102 332	128 284	186 766	5,022 13,093	5,324 11,903	_	1 0	5 9	53 46	58 68
South Carolina [†]	_	1	7	69	86	151	132	704	6,262	6,798	—	1	3	27	28
Virginia† West Virginia	3	8 0	50 5	359 17	446 32	148 18	130 17	288 42	5,161 743	6,628 538	_	1 0	8 4	49 16	43 23
E.S. Central	20	8	40	377	329	205	564	863	22,544	21,249	_	2	7	79	94
Alabama⁺ Kentucky	13 N	4 0	29 0	204 N	149 N	7	183 55	310 132	7,110 2,301	6,880 2,353	_	0 0	5 1	20 4	17 11
Mississippi Tennessee [†]	7	0 4	0 12	173	180	198	141 187	435 237	5,607 7,526	5,401 6,615	_	0 1	1 4	3 52	66
W.S. Central	11	5	31	209	248	291	879	1,430	35,567	34,532	3	1	15	54	95
Arkansas Louisiana	6	2 0	6 3	92 18	66 51	97 123	79 161	142 354	3,241 6,889	3,499 7,183	_	0 0	2 2	7 5	7 32
Oklahoma Texas†	5 N	2 0	24 0	99 N	131 N	71	82 541	764 836	3,440 21,997	3,482 20,368	3	1 0	14 2	40 2	51 5
Mountain	47	30	56	1,230	1,147	362	217	552	8,715	10,447	2	4	8	158	183
Arizona Colorado	 21	3 9	36 33	116 439	108 400	198 84	90 41	201 90	3,541 1,595	3,768 2,459	1	1 1	7 4	73 42	92 37
Idaho† Montana	7	3	11 11	134 79	111 58	15	2	10 20	132 145	84 121	1	0	1	4	4
Nevada [†]	_	1	6	41	83	65	24	194	1,230	2,215	_	0	1	_	14
New Mexico [†] Utah	18	1 7	6 19	47 344	68 299	_	31 17	64 25	1,348 631	1,211 530	_	0 0	4 4	21 15	21 8
Wyoming	1	1	4	30	20	—	2	6	93	59	—	0	1	3	7
Pacific Alaska	30 3	59 1	202 15	2,299 75	2,581 86	83 11	807 11	963 23	30,376 451	31,379 447	1	2 0	15 2	93 9	120 26
California Hawaii	_	43 1	105 3	1,606 37	1,833 52	1	659 18	830 29	24,950 707	26,144 788	_	0	9 1	21 14	50 8
Oregon [†]	7	8	14	308	344	_	28	58	1,016	1,174	1	1	6	47	36
Washington American Samoa	20 U	6 0	90 0	273 U	266 U	71 U	75 0	142 2	3,252 U	2,826 U	U	0	4 0	2 U	 U
C.N.M.I.	Ū	0	0	Ŭ	U	Ū	0	0	U	U	Ŭ	0	0	Ū	U
Guam Puerto Rico	4	0	0 12	62	11 211	_	1 5	15 16	188	71 286	_	1	2	1	7 3
U.S. Virgin Islands	—	0	0	—		—	0	5	30	45	—	0	0	—	_

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending October 7, 2006, and October 8, 2005 (40th Week)*

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-* Incidence data for reporting year 2006 is provisional. * Contains data reported through the National Electronic Disease Surveillance System (NEDSS). Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.

(40th Week)*	,	Hepatitis (viral, acute), by type															
			A	нер	atitis (virai	, acute), by	Legionellosis										
			/ious			-	Previ						vious	-			
Reporting area	Current week	52 w Med	veeks Max	Cum 2006	Cum 2005	Current week	<u>52 we</u> Med	eks Max	Cum 2006	Cum 2005	Current week	<u>52 v</u> Med	veeks Max	Cum 2006	Cum 2005		
United States	17	69	245	2,463	3,245	45	84	597	3,064	4,035	37	44	127	1,636	1,574		
New England	_	3	20	144	371	_	1	9	49	119	9	2	12	99	106		
Connecticut Maine [†]	_	1 0	2 2	34 6	43 3	_	0 0	3 2	 15	39 12	9	0 0	8 2	38 7	22 5		
Massachusetts	_	1	13	51	234	_	0	5	14	39	_	1	6	27	49		
New Hampshire Rhode Island	_	0 0	16 4	36 9	75 10	_	0 0	2 4	11 8	24 1	_	0 0	1 10	1 20	8 16		
Vermont [†]	—	0	2	8	6	—	0	1	1	4	_	0	3	6	6		
Mid. Atlantic New Jersey	4	7 2	18 7	270 61	525 109	3	8 2	55 8	317 80	527 195	16	13 1	42 10	553 61	537 91		
New York (Upstate)	4	1	14	67	81	_	1	43	49	45	11	5	29	230	137		
New York City Pennsylvania	_	2 1	12 5	92 50	252 83	3	2 3	4 9	60 128	110 177	5	1 4	9 17	46 216	83 226		
E.N. Central	2	7	12	224	284	2	8	24	311	438	2	8	25	335	326		
Illinois Indiana	1	1 0	4 5	50 23	103 14	_	2 0	7 17	57 42	124 33	_	1 0	4 3	21 24	46 21		
Michigan	_	2	8	78	88	1	3	7	105	143	_	2	7	86	89		
Ohio Wisconsin	1	1 1	4 5	45 28	41 38	1	2 0	10 4	101 6	104 34	2	4 0	19 5	171 33	141 29		
W.N. Central	1	2	30	103	70	—	4	22	125	213	_	1	15	52	63		
lowa Kansas	_	0 0	2 5	8 24	18 13	_	0 0	3 2	14 8	21 24	_	0 0	3 2	10 3	4 2		
Minnesota Missouri	1	0 1	29 3	9 39	3 28	_	0 2	13 7	17 74	29 111	_	0	11 3	11 18	16 24		
Nebraska [†]	_	0	3	15	8	_	0	1	11	22	_	0	2	6	3		
North Dakota South Dakota	_	0	2 3	8	_	_	0	0 1	1	6	_	0 0	1 6	4	2 12		
S. Atlantic	6	11	29	419	572	12	23	66	889	1,081	6	9	19	322	304		
Delaware District of Columbia	_	0	2 2	10 6	5 3	_	1 0	4 2	35 5	25 10	_	0 0	2 5	8 16	13 9		
Florida	4	4	13	165	228	8	8	19	322	370	2	3	9	130	84		
Georgia Maryland†	1	1	7 6	50 53	108 58	1	3 3	7 10	124 128	166 121	4	0 1	4 6	15 65	27 87		
North Carolina South Carolina [†]	_	0 0	20 2	67 17	70 34	3	0 2	23 7	123 63	128 121	_	0 0	5 1	29 2	24 11		
Virginia†	1	1	11	46	63	_	1	18	43	113	_	1	7	49	35		
West Virginia E.S. Central	2	0 2	3 8	5 97	3 216	4	0 6	18 15	46 249	27 284	- 1	0 1	3 9	8 67	14 62		
Alabama [†]		0	3	13	40	—	1	8	78	67	_	0	2	9	12		
Kentucky Mississippi	_	0	5 1	29 5	22 17	1	1 0	5 2	57 11	54 44	1	0 0	4	23 1	21 3		
Tennessee [†]	2	1	5	50	137	3	2	8	103	119	—	1	7	34	26		
W.S. Central Arkansas	2	4 0	77 9	138 35	376 16	17 1	14 1	315 4	578 37	478 53	_	1 0	32 3	43 3	38 5		
Louisiana	_	0	4	15	56	_	0	3	25	62	_	0	2	4	1		
Oklahoma Texas†	2	0 2	2 73	6 82	4 300	12 4	0 12	17 295	43 473	37 326	_	0 0	3 26	1 35	7 25		
Mountain	_	5	18	192	253	_	4	39	126	429	3	2	7	95	79		
Arizona Colorado	_	2 1	16 4	108 33	135 34	_	1	23 5	33 29	276 44	1	1 0	4 2	32 21	19 17		
Idaho†	—	0	2	9	20	—	0	2 7	10	12	1	0	3	11	3		
Montana Nevada†	_	0 0	3 2	9 7	7 19	_	0 0	4	15	3 42	_	Ō	1 2	5 4	5 17		
New Mexico† Utah	_	0	3 2	12 11	19 18	_	0	3 5	15 24	18 32	1	0 0	1	4 18	3 11		
Wyoming	—	Ő	1	3	1	—	0	1		2	_	0	ò		4		
Pacific Alaska	_	19 0	163 0	876	578 4	7	9 0	61 1	420 5	466 7	_	1 0	9 1	70	59		
California	_	15	162	793	477	_	7	41	317	312	_	1	9	70	57		
Hawaii Oregon†	_	0	2 5	9 37	21 38	1	0 1	1 5	5 54	6 84	N	0 0	1 0	N	2 N		
Washington	_	1	13	37	38	6	0	18	39	57	_	0	0	_	_		
American Samoa C.N.M.I.	U U	0 0	0 0	U U	1 U	U U	0 0	0 0	U U	 U	U U	0 0	0 0	U U	U U		
Guam	_	0	0	_	2	_	0	0	_	18	_	0	0	_	_		
Puerto Rico U.S. Virgin Islands	_	0 0	5 0	23	58	_	1 0	8 0	24	38	_	0 0	1 0	1	_		
U		-	-				-	-				-	-				

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending October 7, 2006, and October 8, 2005 (40th Week)*

Cum: Cumulative year-to-date counts.

Max: Maximum.

Med: Median.

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-* Incidence data for reporting year 2006 is provisional. * Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

Pervious Pervious Current Mad Mark Current Mark <th cols<="" th=""><th>(40th Week)*</th><th></th><th></th><th>Lyme dis</th><th>ease</th><th></th><th></th><th></th><th>Malaria</th><th>1</th><th></th><th></th></th>	<th>(40th Week)*</th> <th></th> <th></th> <th>Lyme dis</th> <th>ease</th> <th></th> <th></th> <th></th> <th>Malaria</th> <th>1</th> <th></th> <th></th>	(40th Week)*			Lyme dis	ease				Malaria	1		
Begeoting area Week Hed Max 2006 2005 week Her Max 2006 2005 United State 101 252 2,153 10,234 17,704 10 24 125 916 1,996 Opennechal 90 13 14 464 233 - 0 1 4 5 Mainsi 16 13 34 164 233 - 0 3 19 34 Mainsi - 1 14 78 132 - 0 3 9 2 Vermont - 1 14 78 10,314 2 4 13 283 699 New Metry - 1 10 122 168 10,314 2 1 13 32 29 New Metry - 1 14 160 10,05 2 2 7 101 117 New M			Pre	evious				Prev	vious				
United States 101 252 2,153 13,234 17,704 13 24 125 916 1.096 New ingland 13 37 783 2,217 3.094 1 14 163 Mained 16 1 34 164 213 0 1 14 15 Massed-inserts 1 35 33 2,123 0 3 9 34 Massed-inserts 1 14 78 41 0 1 1 1 Memode lian 1 17 7.681 10.314 -2 4 13 163 297 New Vor Civit D<	Benarting area												
New Enginal 29 37 770 2,217 3,044 1 11 44 63 Mainal 10 1 33 15,19 431 0 1 14 54 Mainal 10 1 33 15,19 0 3 19 54 Mainal 0 3 19 35 54 54 54 54 54 54 54 54 54 54 54 54 54 54 54 54 54 54 56 56 57 57 57 77 1 3 55 59 56													
Connesfort 11 11 13 753 15.19 491 — 0 5 11 16 Mand' 11 1 33 753 15.19 491 — 0 5 11 16 Mand' 11 1 33 753 16 12 213 — 0 1 4 5 Mary Lampabrie 2 6 59 42 2.183 — 0 8 — 2 Vermont — 1 13 77 78 11 — 0 1 1 1 7 New Jerson 11 7 78 41 — 0 1 1 1 7 New Jerson 12 7 18 12 — 1 3 163 237 New Jerson 12 7 18 12 3 1.17 78 141 — 0 1 1 1 7 New Jerson 12 1 18 224 349 — 2 1 8 64 160 Pennsylvana 11 40 122 2.657 3.770 — 1 3 35 29 Pennsylvana 11 40 122 2.657 3.770 — 1 4 3 163 237 New Jerson 11 40 122 2.657 3.770 — 1 4 42 66 Pennsylvana 11 40 122 2.657 3.770 — 1 4 42 66 Pennsylvana 11 40 122 2.657 3.770 — 1 4 42 66 Pennsylvana 11 40 122 2.657 3.770 — 1 4 42 66 Pennsylvana 11 40 122 2.657 3.770 — 1 4 42 66 Pennsylvana 11 40 12 4 1.161 1.65 2 2 7 10 117 Hillinois — 1 0 1 2 1.066 1.364 — 0 3 27 10 Wisconsin — 1 0 1 2 1.066 1.364 — 0 3 27 10 Wisconsin — 1 0 12 10 166 1.364 — 0 3 27 10 Wisconsin — 1 0 12 10 166 1.364 — 0 3 27 10 Wisconsin — 1 0 12 10 166 1.364 — 0 3 27 10 Wisconsin — 0 1 1 6 3 9 13 — 0 4 6 15 Minasout — 0 3 9 13 — 0 1 6 16 Minasout — 0 3 9 13 — 0 1 6 16 Minasout — 0 3 7 10 Wisconsin — 0 1 1 1 2 — 0 1 1 1 — S.Atanic & 8 32 108 1.116 1.841 6 6 6 15 258 234 Delivic O'olumbia 2 0 7 44 8 — 0 2 2 3 8 Masout — 0 3 1 1 2 — 0 0 1 1 1 — S.Atanic & 8 32 108 1.116 1.841 6 6 6 15 258 234 Delivic O'olumbia 2 0 7 44 8 — 0 2 2 3 8 Mary Masout — 0 3 1 1 2 — 0 1 1 1 — S.Atanic & 8 32 108 1.116 1.841 6 1 6 16 Maryadri — 0 2 4 24 42 — 0 2 3 14 24 South Diskin — 0 3 1 21 11 — S.Atanic & 8 32 108 1.116 1.841 — 0 2 3 1 5 5 7 88 Maryadri — 0 2 4 24 42 — 0 1 4 3 3 Delivic O'olumbia 2 0 7 4 4 8 — 0 2 2 3 8 Delivic O'olumbia 2 0 7 4 4 9 0 10 — 0 8 24 24 South Diskin — 0 1 8 24 29 10 10 — 0 8 24 24 South Diskin — 0 9 1 4 3 3 0 9 10 — 0 9 2 1 4 13 0 1 1 1 9 5 16 0 1 1 1 9													
Massachusetis - 1 35 33 2,129 - 0 3 19 34 Wertangehold - 1 14 78 41 - 0 3 19 54 Wertangehold - 1 14 78 41 - 0 1 1 1 New draft 1 1 67 76 150 3.114 - 1 3 68 69 New draft 1 168 1.666 3.111 - 1 1 32 69 New draft 1 1 10 124 1.666 3.111 - 1 1.66 364 160 1											16		
New Hampshire 2 6 50 4.22 188 - 0 3 - 2 Vermont* - 0 1 14 77 1 10 1													
Phode leakad - 0 5 1 32 - 0 8 - 2 Md. Alantic 62 153 1,176 7,681 10.314 2 4 1													
Hid. Alarkic 62 153 1.76 7.681 10.314 2 4 13 163 297 New York (Dipitate) 51 75 1.150 3.244 3.084 2 1 13 283 39 New York (Dipitate) 11 40 222 2.657 3.770 1 3 35 29 E.N. Central 1 10 24 1.61 1.663 2 2 7 101 117 Binos - 0 2 - 1 44 46 66 Wisconsin - 10 129 1.066 0 3 27 18 Wisconsin - 10 129 1.066 1.844 0 3 27 18 Wisconsin - 10 129 1.864 0 3 27 10 Nisconsin - 0 1 497 646 0 30 1 5 5 16 16	Rhode Island	—	0	5	1	32		0	8	—	2		
New Jersey													
New York (Lipstate) 51 75 1,160 3,244 3,084 2 1 11 36 39 Pennsylvania 1 140 222 2,657 3,770 1 3 35 29 Illinois 0 2 - 101 117 Illinois 0 2 7 101 117 Wisconsin 1 6 44 44 1 4 42 64 Ohio 1 6 38 49 2 0 3 7 10 Wisconsin 1 8 349 2 0 3 2 16 14 11 6 14 11 11 2 14 11 11 2 11 33 3 3 3 11 11 2 11 11 2 11 11 11 11													
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 TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending October 7, 2006, and October 8, 2005

 (40th Week)*

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

 U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-or * Incidence data for reporting year 2006 is provisional.
 * Contains data reported through the National Electronic Disease Surveillance System (NEDSS).
 Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.

(40th Week)*		Meningococcal disease, invasive																	
			All serog	roups				<u> </u>	Inknown				Pertus	ssis					
	Current		vious weeks	Cum	Cum	Current	Previ 52 we		Cum	Cum	Current		/ious /eeks	Cum	Cum				
Reporting area	week	Med	Max	2006	2005	week	Med	Max	2006	2005	week	Med	Max	2006	2005				
United States	5	20	85	840	962	4	13	58	548	587	129	265	2,877	9,945	17,567				
New England	_	1	3	35	61	_	0	2	25	22	1	29	83	959	1,057				
Connecticut Maine [†]	_	0 0	2 1	9 4	12 2	_	0 0	2 1	2 3	1 2	_	1 1	5 11	35 63	52 40				
Massachusetts New Hampshire	_	0 0	2 2	15 5	28 12	_	0 0	2 2	15 5	5 12	_	19 2	43 36	594 129	805 54				
Rhode Island	_	0	1		2	_	0	0		_	_	0	17	45	29				
Vermont [†]		0	1	2	5	_	0	0		2	1	1	14	93	77				
Mid. Atlantic New Jersey	1	3 0	14 2	121 11	118 27	1	2 0	11 2	90 11	90 27	50	34 3	137 13	1,412 156	1,052 143				
New York (Úpstate)		1	7	31	31	_	0	5	4	11	34	14	123	644	402				
New York City Pennsylvania	1	0 1	6 5	42 37	18 42	1	0 0	6 5	42 33	18 34	16	1 11	8 26	64 548	86 421				
E.N. Central	1	3	11	96	120	1	1	6	65	99	23	39	133	1,406	2,986				
Illinois Indiana	_	0 0	4 5	18 19	27 18	_	0 0	4 1	18 6	27 8	5	7 4	35 75	230 189	685 252				
Michigan	_	0	3	19	24	_	0	3	8	15	4	7	24	389	246				
Ohio Wisconsin	1	1 0	5 2	37 3	32 19	1	1 0	4 2	30 3	30 19	14	14 4	30 41	459 139	907 896				
W.N. Central	_	1	4	45	64	_	0	3	15	28	16	28	552	955	2,908				
lowa Kansas	_	0 0	2 1	13 1	15 9	_	0 0	1 1	5 1	1 9	_	6 7	63 28	212 226	733 334				
Minnesota	_	0	2	11	11	_	0	1	3	4	14	0	485	160	966				
Missouri Nebraska†	_	0	2 2	13 5	22 4	_	0	1	2 3	11 3	1	7 2	42 9	241 73	361 232				
North Dakota	—	0	1	1	—	_	0	1	1	_	—	0	25	26	112				
South Dakota S. Atlantic	2	0 3	1 14	1 149	3 182	2	0 2	0 7	61		9	0 20	4 46	17 758	170				
Delaware		0	14	4	4		0	1	4	4	_	0	1	3	1,133 15				
District of Columbia Florida	2	0 1	1 6	1 59	5 69	2	0	1 5	1 21	4 27	2 3	0 4	3 9	6 172	7 165				
Georgia	_	0	2	12	14	—	0	2	12	14	_	0	3	15	41				
Maryland† North Carolina	_	0 0	2 11	11 24	19 28	_	0 0	1 3	2 7	3 6	1 1	3 0	9 22	97 155	162 98				
South Carolina† Virginia†	_	0 0	2 4	18 15	13 24	_	0 0	2 3	8 6	8 9	2	3 2	22 27	129 155	322 284				
West Virginia	_	0	2	5	6	_	0	0		2	_	0	9	26	39				
E.S. Central	_	1	4	31	47	_	1	4	25	36	1	7	16	261	435				
Alabama [†] Kentucky	_	0 0	1 2	5 7	5 16	_	0 0	1 2	4 7	3 16	_	1 1	7 5	54 53	72 130				
Mississippi Tennessee [†]	_	0 0	1 2	3 16	5 21	_	0 0	1 2	3 11	5 12	1	1 2	4 10	35 119	48 185				
W.S. Central	_	1	23	52	21 93	_	0	2 6	23	23	- -	2 16	360	520	1,842				
Arkansas	_	0	3	9	12	_	0	2	6	3	_	1	21	47	245				
Louisiana Oklahoma	_	0 0	2 4	6 8	28 14	_	0 0	1 0	3	5 2	_	0 0	3 124	11 18	44 1				
Texas [†]	—	1	16	29	39	—	0	4	14	13	—	14	215	444	1,552				
Mountain	—	1	5	57 16	80	_	0 0	4	27	21	22	61	230	2,099	3,260				
Arizona Colorado	_	0 0	3 2	16 19	31 17	_	0	3 1	16 2	10	2 5	9 20	177 40	402 650	819 1,040				
Idaho† Montana	_	0 0	2 1	3 4		_	0 0	2 1	2 2	3	_2	2 2	8 9	74 96	174 549				
Nevada [†]	_	0	1	3	12	_	0	Ó	—	2	—	0	9	39	43				
New Mexico† Utah	_	0 0	1 1	3 5	5 11	_	0 0	1 0	1	4 2	13	2 15	6 39	60 716	152 439				
Wyoming	—	0	2	4	—	_	0	2	4	—	—	1	8	62	44				
Pacific Alaska	1	5 0	29 1	254 2	197 3	_	5 0	25 1	217 2	191 3	7	42 2	1,334 15	1,575 61	2,894 110				
California	_	3	14	156	128	_	3	14	156	128	—	27	1,136	1,099	1,359				
Hawaii Oregon†	_	0 1	1 7	7 60	11 36	_	0 1	1 4	7 41	6 36	_	2 2	4 8	65 94	142 598				
Washington	1	0	25	29	19	_	0	11	11	18	7	7	195	256	685				
American Samoa C.N.M.I.	U U	0 0	0 0	_	_	U U	0 0	0 0	U U	U U	U U	0 0	0 0	U U	U U				
Guam		0	0	_	1	_	0	0	—	1	_	0	0	—	2				
Puerto Rico U.S. Virgin Islands	_	0	1 0	4	6	_	0 0	1 0	4	6	_	0 0	1 0	1	5				
		-	-				-	-				-	-						

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending October 7, 2006, and October 8, 2005 (40th Week)*

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-* Incidence data for reporting year 2006 is provisional. * Contains data reported through the National Electronic Disease Surveillance System (NEDSS). Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.

(40th Week)*	,					,		,							
		Ra Prev	abies, ani	mal		Roc	ky Mour Prev	<u> </u>	tted feve	r			almonello vious	osis	
	Current		eeks	Cum	Cum	Current	52 w		Cum	Cum	Current		vious veeks	Cum	Cum
Reporting area	week	Med	Max	2006	2005	week	Med	Max	2006	2005	week	Med	Max	2006	2005
United States	30	109	173	4,654	4,807	3	37	246	1,599	1,349	640	809	2,291	30,471	33,347
New England Connecticut	7 4	11 3	26 14	529 160	574 156	_	0 0	2 0	2	7	6	30 0	383 375	1,533 375	1,776 392
Maine [†]	—	2	7	84	50	Ν	0	0	N	N	—	2	10	91 782	139
Massachusetts New Hampshire	1	0	17 5	178 39	288 12	_	0	1 1	1 1	5 1	4	18 2	53 24	160	941 144
Rhode Island Vermont [†]	2	0 1	4 4	20 48	20 48	_	0 0	2 0	_	1	2	0 1	17 5	73 52	81 79
Mid. Atlantic	- 1	23	59	1,103	779	1	1	6	56	79	59	86	272	3,551	4,069
New Jersey New York (Upstate)	N	0 11	0 22	N 416	N 435	_	0 0	2 2	7 4	25 1	37	14 22	43 233	630 966	805 965
New York City	1	0	5	20	23	_	0	4	13	6	2	18	36	694	955
Pennsylvania	_	14	42	667	321	1	1	3	32	47	20	29	67	1,261	1,344
E.N. Central Illinois	1 1	1 0	18 7	141 44	163 47	_	0 0	6 1	32 3	37 11	72	99 25	172 45	3,926 854	4,597 1,527
Indiana Michigan	_	0 0	2 5	11 41	11 35	_	0 0	1 1	5 2	5	35 3	15 17	67 32	711 749	479 749
Ohio		0	9	45	70	_	0	4	21	19	34	23	56	983	1,058
Wisconsin	N	0 5	0	N OF 1	N		0	1 15	1	2		15	27	629	784
W.N. Central Iowa	4	0	20 7	251 52	281	1	2 0	1	190 4	141 5	35	42 7	107 21	1,994 335	2,040 339
Kansas Minnesota	_	1	5 6	61 36	70 61	_	0 0	1 2	2 4	5 2	22	6 10	16 60	259 552	296 443
Missouri	4	1	8	65	65	1	2	11	159	117	12	14	36	587	634
Nebraska† North Dakota	_	0 0	0 7	16	28	_	0 0	5 1	21	7	1	4 0	9 46	142 19	170 28
South Dakota	—	0	4	21	57	—	0	0	—	5	—	3	7	100	130
S. Atlantic Delaware	_	36 0	118 0	1,562	1,719	_	16 0	94 3	894 18	674 7	321	207 2	450 9	8,192 117	9,223 105
District of Columbia	—	0	0			—	0	1	1	2	2	1	7	50	45
Florida Georgia	_	0 2	99 9	136 100	201 214	_	0 0	3 3	15 28	13 84	139 52	95 27	228 100	3,449 1,243	3,593 1,464
Maryland [†] North Carolina	_	7 9	13 22	254 397	308 391	_	1 15	6 87	60 663	60 356	18 85	12 32	29 130	540 1,231	645 1,219
South Carolina [†]	_	3	10	129	176	_	0	6	23	61	21	19	51	720	1,113
Virginia† West Virginia	_	10 1	27 13	458 88	383 46	_	2 0	13 2	83 3	86 5	4	20 2	57 19	751 91	905 134
E.S. Central	8	4	16	197	128	_	5	26	273	247	30	54	149	2,197	2,329
Alabama† Kentucky	8	1 0	7 5	69 23	67 16	_	1 0	8 1	78 1	64 3	5 7	14 8	71 22	729 350	550 393
Mississippi Tennessee [†]		0	2 9	4 101	5 40	_	0 3	1 20	2 192	13 167	 18	12 14	47 31	541 577	729 657
W.S. Central	1	14	34	549	744	1	1	161	192	137	62	83	922	2,904	3,226
Arkansas	1	0	4	26	31	—	0	10	46	98	44	14	45	703	566
Louisiana Oklahoma	_	0 1	0 9	52	67	_	0 0	1 154	1 35	6 7	2 16	12 7	35 48	400 382	734 326
Texas [†]	_	12	29	471	646	1	0	3	19	26		48	839	1,419	1,600
Mountain Arizona	8 6	3 2	12 10	156 119	236 152	_	0 0	6 6	44 8	25 12	19 2	50 15	84 67	1,864 587	1,836 499
Colorado		0	1	2	16	—	0	1	2	4 3	6	12	30 9	515	472
Idaho† Montana	_	0 0	12 2	13	15	_	0	3 2	13 2	3	3	3 3	16	137 107	114 69
Nevada [†] New Mexico [†]	_	0	1 2	1 7	14 9	_	0 0	0 2	7	3	_	1 4	17 12	72 179	146 212
Utah	1	0	1	9	14	—	0	2	6	_	8	5	15	230	255
Wyoming Pacific	1	0 4	2 10	5 166	16 183	_	0 0	1	6 7	2 2		1 110	5 426	37 4,310	69 4,251
Alaska	_	0	4	14	1	_	0	0	_	_	1	1	7	62	45
California Hawaii	_	3 0	10 0	135	176	_	0 0	1 0	5	_	_	88 5	292 10	3,369 181	3,217 237
Oregon [†]		0	4	17	6		0	1	2	2	1	7	16	324	331
Washington American Samoa	U U	0 0	0	U U	U U	N U	0	0 0	N U	N U	34 U	7 0	124 1	374 U	421 7
C.N.M.I.	U	0	Ō	U	Ū	Ŭ	Ō	0	U	Ū	U	0	0	U	U
Guam Puerto Rico	_	0 1	0 6	66	 55	N	0 0	0 0	N	N		1 6	3 35	189	30 506
U.S. Virgin Islands	—	0	0	_	_	_	0	0	_	_	_	Ō	0	_	_

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending October 7, 2006, and October 8, 2005 (40th Week)*

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: No U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-* Incidence data for reporting year 2006 is provisional. * Contains data reported through the National Electronic Disease Surveillance System (NEDSS). Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.

(40th Week)*	Shiga toxin-producing <i>E. coli</i> (STEC) [†]					Shigellosis					Streptococcal disease, invasive, group A				
	Shiga	Prev		<i>E. COII</i> (5)			Prev	0			Previous				
Dementing and	Current	52 w	eeks	Cum	Cum	Current	52 w	eeks	Cum	Cum	Current	52 w	eeks	Cum	Cum
Reporting area United States	week 54	Med 56	Max 297	2006 2,264	2005 2,419	220	Med 242	Max 1,013	2006 8,895	2005 11,307	42	Med 87	283	2006 3,828	2005 3,635
New England Connecticut Maine ⁶ Massachusetts New Hampshire Rhode Island Vermont ⁶	1 - 1 	3 0 1 0 0	59 58 8 9 3 2 2	217 58 30 82 22 8 2	184 50 28 71 14 5 16		4 0 3 0 0 0	57 51 2 11 4 6 2	205 51 3 128 7 11 5	259 47 13 157 12 14 16	2 U 2 	4 0 2 0 0 0	15 3 2 6 9 3 2	177 U 15 101 44 5 12	235 82 12 107 16 9 9
Mid. Atlantic New Jersey New York (Upstate) New York City Pennsylvania	3 	5 0 0 0 0	107 3 103 4 5	158 3 12 27 5	286 62 110 13 101	5 4 1	14 4 4 2	72 26 60 12 5	589 206 188 128 67	1,046 267 219 347 213	5 4 1	15 3 4 1 6	43 8 32 9 13	699 122 251 74 252	732 151 209 143 229
E.N. Central Illinois Indiana Michigan Ohio Wisconsin	11 4 7	11 1 1 3 2	52 7 8 7 18 39	503 59 68 69 150 157	503 120 50 78 126 129	5 - 3 2	20 7 2 3 3 3	38 16 18 10 11 9	703 229 113 119 130 112	890 305 121 192 82 190	3 — — 3 —	14 4 2 3 4 1	43 11 11 12 19 4	666 144 92 183 205 42	757 252 86 179 161 79
W.N. Central lowa Kansas Minnesota Missouri Nebraska [§] North Dakota South Dakota	10 — 7 5 —	8 2 3 2 1 0 0	35 8 3 27 13 8 15 5	335 108 	397 82 39 120 82 42 6 26	21 — 17 4 —	34 2 3 2 13 2 0 4	77 10 20 10 69 14 18 21	1,225 77 103 122 574 102 63 184	1,220 68 167 70 783 83 4 45	10 N 9 1 	5 0 1 0 1 0 0	57 0 52 5 4 5 3	285 N 46 136 61 25 9 8	225 N 35 86 57 18 9 20
S. Atlantic Delaware District of Columbia Florida Georgia Maryland [§] North Carolina South Carolina [§] Virginia [§] West Virginia	3 - 1 2 7 	7 0 2 1 1 1 0 0	39 2 1 29 6 8 10 2 8 2	350 7 2 75 69 69 90 6 - 7	322 8 75 44 66 44 9 74 2	118 1 73 43 - 1 - -	54 0 26 17 2 1 1 1 0	122 2 66 41 10 21 9 8 2	2,150 8 14 1,064 714 94 125 69 60 2	1,682 10 9 818 435 68 149 84 108 1	12 2 9 1 	22 0 6 5 4 0 1 2 0	43 2 16 11 12 26 6 11 6	927 10 13 234 176 169 138 51 110 26	725 5 8 188 153 143 104 30 72 22
E.S. Central Alabama [§] Kentucky Mississippi Tennessee [§]	5 1 3 	3 0 1 0 0	15 5 8 1 4	171 29 69 24	135 26 51 8 50	32 17 3 — 12	13 3 4 1 3	31 17 12 8 10	530 176 171 61 122	1,017 194 257 75 491	N 	3 0 0 0 3	11 0 5 0 9	161 N 33 128	141 N 28 — 113
W.S. Central Arkansas Louisiana Oklahoma Texas [§]	8 6 2	1 0 0 1	52 3 1 8 44	37 19 — 18 64	84 11 18 22 33	6 1 5	34 1 3 29	596 7 25 286 308	1,153 81 90 100 882	2,813 51 119 524 2,119	2 — — 2	7 0 0 2 4	58 5 1 14 43	304 24 5 81 194	254 15 5 93 141
Mountain Arizona Colorado Idaho [§] Montana Nevada [§] New Mexico [§] Utah Wyoming	1 1 5 	5 1 1 0 0 1 0	16 8 7 1 3 1 13 3	233 76 87 56 — 10 4 103 17	239 23 62 32 14 18 22 61 7	20 4 10 5 1	22 11 3 0 0 2 1 0	60 30 18 4 6 8 10 6 3	889 468 180 15 12 30 114 62 8	641 340 108 10 5 44 96 33 5	6 5 — — 1	11 6 3 0 0 1 1 0	78 57 8 2 0 2 7 7 1	523 277 112 8 — 63 60 3	486 206 149 3 — 8 68 49 3
Pacific Alaska California Hawaii Oregon [§] Washington	12 — — 3 12	7 0 4 0 2 1	55 1 18 2 47 32	260 — 161 12 99 87	269 9 103 10 72 75	13 — — 1 12	40 0 32 1 2 2	148 2 104 4 31 43	1,451 9 1,189 33 110 110	1,739 11 1,488 28 112 100	2 2 	2 0 2 0 0	9 0 9 0	86 — 86 N N	80 — 80 N N
American Samoa C.N.M.I. Guam Puerto Rico U.S. Virgin Islands	U U —	0 0 0 0	0 0 0 0	U U —	U U 2	U U 1	0 0 0 0	0 0 3 2 0	U U 12	7 U 16 5	U U N	0 0 0 0	0 0 0 0	U U N	U U N

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending October 7, 2006, and October 8, 2005 (40th Week)*

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

Cum: Cumulative year-to-date counts.

Max: Maximum.

Med: Median.

¹ Incidence data for reporting year 2006 is provisional.
 ¹ Incidence data for reporting year 2006 is provisional.
 ¹ Incidence *E. coli* O157:H7; Shiga toxin positive, serogroup non-0157; and Shiga toxin positive, not serogrouped.
 ⁸ Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

Max: Maximum.

(40th Week)*	Streptococcus pneumoniae, invasive disease Drug resistant, all ages					Sypl	nilis, prin	nary and	seconda	ry	Varicella (chickenpox)					
		Prev		•			Previ						/ious			
Reporting area	Current week	52 w Med	еекs Max	Cum 2006	Cum 2005	Current week	52 we Med	<u>екs</u> Max	Cum 2006	Cum 2005	Current week	Med	veeks Max	Cum 2006	Cum 2005	
United States	84	50	334	1,976	2,040	79	172	334	6,711	6,463	442	802	3,204	31,532	21,295	
New England		1	24	30	176	2	4	17	157	159	16	39	144	1,147	4,043	
Connecticut Maine [†]	<u> </u>	0 0	7 2	U 8	74 N	1	0 0	11 2	34 7	34 1	U	0 5	58 20	U 151	1,196 240	
Massachusetts New Hampshire	_	0 0	6 0	_	76	1	2 0	6 2	97 10	98 13	10	0 7	54 47	94 376	1,816 231	
Rhode Island	_	0	11	10	17	_	0	6	7	12		0	0	_	_	
Vermont [†]	_	0	2	12	9		0	1	2	1	6	12	50	526	560	
Mid. Atlantic New Jersey	1 N	3 0	15 0	125 N	169 N	13	21 3	35 7	855 128	801 111	84	103 0	183 0	3,659	3,605	
New York (Upstate) New York City	 U	1 0	10 0	44 U	65 U	5 4	2 10	14 23	117 410	62 481	_	0 0	0 0	_	_	
Pennsylvania	1	2	9	81	104	4	5	9	200	147	84	103	183	3,659	3,605	
E.N. Central Illinois	7	11 0	41 3	442 15	510 26	8 1	17 8	38 23	678 313	700 393	121	237 2	587 7	11,255 68	4,425 78	
Indiana	1	2	21	117	160	2	1	4	68	51	_	0	475	475	251	
Michigan Ohio	6	0 6	4 32	17 293	33 291	1 4	2 4	19 8	90 159	63 167	33 88	102 93	174 420	3,266 6,816	2,637 1,118	
Wisconsin	N	0	0	Ν	Ν	—	1	4	48	26	—	12	52	630	341	
W.N. Central Iowa	62 N	1 0	191 0	96 N	34 N	1	5 0	10 2	194 11	195 8	7 N	23 0	84 0	1,108 N	350 N	
Kansas	N	0	0	N	Ν	—	0	2	16	15	_	0	8	20	_	
Minnesota Missouri	60 2	0 1	191 3	60 35	27	_	0 3	3 8	21 130	57 110	7	0 19	0 82	1,006	242	
Nebraska† North Dakota	_	0	0 1	_	2 2	_	0 0	1 1	3 1	4	_	0 0	0 25	44	20	
South Dakota	—	Ő	1	1	3	1	Ő	3	12	1	_	1	12	38	88	
S. Atlantic Delaware	13	26 0	53 2	1,035	835 1	25	42 0	186 2	1,619 16	1,584 9	22	90 1	860 5	3,328 52	1,646 25	
District of Columbia	1	0	3	23	13	5	2	9	101	86	2	0	5	30	28	
Florida Georgia	10 2	14 7	36 29	574 343	460 265	10 1	15 7	29 147	575 277	537 332	_	0 0	0 0	_	_	
Maryland† North Carolina	N	0 0	0 0	N	N	1 5	5 5	19 17	229 229	247 205	_	0 0	0 0	_	_	
South Carolina [†]	—	0	0	—	—	1	1	7	54	58	10	15	53	809	439	
Virginia† West Virginia	N	0 1	0 14	N 95	N 96	1 1	3 0	12 1	133 5	107 3	10	30 26	812 70	1,287 1,150	350 804	
E.S. Central		3	13	152	143	7	13	25	547	356	—	1	70	91	95	
Alabama [†] Kentucky	<u>N</u>	0 0	0 5	N 29	N 26	1	4 1	19 8	238 56	115 36	N	1 0	70 0	90 N	95 N	
Mississippi Tennessee†	_	0 3	0 13	123	1 116	6	0 5	6 13	47 206	39 166	N	0 0	1 0	1 N	N	
W.S. Central	_	0	4	17	99	12	27	43	1,153	953	134	183	1,757	8,838	5,089	
Arkansas Louisiana	_	0 0	3 4	12 5	12 87	1 10	1 4	5 17	60 190	40 199	_	7 0	110 8	590 44	110	
Oklahoma	Ν	0	0	N	N	1	1	6	57	29	_	0	0	—	_	
Texas [†] Mountain	N 1	0 1	0 8	N 79	N 74	 10	21 7	36 25	846 322	685 332	134 58	170 54	1,647 138	8,204 2,106	4,979 2,042	
Arizona	N	0	0	N	Ν	7	3	16	144	132		0	0	´ —	_	
Colorado Idaho†	N N	0 0	0 0	N N	N N	_	1 0	3 1	32 2	36 20	35	33 0	76 0	1,152	1,408	
Montana Nevada†	—	0 0	1 3	4	29	3	0 1	1 12	1 83	5 91	_	0 0	2 2	2 4	2	
New Mexico [†]	_	0	1	1	—		1	5	52	40	_	3	34	307	172	
Utah Wyoming	1	0 1	8 4	34 40	23 22	_	0 0	1 0	8	8	23	10 0	55 8	608 33	409 51	
Pacific	_	0	0	_	_	1	33	49	1,186	1,383	_	0	0	_	_	
Alaska California	N	0	0	N	N	_	0 28	4 39	9 1,007	6 1,235	_	0 0	0 0	_	_	
Hawaii	—	0	Ō	_	—	_	0	2	15	9	N	0	0	N	Ν	
Oregon [†] Washington	N N	0 0	0 0	N N	N N	1	0 3	6 10	13 142	26 107	N N	0 0	0 0	N N	N N	
American Samoa	—	0	0	—	—	U	0	0	U	U	U	0	0	U	U	
C.N.M.I. Guam	_	0 0	0 0	_	_	U 	0 0	0 0	U	U 3	U 	0 4	0 12	U	U 389	
Puerto Rico U.S. Virgin Islands	N	0	0 0	N	N	_	2 0	10 0	86	164	3	8 0	47 0	284	554	
		0	v				v	0				0	0			

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending October 7, 2006, and October 8, 2005 (40th Week)*

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.

U: Unavailable. Cum: Cumulative year-to-o
 * Incidence data for reporting year 2006 is provisional.
 Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

(40th Week)*		West Nile virus disease [†]													
			Neuroinva	sive	west mile v	Non-neuroinvasive									
			vious			_			vious						
Reporting area	Current week	52 w	eeks Max	Cum 2006	Cum 2005	C	Current week	<u>52 w</u> Med	reeks Max	Cum 2006	Cum 2005				
United States	_	1	161	1,119	1,246		_	1	344	1,895	1,653				
New England	_	0	3	8	9		_	0	2	3	4				
Connecticut Maine§	_	0	2 0	6	4		_	0 0	1 0	2	2				
Vassachusetts	_	Ő	1	2	4		_	0	1	1	2				
New Hampshire	—	0	0	_	_		—	0	0	_	—				
Rhode Island /ermont [§]	_	0	0 0	_	1		_	0 0	0 0	_	_				
Vid. Atlantic	_	0	6	16	47		_	0	3	6	22				
New Jersey	_	0	2	2	3		_	0	1	2	3				
New York (Upstate)	—	0 0	1 4	7	19		_	0 0	1 2	3	5				
New York City Pennsylvania	_	0	4	7	11 14		_	0	2	3	3 11				
E.N. Central	_	0	35	188	255		_	0	18	77	153				
Illinois	—	0	21	111	136		_	0	16	55	114				
Indiana Michigan	—	0	4 7	11	10 54		—	0 0	2	5 2	11 8				
Michigan Ohio	_	0 0	11	29 27	54 45		_	0	1 3	2 7	8 14				
Visconsin	—	õ	2	10	10		—	õ	2	8	6				
W.N. Central	_	0	31	191	161		_	0	73	374	461				
owa Kansas	_	0	2 3	17 14	13 13		—	0 0	4 3	12 10	23 N				
Kansas Minnesota	_	0	6	14 29	13		_	0	3	10 34	N 27				
Missouri	—	0	12	41	17		_	0	3	9	13				
Nebraska§ North Dakota	_	0	7 5	33 20	53 12		_	0 0	24 27	123 115	131 74				
South Dakota	_	0	7	37	35		_	0	22	71	193				
S. Atlantic	_	0	3	8	29		_	0	3	6	26				
Delaware	_	0	0	_	1		_	0	1	_					
District of Columbia Florida	_	0 0	0 2	3	3 8		_	0 0	1 0	1	1 11				
Georgia	_	Ő	1	2	7		_	ŏ	2	4	10				
Maryland [§]	—	0	1	2	4		—	0	1	1	1				
North Carolina South Carolina§	_	0 0	0 1	_	2 4		_	0 0	0 0	_	2				
Virginia§	_	0	Ö	_	_		_	õ	0	_	1				
West Virginia	—	0	1	1	—		Ν	0	0	N	N				
E.S. Central	—	0	12	86	62		—	0	15	81	36				
Alabama [§] Kentucky	_	0 0	1 1	4 3	6 4		_	0 0	2 1	1	2				
Mississippi	_	Ő	9	72	38		_	0	15	79	31				
Tennessee§	—	0	3	7	14		—	0	1	1	3				
W.S. Central	—	1	55	283	249		—	0	25	147	145				
Arkansas Louisiana	_	0 0	4 14	21 66	12 106		_	0 0	2 8	5 49	15 53				
Oklahoma	_	0	6	21	16		_	0	3	12	11				
Texas§	—	0	35	175	115		—	0	14	81	66				
Mountain	_	0	59	270	134		_	0	196	993	228				
Arizona Colorado	_	0 0	4 10	15 54	44 20		_	0 0	5 43	14 219	49 85				
ldaho§	_	0	29	94	3		_	0	128	542	10				
Montana	—	0	3 9	10 34	8 14		—	0 0	7 13	19 73	17 17				
Nevada§ New Mexico§	_	0	9	34 1	14		_	0	13	73	17				
Utah	—	0	8	48	21		_	0	17	88	31				
Wyoming	—	0	7	14	5		_	0	7	36	6				
Pacific	—	0	15	69	300		_	0	45	208	578				
Alaska California	_	0 0	0 15	65	299		_	0 0	0 33	164	572				
Hawaii	—	0	0	—	_		—	0	0	_	—				
Oregon [§] Washington	_	0	2 0	4	1		_	0 0	12 2	42 2	6				
American Samoa	 U	0	0	 U	U		U	0	0	2 U	 U				
C.N.M.I.	U	0	0	U	U		U	0	0	U	U				
Guam	_	0	0	_	_		_	0	0	_	_				
Puerto Rico U.S. Virgin Islands	_	0 0	0 0	_	_		_	0 0	0 0	_	_				
o.o. virgin islanus		U	U	_			_	0	U	_	_				

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending October 7, 2006, and October 8, 2005 (40th Week)*

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.

 ¹ Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Zoonotic, Vector-Borne, and Enteric Diseases (proposed) (ArboNET § Surveillance). § Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

TABLE III. Deaths in 122 U.S. cities,* week ending October 7, 2006 (40th Week)

TABLE III. Deaths				y age (ye		.,.0			All causes, by age (years)						
Reporting Area	All Ages	<u>≥</u> 65	45-64	25-44	1-24	<1	P&l⁺ Total	Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	P&I [†] Total
New England	516	358	115	25	14	4	41	S. Atlantic	988	612	245	75	31	25	65
Boston, MA	124	79	27	7	8	3	9	Atlanta, GA	63	40	18	3	2		2
Bridgeport, CT	35	21	12	2	—	—	1	Baltimore, MD	116	64	32	10	7	3	12
Cambridge, MA	7	5	2	—	—	—	—	Charlotte, NC	119	73	23	12	7	4	14
Fall River, MA	27	22	4	_	1	_	1	Jacksonville, FL	114	64	30	11	6	3	6
Hartford, CT	48	36	11	_	1	_	6	Miami, FL	97	69	21	4	_	3	11
Lowell, MA	25	18 10	5 3	1	1	_	1 2	Norfolk, VA	51	29 25	11 14	6	2	3	2 3
Lynn, MA New Bedford, MA	13 19	10	3 5	_	_	_		Richmond, VA Savannah, GA	42 50	25 31	14	3 5	_	1	2
New Haven, CT	44	29	11	3	_	1	6	St. Petersburg, FL	47	32	8	3	1	3	5
Providence, RI	60	43	12	4	1		7	Tampa, FL	172	110	42	14	4	2	5
Somerville, MA	5	3	2	_	_	_	_	Washington, D.C.	100	62	30	3	2	3	1
Springfield, MA	30	21	5	4	_	_	_	Wilmington, DE	17	13	3	1	_	_	2
Waterbury, CT	34	24	6	3	1	_	4	E.S. Central	874	542	235	62	18	17	63
Worcester, MA	45	33	10	1	1	—	4	Birmingham, AL	173	106	235 47	13	2	5	14
Mid. Atlantic	1,995	1,362	436	129	36	31	108	Chattanooga, TN	58	39	11	5	1	2	3
Albany, NY	40	26	10	1	3	_	4	Knoxville, TN	101	70	24	6	_	1	5
Allentown, PA	25	17	6	1	_	1	1	Lexington, KY	94	65	22	3	2	2	6
Buffalo, NY	62	40	17	3	2	_	6	Memphis, TN	137	82	33	14	3	5	13
Camden, NJ	19	8	7	3	_	1	_	Mobile, AL	91	59	18	11	3	_	6
Elizabeth, NJ	10	8	2	—	—	—	—	Montgomery, AL	65	37	24	3	1	—	7
Erie, PA	38	31	5	1	_	1	3	Nashville, TN	155	84	56	7	6	2	9
Jersey City, NJ	19	12	7				2	W.S. Central	1,239	775	307	93	29	35	47
New York City, NY	1,024	705	219	69	16	14	41	Austin, TX	77	46	22	6	2	1	1
Newark, NJ Paterson, NJ	49 26	19 15	15 8	8 2	2	5 1	1	Baton Rouge, LA	64	35	21	6	2	_	1
Philadelphia, PA	321	201	89	21	8	2	19	Corpus Christi, TX	48	35	10	3	_	_	2
Pittsburgh, PA§	24	18	4	1		1		Dallas, TX	181	103	43	14	3	18	9
Reading, PA	25	18	3	2	1	1	2	El Paso, TX	74	53	13	5	2	1	3
Rochester, NY	119	85	25	6	1	2	10	Fort Worth, TX	94	63	21	3	2	5	2
Schenectady, NY	21	16	1	4	_	_	2	Houston, TX	344	199	99	29	11	6	11
Scranton, PA	25	23	2	_	_	_	2	Little Rock, AR New Orleans, LA ¹	71 U	41 U	20 U	4 U	3 U	3 U	
Syracuse, NY	85	71	7	3	2	2	8	San Antonio, TX	141	90	38	11	1	1	8
Trenton, NJ	21	14	6	1	—	—		Shreveport, LA	55	90 41	9	3	2	_	8
Utica, NY	14	12	1	1		_	2	Tulsa, OK	90	69	11	9	1	_	2
Yonkers, NY	28	23	2	2	1	_	4								
E.N. Central	1,967	1,314	436	134	33	50	136	Mountain Albuquerque, NM	966 133	611 86	233 38	68 7	27 2	27	60 7
Akron, OH	48	28	13	2	1	4	4	Boise, ID	63	41	12	5	2	3	4
Canton, OH	25	19	6				3	Colorado Springs, CO		40	6	2	_	3	2
Chicago, IL	330	199	72	34	13	12	25	Denver, CO	81	48	27	3	2	1	4
Cincinnati, OH	101	64	27	5	1	4	10	Las Vegas, NV	244	164	57	16	4	3	17
Cleveland, OH Columbus, OH	232 196	169 144	47 33	13 11	2	3 6	7 22	Ogden, UT	24	16	4	3	1	_	1
Dayton, OH	1190	84	30	3	1	1	8	Phoenix, AZ	189	92	54	26	10	7	14
Detroit, MI	152	75	53	19	2	3	11	Pueblo, CO	25	20	4	1			2
Evansville, IN	49	38	7	3	1	_	2	Salt Like City, UT	156	104	31	5	6	10	9
Fort Wayne, IN	36	20	12	3	1		1	Tucson, AZ	U	U	U	U	U	U	U
Gary, IN	11	4	4	2	1	—	_	Pacific	1,602	1,096	338	93	42	33	117
Grand Rapids, MI	45	29	11	2	_	3	5	Berkeley, CA	14	11	2	_	_	1	3
Indianapolis, IN	187	124	47	10	2	4	8	Fresno, CA	152	97	35	11	4	5	5
Lansing, MI	44	34	9	1	_	_	2	Glendale, CA	8	6	2	_	_		3
Milwaukee, WI	95	71	13	6	2	3	3	Honolulu, HI	66	52	8	3	2	1	5
Peoria, IL Rockford, IL	36 54	24 41	9 8	1 2	1 2	1	3 4	Long Beach, CA Los Angeles, CA	52	40	6 55	3 19	2	1 6	8
South Bend, IN	54 52	31	11	2 5	1	4	4	Pasadena, CA	211 27	120 18	55 8	19	11	0	16 1
Toledo, OH	104	75	16	10	2	1	11	Portland, OR	130	90	29	7	3	1	5
Youngstown, OH	51	41	8	2	_		4	Sacramento, CA	236	165	51	9	6	5	15
8					10	00	04	San Diego, CA	170	118	34	13	2	3	15
W.N. Central	566	362	127	32	19	26	34	San Francisco, CA	104	79	21	3	_	1	17
Des Moines, IA Duluth, MN	53 24	36 19	16 5	1	_	_	3 2	San Jose, CA	161	115	28	11	3	4	11
Kansas City, KS	24 26	19	5 5	1	2	_	2 1	Santa Cruz, CA	25	17	4	1	2	1	1
Kansas City, MO	107	70	24	5	2	6	6	Seattle, WA	89	54	21	7	5	2	5
Lincoln, NE	34	21	10	3		_	2	Spokane, WA	65	47	13	2	1	2	4
Minneapolis, MN	53	34	7	7	3	2	4	Tacoma, WA	92	67	21	3	1	—	3
Omaha, NE	72	51	11	4	2	4	5	Total	10,713**	7,032	2,472	711	249	248	671
St. Louis, MO	78	35	24	7	5	7	4		, -	, -			-	-	
St. Paul, MN	59	41	12	1	3	2	3								
Wichita, KS	60	37	13	3	2	5	4								
	No ropor														

U: Unavailable.

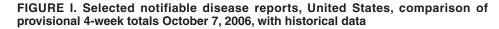
 $\frac{1}{2}$: Unavailable. -:No reported cases. * Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of \geq 100,000. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included. [†] Pneumonia and influenza.

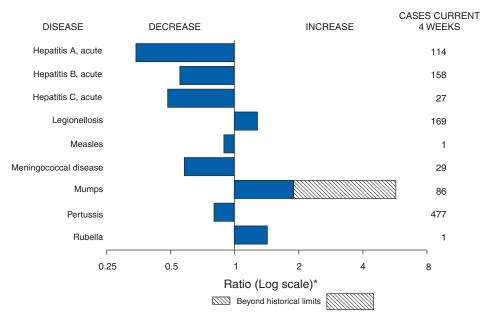
¹Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks. ¹Because of Hurricane Katrina, weekly reporting of deaths has been temporarily disrupted. ** Total includes unknown ages.

TABLE IV. Provisional cases of selected notifiable diseases,* United States, quarter ending September 30, 2006 (39th Week)

States, quarter ending September 30, 2006 (39th Week) Tuberculosis											
			/ious								
Demonstration and a	Current		arters	Cum	Cum						
Reporting area United States	quarter	Min	Max	2006	2005						
	3,090	2,384	3,589	8,100	9,661						
New England	90	60	167	224	278						
Connecticut	22	11	35	52	60						
Maine	3	2	4	8	13						
Massachusetts	53	31	113	142	161						
New Hampshire	5	0	5	9	4						
Rhode Island	7	0	12	10	35						
Vermont	—	0	3	3	5						
Mid. Atlantic	505	472	605	1,501	1,491						
New Jersey	137	101	137	370	361						
New York (Upstate)	61	50	110	172	192						
New York City	250	232	269	719	715						
Pennsylvania	57	57	102	240	223						
E.N. Central	284	219	371	815	948						
Illinois	120	91	158	369	453						
Indiana	39	24	40	91	106						
Michigan	44	30	93	129	153						
Ohio	68	54	83	180	177						
Wisconsin	13	13	19	46	59						
W.N. Central Iowa Kansas Minnesota Missouri Nebraska North Dakota South Dakota	131 5 28 62 30 6 	95 4 15 34 15 1 0 0	142 19 31 62 38 6 4 5	331 18 80 150 68 10 5	348 36 143 70 40 2 11						
S. Atlantic	608	369	839	1,549	2,041						
Delaware	9	3	9	18	19						
District of Columbia	18	10	18	48	42						
Florida	247	177	354	646	740						
Georgia	64	8	131	203	418						
Maryland	48	26	79	153	212						
North Carolina	100	57	126	231	203						
South Carolina	33	6	47	47	164						
Virginia	83	49	131	188	224						
West Virginia	6	4	7	15	19						
E.S. Central	147	147	211	473	531						
Alabama	44	44	52	145	168						
Kentucky	21	12	43	59	81						
Mississippi	23	23	36	82	67						
Tennessee	59	59	84	187	215						
W.S. Central	410	136	462	1,008	1,338						
Arkansas	26	19	37	82	79						
Louisiana		0	0								
Oklahoma	27	23	55	105	102						
Texas	357	76	388	821	1,157						
Mountain Arizona Colorado Idaho Montana Nevada New Mexico Utah Wyoming	139 90 3 — 21 13 11 1	65 25 0 0 9 7 4 0	193 115 34 7 2 24 13 11 1	296 176 17 — 45 31 25 2	402 166 67 16 8 88 32 25 						
Pacific	776	528	776	1,903	2,284						
Alaska	12	11	19	42	46						
California	652	446	652	1,578	1,891						
Hawaii	43	18	43	92	87						
Oregon	—	0	28	—	75						
Washington	69	45	77	191	185						
American Samoa C.N.M.I. Guam Puerto Rico U.S. Virgin Islands	U 62 	0 0 0 0	2 0 8 62 0	U 79 	U 55 76						

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: Not notifiable. Cum: Cumulative year-to-date counts. Min: Minimum. Max: Maximum. * AIDS and HIV/AIDS data are not updated for this quarter because of upgrading of the national HIV/AIDS surveillance data management system.





* 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.

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