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Performance of Rapid Influenza Diagnostic Tests During Two School Outbreaks of 2009 Pandemic Influenza A (H1N1) Virus Infection – Connecticut, 2009

During May 2009, a few weeks after 2009 pandemic influenza A (H1N1) infection was first detected in the United States (1), outbreaks among students from two schools were detected in Greenwich, Connecticut. Staff members from Greenwich Hospital and the Connecticut Department of Public Health collected data on symptoms for 63 patients and submitted nasopharyngeal washings for testing using a rapid influenza diagnostic test (RIDT) for influenza A and B and real-time reverse transcription-polymerase chain reaction (rRT-PCR) assay, thereby affording an opportunity to assess the field performance of the RIDT. A total of 49 patients had infections with pandemic influenza A (H1N1) confirmed by rRT-PCR. This report summarizes the findings from this performance assessment, which indicated that, compared with rRT-PCR, the sensitivity of the RIDT for detecting infection in patients with 2009 pandemic influenza A (H1N1) was 47%, and the specificity was 86%. Sensitivity and specificity did not vary substantially by the presence or absence of CDC-defined influenza-like illness (ILI) or by time from symptom onset to specimen acquisition. In this group of patients, although positive RIDT results performed well in predicting confirmed infection with pandemic H1N1 virus (positive predictive value: 92%), negative tests did not accurately predict the absence of infection (negative predictive value: 32%). These results affirm recent CDC recommendations against using negative RIDT results for management of patients with possible 2009 pandemic influenza A (H1N1) infection (2).

During April 29–May 1, 2009, 78 students from a private school (school A) near Greenwich, Connecticut, participated in a class trip to Pennsylvania. Several students became sick with a respiratory illness. Because infection with 2009 pandemic influenza A (H1N1) was suspected, upon returning home, 11 of the students, a sibling, and two other students went to the Greenwich Hospital for outpatient influenza testing and treatment.

During May 18–20, 133 students and eight teachers from a public school (school B) in Greenwich traveled to a camp in Connecticut. Among these students, 36 visited the camp infirmary with fever, headache, or fatigue. The Greenwich Health Department asked physicians at the hospital to assist with testing the students for pandemic H1N1. A total of 67 students and staff from school B became ill, and 49 of these patients went to the hospital for influenza testing.

A total of 63 patients (14 students from school A and 49 students and staff from school B) were tested for influenza at the hospital. A standard symptom survey was completed by a physician for each patient after which a nasopharyngeal washing was performed by an experienced respiratory therapist trained in the procedure. All samples were placed in viral transport media and sent to the Connecticut Department of Public Health laboratory for influenza A and B was performed concurrently at the hospital laboratory using the Remel Xpect Flu A&B test (Remel Products, Lenexa, Kansas) according to

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manufacturer's instructions (3). Although the number of ill persons who eventually received antiviral therapy is unknown, all nasopharyngeal washings were obtained before initiation of therapy.

Of the 63 patients tested by RIDT, 49 patients, 11 (79%) from school A and 38 (78%) from school B, were found to have 2009 pandemic influenza A (H1N1) infection by rRT-PCR (Figure). Of the 49 patients with confirmed infection, 23 (47%) tested positive (eight from school A and 15 from school B) and 26 (53%) tested negative for 2009 pandemic influenza A (H1N1) by RIDT. Among 11 patients with positive rRT-PCR tests from school A and 38 from school B, the numbers of positive RIDT tests were 8 (73%) and 15 (39%) respectively.

Among the 14 patient samples from both schools that tested negative by rRT-PCR, three were from students at school A, and 11 were from school B. Of the 14 rRT-PCR negative specimens, two tested positive by RIDT (one from school A and one from school B). The overall sensitivity of the RIDT was 23 of 49 (47%), and the specificity was 12 of 14 (86%). The positive predictive value was 23 of 25 (92%), and the negative predictive value was 12 of 38 (32%).

The schools did not differ significantly with respect to percentage of patients with confirmed pandemic H1N1 by rRT-PCR, severity of symptoms, interval between the onset of symptoms and collection of specimens for testing, or overall RIDT positivity rate. Among all the patients tested by RIDT, no significant differences between true positives and false negatives were seen with respect to ILI.* In RIDT positive and RIDT negative patients with pandemic H1N1, the median interval from symptom onset to specimen collection was 36 hours. Of the 34 patients with washings obtained \leq 36 hours from the onset of symptoms, 16 (47%) were RIDT positive; of the 15 patients with washings collected after 36 hours of symptoms, seven (47%) were positive. RIDT test performance was assessed for patients with and without CDC-defined ILI (Table). The sensitivity and specificity were approximately the same for the two groups (48% versus 44% and 88% versus 83%, respectively).

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^{*} CDC ILI surveillance case definition: fever (≥100°F [≥37.8°C]), plus cough, sore throat, or both in the absence of another known cause of illness.



FIGURE. Number of confirmed* cases of 2009 pandemic influenza A (H1N1) virus infections after school trips, by school, date of hospital visit, and result of rapid influenza diagnostic test[†] — Connecticut, May 2009

* By real-time reverse transcription–polymerase chain reaction assay; all patients tested negative for seasonal influenza. † Remel Xpect Flu A&B test (Remel Products, Lenexa, Kansas).

TABLE. Performance of a rapid influenza detection test (RIDT)* in patients with suspected and confirmed [†] 2009 pandemic influenze	za
A (H1N1) virus infection, by clinical syndrome consistent with CDC-defined influenza-like illness (ILI) [§] — Connecticut, 2009	

		rRT-PCR positive		rRT-PCR	negative				
	Total	RIDT positive	RIDT negative	RIDT positive	RIDT negative	Sensitivity %	Specificity %	PPV [¶] %	NPV** %
Overall	63	23	26	2	12	47	86	92	32
CDC-defined ILI**	48	19	21	1	7	48	88	95	25
No CDC-defined ILI	15	4	5	1	5	44	83	80	50

* Remel Xpect Flu A&B test (Remel Products, Lenexa, Kansas).

[†] By real-time reverse transcription-polymerase chain reaction (rRT-PCR); all patients tested negative for seasonal influenza.

§ CDC ILI surveillance case definition: fever (≥100°F [≥37.8°C]) plus cough, sore throat, or both in the absence of another known cause of illness.

[¶] Positive predictive value.

** Negative predictive value.

Editorial Note: When cases of 2009 pandemic influenza A (H1N1) began appearing in the United States in April 2009, several RIDTs had been in common use in the United States as point-of-care tests for seasonal influenza, but the performance of these tests in patients infected with 2009 pandemic influenza A (H1N1) virus was unknown. CDC has since reported varying sensitivities of RIDTs in retrospective analyses of rRT-PCR positive respiratory samples, from 40%–69%. In these

analyses, RIDT sensitivity was positively associated with the titer of virus in the sample (4).

The analysis in this report of pandemic H1N1 cases at two schools determined that the RIDT used detected less than half the cases confirmed by rRT-PCR. The low sensitivity and low negative predictive value of the test during these outbreaks highlight the limitations of using this test alone to establish diagnosis and aid clinical management. These results affirm current recommendations not to use negative RIDT results to rule out pandemic H1N1 or to make infection control decisions (2).

Rapid tests differ in their sensitivity and specificity for detecting seasonal influenza in respiratory specimens but generally have low to moderate sensitivity compared with viral culture or rRT-PCR. Previous RIDT studies have described the performance of the QuickVue Influenza A+B test (Quidel Corporation, San Diego, California) for detecting seasonal influenza in three different populations during 2008. Sensitivity when compared with rRT-PCR was low for all populations (median: 27%; range: 19%–32%) (5).

The RIDT used in the current study has a reported sensitivity of 92.5% and a specificity of 100% for the diagnosis of seasonal influenza A by nasopharyngeal wash (3). This investigation yielded much lower sensitivity (47%) and specificity (86%) in patients having confirmed infection with 2009 pandemic influenza A (H1N1) virus.

The findings in this report are comparable to recently reported observations of low performance of RIDTs in patients with pandemic H1N1. In a report of hospitalized patients in California, rapid antigen test results were positive in 67% of cases of pandemic H1N1 tested (6). In an assessment of rapid testing compared with rRT-PCR conducted on 6,090 patient samples from the New York City area, the sensitivity and specificity for the detection of 2009 pandemic influenza A (H1N1) virus by rapid antigen testing, using the BinaxNOW Influenza A&B test (Binax, Inc., Scarborough, Maine) and the 3M Rapid Detection Flu A+B test (3M, St. Paul, Minnesota) were 17.8% and 93.6% respectively (7). A recent report from the Naval Health Research Center described screening 3,066 clinical samples from service personal with influenza-like illness; of those screened, 767 rapid test results by QuickVue Influenza A+B test were available for comparison with rRT-PCR results (8). Of 39 patients with pandemic H1N1, 20 were RIDT positive, with a 51% sensitivity; for seasonal influenza A the sensitivity was 63% for H1N1 and 31% for H3N2. Specificity was 99% for all three subtypes when compared with rRT-PCR.

The results of these studies and the findings in this report affirm that a negative result for this rapid test does not rule out 2009 pandemic influenza A (H1N1) virus infection in an individual with symptoms consistent with influenza. Factors that might decrease the performance of rapid influenza antigen tests include improper specimen collection, not testing the recommended clinical sample (e.g., nasal versus nasopharyngeal swab), quality of the specimen, prolonged time from illness onset to specimen collection (because viral shedding decreases over time), and improper handling and storage of the specimen before testing. The reason for the suboptimal detection of 2009 pandemic influenza A (H1N1) by the RIDT used in this study was not specifically determined but did not appear to be related to differences in the interval (median: 36 hours for both groups) from onset of symptoms to specimen collection or to the severity of symptoms. As with all screening tests, the positive and negative predictive values of RIDTs are dependent on the prevalence of the disease in the population.

The findings in this report are subject to at least one limitation. The assessment involved a limited number of patients from two small outbreaks. The results should be viewed in this context. In other field situations (e.g., with other disease prevalences, collection and transport methods, or using other RIDTs), RIDTs might have different performance characteristics.

RIDTs can be an important tool for patient care during the normal influenza season because they usually provide results within 30 minutes. In addition, these tests can be used to make decisions about isolating or cohorting patients in health-care settings and recommending or restricting patient movements in outpatient settings. They might be especially important for hospitals limited by the expense of rRT-PCR and in identifying influenza during outbreaks in defined patient groups, such as those in schools or nursing homes. However, if used for management of patients with possible pandemic H1N1 virus infection, false-negative test reports might result in inappropriate exposure of susceptible persons to infected patients. Additional large studies to better characterize the performance of RIDTs for detection of infection in patients with pandemic H1N1 virus and improvements in rapid testing for pandemic H1N1 are needed.

Acknowledgment

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Anaplasmosis and Ehrlichiosis – Maine, 2008

Anaplasmosis and ehrlichiosis are rickettsial tickborne diseases that have had at least a twofold increase in prevalence in the United States since 2000 (1,2). Despite similar clinical presentations, the causative organisms are carried by different ticks with distinct geographic and ecologic associations (3). Surveillance efforts are complicated by ambiguous terminology and serologic testing with antibody cross-reactivity. Although anaplasmosis historically has been reported in Maine, ehrlichiosis has been reported infrequently. During 2007-2008, the number of physician-reported anaplasmosis cases nearly doubled in Maine, and ehrlichiosis cases increased more than fourfold. To examine this increase, the Maine Department of Health and Human Services (MDHHS) analyzed available data on tick burden and physician-reported cases of anaplasmosis and ehrlichiosis during 2000-2008. This report describes the results of that analysis, which indicated that Ixodes scapularis (the tick vector for Anaplasma phagocytophilum) was broadly distributed in Maine, whereas Amblyomma americanum (the tick vector for Erhlichia chaffeenisis) was scarce. Moreover, 95% of physician-reported ehrlichiosis cases lacked a concurrent serologic assessment to exclude anaplasmosis, suggesting that antibody cross-reactivity might have resulted in misclassification. In 2008, Maine modified case classification to enhance specificity; ehrlichiosis cases that lack a concurrent test for anaplasmosis are now classified as suspect rather than probable and therefore are not included in national surveillance summaries. The accuracy of case classification and surveillance can be improved by educating health-care providers regarding 1) the expected geographic distribution of tick vectors and 2) recommendations for confirmatory testing to distinguish between the causative organisms of anaplasmosis and ehrlichiosis.

In Maine, laboratories electronically report positive anaplasmosis and ehrlichiosis results to the health department (referred to as physician reported). Field epidemiology personnel follow up positive results by interviewing physicians and patients and by obtaining clinical, laboratory, and epidemiologic information required to complete the CDC tickborne rickettsial disease case report form.* MDHHS conducted a review of available data on tick burden in the state and reviewed the clinical and public health surveillance data for physician-reported human ehrlichiosis and anaplasmosis during 2000–2008. Maine classified cases according to Council of State and Territorial Epidemiologists (CSTE) case definition[†] with the exception that the 2008 cases were classified according to a modified ehrlichiosis case definition that had increased specificity.

Tick Surveillance Data

During 2000–2008, the Vector Borne Disease Laboratory of the Maine Medical Center Research Institute conducted active surveillance of ticks in Maine (through flagging and trapping) and passive surveillance (through receipt of ticks submitted by state residents through the mail) (4). A total of 5,089 *I. scapularis* were collected, but only 15 *A. americanum* ticks were detected. All life stages of *I. scapularis* (larvae, nymphs, and adults) were identified; the tick distribution increased and expanded along the southern coastline and up the river valleys, corresponding to areas of increasing settlement of human populations in this geographic distribution. During 2007–2008, Maine residents submitted 1,968 *I. scapularis* and only six *A. americanum*. The surveillance results suggested that *A. americanum*, the ehrlichiosis vector, had only a sparse and sporadic distribution in Maine.

Human Anaplasmosis Surveillance Data

During 2000–2008, a total of 45 cases of anaplasmosis cases were reported in Maine. Fifteen (33%) cases were confirmed, 30 (67%) were probable, and no suspect cases were reported (Tables 1 and 2). Among the 15 confirmed cases, three (20%) patients were diagnosed by demonstration in paired sera of a fourfold or greater increase in antibodies to A. phagocytophilum in acute versus convalescent samples; 12 (80%) patients were diagnosed by polmerase chain reaction (PCR) detection of A. phagocytophilum DNA, including two patients who also had positive single A. phagocytophilum serologic test. Among the 30 probable cases, 23 (77%) patients were diagnosed only by a single test for antibodies to A. phagocytophilum, including one (3%) patient who also had detection of morulae consistent with A. phagocytophilum on a blood smear. Seven (23%) patients were tested for antibodies to both A. phagocytophilum and E. chaffeensis, and all showed higher antibody titers to A. phagocytophilum. The median patient age among all confirmed and probable cases was 57 years (range: 21-89 years); 28 patients (62%) were males. Seventeen (38%) patients were hospitalized, and one (2%) patient died from renal failure relating to infection. Two (4%) patients were diagnosed with concurrent Lyme disease, and two (4%) with concurrent babesiosis. Reported anaplasmosis cases occurred during April-December; 30 (67%) of 45 patients had onset dates during May-September. Anaplasmosis was reported in six (38%) of 16 counties; the majority occurred in southern coastal Maine. One patient with confirmed anaplasmosis had traveled to New York, an anaplasmosis-endemic state, during the preceding month.

^{*}Available at http://www.cdc.gov/ncidod/dvrd/rmsf/case_rep_fm.pdf.

[†] Available at http://www.cdc.gov/ncphi/disss/nndss/casedef/ehrlichiosis_2008.htm.

Human Ehrlichiosis Surveillance Data

During 2000-2008, a total of 20 cases of ehrlichiosis were reported in Maine (Tables 1 and 2). The single confirmed case, which was diagnosed by PCR, occurred in a male aged 58 years who worked as an interstate truck driver; therefore, outof-state exposure to E. chaffeensis was possible. An additional 19 ehrlichiosis cases were reported during this same period (including six cases reported during 2005-2007 and 13 cases reported during 2008). All 19 cases were diagnosed by a single positive Ehrlichia serologic assay, and none had accompanying serologic tests to exclude anaplasmosis. Although all 13 cases reported in 2008 would have met the CSTE case definition for probable ehrlichiosis, beginning in that year, Maine had adopted a modified ehrlichiosis case definition to increase specificity; therefore, these 13 case were classified as suspect. Ten of the 20 cases were in persons who had either concurrent Lyme disease (seven persons) or babesiosis (three persons), which, like Anaplasma, are transmitted by I. scapularis.

2008 Classification of Ehrlichiosis Cases

Based on the lack of evidence for a sustained tick vector population in the state, lack of travel history among patients, and the cross-reactive serologic tests for ehrlichiosis and anaplasmosis, MDHHS implemented a new ehrlichiosis case classification strategy using a modified CSTE case definition in 2008 (5). Probable ehrlichiosis cases were defined as clinically compatible with one positive immunoglobulin G (IgG) serologic result for E. chaffeensis and either a concurrent lower titer serologic test for A. phagocytophilum or visualization of intracytoplasmic morulae in peripheral monocytes or macrophages. For cases having serologic reactivity to both agents, the higher antibody level was used to identify the most likely infection (5). Ehrlichiosis reports that did not meet this new more stringent probable case definition (i.e., those that were only tested for ehrlichiosis) were classified as suspect cases, which are excluded from national notifiable disease surveillance summaries.

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Editorial Note: The findings in this report underscore that the use of cross-reactive serologic assays, which test for ehrlichiosis alone in anaplasmosis-endemic areas, can result in an inaccurately high ehrlichiosis incidence and contribute to underrecognition of actual anaplasmosis cases. Serologic assays for *A. phagocytophilum* and *E. chaffeensis* have >50% cross reactiv-

TABLE 1. Number and percentage of anaplasmosis and ehrlichiosis cases*, by selected characteristics — Maine, 2000–2008

	Anaplasmo	sis (n = 45)	Ehrlichio	sis (n = 20)
Characteristic	No.	(%)	No.	(%)
Classification				
Confirmed	15	(33)	1	(5)
Probable	30	(67)	6	(30)
Suspect	0	_	13	(65)
Year				
2000	1	(2)	0	_
2001	1	(2)	0	_
2002	1	(2)	0	_
2003	1	(2)	0	_
2004	1	(2)	0	_
2005	5	(12)	1	(5)
2006	9	(20)	2	(10)
2007	9	(20)	3	(15)
2008	17	(38)	14	(70)
Sex				
Male	28	(62)	9	(45)
Female	17	(38)	11	(55)
Age group (yrs)				
<20	0	_	0	_
20–29	2	(4)	2	(10)
30–39	4	(9)	2	(10)
40–49	11	(24)	6	(30)
50–59	10	(22)	5	(25)
≥60	17	(38)	5	(25)
Unknown	1	(2)	0	_
Coinfections				
Lyme disease	2	(4)	7	(35)
Babesiosis	2	(4)	3	(15)
Outcome				
Hospitalized	17	(38)	2	(10)
Complications [†]	2	(4)	1	(5)
Death	1	(2)	0	

* Cases reported during 2000–2007 were classified based on Council of State and Territorial Epidemiologists (CSTE) case definitions (available at http://www.cdc.gov/ncphi/disss/nndss/casedef/ehrlichiosis_2008.htm). However, beginning in 2008, Maine modified the case definition to increase specificity regarding ehrlichiosis; reports with only one serologic test result for ehrlichiosis and no concurrent anaplasmosis test result were classified as suspect in Maine.

[†] Complications related to infection included renal failure, polymyositis, and meningitis.

ity; thus, differentiating between ehrlichiosis or anaplasmosis based on single serologic assay is not possible (6–8). In 2008, Maine classified 13 ehrlichiosis cases as suspect because they more likely represent infection with *A. phagocytophilum* given that tick data did not support a sustained ehrlichiosis vector in the state and confirmatory laboratory testing and supporting travel history for ehrlichiosis infection were lacking. The likelihood these suspect cases are anaplasmosis cases is further supported by the fact that 54% of suspect ehrlichiosis cases occurred in persons who had either concurrent Lyme disease or babesiosis, which, like *Anaplasma*, are transmitted by *I. scapularis*. Whether the emergence of anaplasmosis in Maine

			Anaplasmosis	(n = 45)	Ehrlichiosis (n = 20)						
Diagnostic test used	No.	(%)	Confirmed	Probable	Suspect	No.	(%)	Confirmed	Probable	Suspect	
Single serology [†]	22	(49)		22	_	19	(95)		6	13	
Single serology for both infections	7	(16)	_	7§		0	_	_	_	_	
Paired serology [¶]	3	(7)	3			0	_	_	_	_	
PCR**	10	(22)	10			1	(5)	1	_	_	
PCR + single serology	2	(4)	2			0		_	_	_	
Smear ^{††} + single serology	1	(2)	_	1		0	_		_	_	

TABLE 2. Number and percentage of anaplasmosis and ehrlichiosis cases*, by diagnostic test used and case classification — Maine, 2000–2008

* Cases reported during 2000–2007 were classified based on Council of State and Territorial Epidemiologists (CSTE) case definitions (available at http:// www.cdc.gov/ncphi/disss/nndss/casedef/ehrlichiosis_2008.htm). However, beginning in 2008, Maine modified the case definition to increase specificity regarding ehrlichiosis; reports with only one serologic test result for ehrlichiosis and no concurrent anaplasmosis test result were classified as suspect in Maine.

[†] Serum tested with Anaplasma phagocytophilum (for anaplasmosis) or Ehrlichia chaffeensis (for ehrlichiosis) antigen, but not both.

§ Seven patients were tested for antibodies to both A. phagocytophilum and E. chaffeensis concurrently, and all showed higher antibody titers to A. phagocytophilum.

¹ Diagnosed by demonstration in paired sera of a fourfold or greater increase in antibodies to *A. phagocytophilum* in acute versus convalescent samples. ** Polymerase chain reaction.

⁺⁺ Visualization of intracytoplasmic morulae in granulocytes for anaplasmosis or peripheral monocytes or macrophages for ehrlichiosis.

and nationwide is an actual increase in incidence or an increase in awareness and testing is unclear. Reports of anaplasmosis have increased threefold (from 351 cases in 2000 to 1,053 cases in 2008), and reports of ehrlichiosis have increased more than fourfold (from 200 cases in 2000 to approximately 800 cases in 2008) (*1*; CDC, unpublished data, 2009). Most cases of ehrlichiosis have been reported from the southern and south-central United States, corresponding to the geographic distribution of the tick vector, *A. americanum*. However, during 2008–2009, a concerning trend of increased ehrlichiosis case reports from some northern-area states, including Maine, has been noted (CDC, unpublished data, 2009). Possible explanations for this increase include expanding geographic ranges of the tick vector *A. americanum* or misclassification of cases.

Anaplasmosis, referred to as human granulocytic anaplasmosis, is caused by A. phagocytophilum. Before a taxonomic reorganization in 2001, this organism was called Ehrlichia phagocytophilum, and the infection was described as human granulocytic ehrlichiosis. *I. scapularis* (the black-legged tick), the vector for anaplasmosis, is reported commonly from northern and northeastern states. Ehrlichiosis, known as human monocytic ehrlichiosis, is caused by E. chaffeensis and is transmitted by A. americanum (the lone star tick). E. chaffeensis is commonly reported in the southern and south-central states, where the vector is common. Both anaplasmosis and ehrlichiosis are nationally notifiable diseases. In Maine, the vector A. americanum responsible for transmission of E. chaffeensis is not endemic. Conversely, A. phagocytophilum DNA has been detected in 16% of 94 I. scapularis ticks tested in 2008 (9). The fact that 95% of physician-reported ehrlichiosis cases lacked a concurrent serologic assessment to exclude anaplasmosis supports the likelihood that antibody cross-reactivity could have resulted in misclassification. One factor contributing to this misclassification might have been confusion among physicians regarding the recent change in terminology for *A. phagocytophilum* infection (from human granulocytic ehrlichiosis to anaplasmosis) and a lack of understanding of appropriate testing strategies. Since taxonomic changes were adopted in 2001, the term "anaplasmosis" has gradually replaced the term "human granulocytic ehrlichiosis" to describe human infections with *A. phagocytophilum*. However, some medical references and commercial test names still use the term "ehrlichiosis," which might cause confusion among physicians regarding the selection of appropriate diagnostic tests.

Health-care providers should assess clinical and ecologic features and, as indicated, include concurrent confirmatory testing for both anaplasmosis and ehrlichiosis or other tickborne diseases when evaluating patients with suspected tickborne illness. Compared with anaplasmosis patients, ehrlichiosis patients might have a higher potential for severe or fatal outcome, and a higher proportion (up to 30%) of ehrlichiosis patients have rash; thus, these diagnostic clues also can prompt physicians to request concurrent testing for ehrlichiosis (3). If serologic testing is selected to evaluate patients, serology should include 1) concurrent testing for both A. phagocytophilum and E. chaffeensis and 2) testing of paired acute and convalescent sera whenever possible. PCR is considered a confirmatory test and is the recommended diagnostic tool preferred over serology because it can differentiate between the two infections (4, 10). Patients with suspected anaplasmosis or ehrlichiosis should be treated promptly with doxycycline, without regard to initial serologic test results, because antibodies in the first week of illness frequently are not detected.

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Progress Toward Measles Control – African Region, 2001–2008

In 2001, the countries of the World Health Organization (WHO) African Region (AFR) became part of a global initiative with a goal of reducing the number of measles deaths by 50% by 2005, compared with 1999. Recommended strategies for measles mortality reduction included 1) increasing routine coverage for the first dose of measles-containing vaccine (MCV1) for all children, 2) providing a second opportunity for measles vaccination through supplemental immunization activities (SIAs), 3) improving measles case management, and 4) establishing case-based surveillance with laboratory confirmation of all suspected measles cases (1). Before introduction of MCV throughout AFR, approximately 1 million measles cases had been reported each year in the early 1980s (2). After strengthening measles-control activities, annual reported cases declined to an estimated 300,000-580,000 during the 1990s. This report summarizes the progress made during 2001–2008 toward improving measles control in AFR. During 2001–2008 estimated MCV1 coverage increased from 57% to 73%, SIAs vaccinated approximately 398 million children, and reported measles cases decreased by 93%, from 492,116 in 2001 to 32,278 in 2008. By 2005, global measles deaths had decreased by 60%, and the AFR goal had been achieved (*3*); AFR adopted a new goal to reduce deaths by 90%, compared with 2000, and that goal was achieved in 2006 (*3*,*4*). However, inaccuracies in reported vaccination coverage exist, surveillance is suboptimal, and measles outbreaks continue to occur in AFR countries. Further progress in measles control will require full implementation of recommended strategies, including validation of vaccination coverage.

Since the 1980s, AFR countries have reported measles vaccination coverage and the number of measles cases each year to the WHO African Regional Office (AFRO), using the WHO and United Nations Children's Fund (UNICEF) Joint Reporting Form. These data are collected through administrative reports from routine vaccination programs and SIAs and routine surveillance systems that provide aggregated case counts based on clinical diagnosis. Estimates of routine coverage with MCV1 are based on review of coverage data from administrative records, surveys, national reports, and consultation with local and regional experts. Coverage achieved during nationwide SIAs against measles are reported on the basis of the reported number of doses administered, divided by the target population.

In 1999, as part of the measles mortality reduction strategy, case-based surveillance with laboratory testing for all suspected measles cases was introduced with support from WHO AFRO. A suspected measles case is defined as 1) any person with generalized maculo-papular rash and fever plus cough or coryza or conjunctivitis or 2) any person in whom a clinician suspects measles. Each suspected measles case should be reported using an individual case-investigation form, and a blood specimen should be collected and sent to the laboratory for measles-specific immunoglobulin M testing. Laboratory confirmation of individual cases is discontinued after an outbreak has been confirmed as measles. An outbreak is confirmed when three or more measles laboratory-confirmed cases are detected in a health facility or district in 1 month; subsequent cases are confirmed by epidemiologic link. An epidemiologic link is defined as a suspected measles case that did not have a specimen collected for laboratory testing and is linked in person, place, and time to a laboratory-confirmed case (i.e., in a patient living in the same district or an adjacent district with a patient with laboratory-confirmed measles where a likelihood of transmission and onset of rash in the two patients within 30 days of each other exists) (5). Case-based surveillance data from AFR countries are shared regularly with WHO AFRO. Data quality is monitored using annualized performance

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indicators that include the 1) percentage of districts reporting one or more suspected case with a blood specimen (target: >80%) and 2) nonmeasles febrile rash illness rate (target: >2 cases per 100,000).

Routine Vaccination Activities

In AFR, MCV1 is administered through routine services to children at age 9 months. According to WHO and UNICEF estimates, AFR MCV1 coverage increased from 57% in 2001 to 73% in 2008 (Figure). In 2008, among the 46 AFR countries,* three (7%) had MCV1 coverage of <60%, 13 (28%) had coverage of 60%–69%, 11 (24%) had coverage of 70–79%, 10 (22%) had coverage of 80–89%, and nine (20%) had coverage of ≥90% (Table 1). As of 2008, five (10%) countries provided a second dose of MCV (MCV2) through routine services: South Africa and Swaziland reported MCV2 coverage of 70%, Lesotho reported MCV2 coverage of 80%, and Algeria and Seychelles reported MCV2 coverage of >95% in 2008.

SIA Results

SIAs provide a second opportunity for measles immunization to all children, including those not vaccinated with MCV1 and those previously vaccinated; approximately 15% of children vaccinated with a single dose at age 9 months will not develop immunity to measles. The SIA strategy generally consists of a one-time catch-up SIA, targeted to a wide age range, which aims to reduce susceptibility to measles in the population. This is followed by periodic follow-up SIAs targeting children born since the last SIA, thus reducing the accumulation of susceptible children in new birth cohorts.

Before 2000, seven (15%) AFR countries (Botswana, Lesotho, Malawi, Namibia, South Africa, Swaziland, and Zimbabwe) had completed a catch-up SIA, and Namibia and South Africa had completed a follow-up SIA (6). By the end of 2008, 43 AFR countries (all except Algeria, Mauritius, and Seychelles) had completed a catch-up SIA, and all but Comoros and Guinea-Bissau had completed at least one follow-up SIA (Table 2). During 2001–2008, approximately 398 million children were vaccinated during measles SIAs in AFR: 237 million (60%) during catch-up SIAs in 34 countries, and 161 million (40%) during follow-up SIAs in 39 countries (Table 2). Nine countries (Benin, Cameroon, Chad, the Democratic

FIGURE. Number of reported measles cases* and coverage with the first dose of measles-containing vaccine (MCV1) among children aged <1 year[†] — World Health Organization (WHO) African Region, 2001–2008



* N = 1.9 million. Confirmed cases of measles reported by member states to WHO and the United Nations Children's Fund (UNICEF) through the Joint Reporting Form.

[†] Data are from WHO and UNICEF measles vaccination coverage estimates; these estimates are based on reviews of surveys and national reports of administrative coverage. Administrative coverage is calculated by dividing the number of doses of vaccine administered through routine health services by the birth cohort of the previous year.

Republic of Congo, Ethiopia, Ghana, Niger, Nigeria, and Tanzania) conducted nationwide SIAs in phases covering different geographic areas implemented over ≥2 years.

Measles Surveillance

By December 2008, all AFR countries except Algeria, Comoros, Guinea Bissau, Mauritius, Sao Tome & Principe, and Seychelles had established measles case-based surveillance in accordance with the WHO AFRO measles surveillance guidelines (5). In 2008, of the 40 countries with case-based surveillance, 21 (53%) met the target of >80% of districts reporting one or more suspected cases; 24 (60%) had a nonmeasles febrile rash illness rate of >2 cases per 100,000 population; and 16 (40%) met both targets.

Monitoring Measles Incidence

Following implementation of the measles mortality reduction strategies during 2001–2008, including introduction of case-based measles surveillance, the number of reported measles cases decreased 93%, from 492,116 in 2001 to 32,278 in 2008 (Figure). Average annual measles incidence in AFR decreased 66%, from 50.2 per 100,000 population during 2001–2004 to 17.2 during 2005–2008 (Table 1). Despite this decrease, during 2005–2008, 14 countries[†] reported outbreaks. Outbreak field investigations conducted during 2003–2007 in South

^{*} Algeria, Angola, Benin, Botswana, Burkina Faso, Burundi, Cameroon, Cape Verde, Central African Republic, Chad, Comoros, Congo, Côte d'Ivoire, Democratic Republic of Congo, Equatorial Guinea, Eritrea, Ethiopia, Gabon, Gambia, Ghana, Guinea, Guinea-Bissau, Kenya, Lesotho, Liberia, Madagascar, Malawi, Mali, Mauritania, Mauritius, Mozambique, Namibia, Niger, Nigeria, Rwanda, Sao Tome and Principe, Senegal, Seychelles, Sierra Leone, South Africa, Swaziland, Togo, Uganda, United Republic of Tanzania, Zambia, and Zimbabwe.

Africa (1,676 cases, 2003–2005) (7), Kenya (2,544 cases, 2005–2007) (8), and Tanzania (1,533 cases, 2006–2007) (9) found that failure to vaccinate was the primary cause. In 2008, outbreaks also contributed to annual case counts in Burkina Faso (395), Cameroon (495), the Democratic Republic of Congo (12,461), Ethiopia (3,511), Niger (1,317), and Nigeria (9,960) (2).

Reported by: Countries in the World Health Organization African Region; Immunization and Vaccine Development, World Health Organization Regional Office for Africa. Dept of Immunization, Vaccines, and Biologicals, World Health Organization, Geneva, Switzerland. Global Immunization Div, National Center for Immunization and Respiratory Diseases, CDC.

Editorial Note: In 2008, after implementation of the measles mortality reduction strategy, routine measles vaccination coverage in AFR reached 73%, SIAs were conducted in nearly all AFR countries, and reported measles cases decreased to a historic low of 32,278. According to previously published WHO estimates, by 2006 AFR had achieved approximately 90% reduction in measles deaths, compared with 2000 (3). However, despite this progress, vaccination coverage reports remain imprecise, disease surveillance remains suboptimal, and outbreaks continue to occur, even in countries that reported implementation of all recommended components of the measles strategy. Available mathematical models likely overestimate the disease burden and underreporting of measles cases is common, even with high-performing surveillance systems; therefore, caution is recommended when drawing comparisons between reported incidence of measles and estimates of measles deaths generated from models.

SIAs are recommended to provide a second opportunity for immunization and increase the likelihood of vaccinating hard-to-reach children. SIA coverage usually is estimated by an administrative method relying on the reported number of vaccine doses administered and available target population denominator data, both of which often are imprecise. For example, during 2001–2008, several countries reported vaccinating >100% of children targets in SIAs. Improved methods for determining the actual target population size for SIAs are needed; reported coverage also should be routinely validated by independent surveys. In addition, detailed field investigations of outbreaks should be undertaken to identify post-SIA risk factors for measles, and help refine vaccination strategies.

The findings in this report are subject to at least two limitations. First, a change in measles surveillance methods might result in underestimates or overestimates of the disease burden over time. For example, in 1999, AFR countries routinely

TABLE 1. Routine measles vaccination coverage* and measles
incidence, [†] by country — World Health Organization (WHO)
African Region, 2001–2008

	% cove with firs measles (MC)	erage t dose vaccine /1)	Average annual measles incidence per 100,000 population				
Country	2001	2008	2001–2004	2005–2008			
WHO African Region	54	73	50.2	17.2			
Algeria	81	83	21.4	2.6			
Angola	72	79	37.0	3.3			
Benin	70	61	28.5	4.9			
Botswana	91	94	0.9	0.2			
Burkina Faso	54	75	18.0	1.3			
Burundi	76	84	4.6	3.3			
Cameroon	47	80	40.9	1.9			
Cape Verde	75	96	0.0	0.0			
Central African Republic	35	62	36.4	3.2			
Chad	26	23	160.4	5.0			
Comoros	70	76	0.0	40.4			
Congo	35	79	94.1	2.6			
Côte d'Ivoire	75	63	31.1	0.2			
Democratic Rep. of Congo	49	67	47.5	137.2			
Equatorial Guinea	51	51	64.9	16.7			
Eritrea	84	95	6.7	1.1			
Ethiopia	53	74	2.2	2.1			
Gabon	55	55	105.0	1.7			
Gambia	89	91	6.7	0.0			
Ghana	81	86	34.2	1.1			
Guinea	44	64	34.9	0.5			
Guinea-Bissau	72	76	89.7	0.2			
Kenva	73	90	9.4	3.2			
Lesotho	70	85	3.2	0.0			
Liberia	58	64	13.9	0.2			
Madagascar	57	81	176.8	0.0			
Malawi	82	88	2.9	0.6			
Mali	53	68	12.9	0.5			
Mauritania	58	65	96.3	1.4			
Mauritius	98	90	16.3	0.7			
Mozambique	74	77	66.8	15.6			
Namibia	58	73	25.9	0.3			
Niger	37	80	436.8	70			
Nigeria	35	62	72.9	21.9			
Bwanda	69	92	14.9	18			
Sao Tome & Principe	75	93	0.0	0.0			
Senegal	48	77	99.6	0.0			
Sevchelles	99	95	0.0	3.3			
Sierra Leone	50	60	10.0	0.5			
South Africa	69	62	18	0.0			
Swaziland	72	95	9.9	0.0			
Tanzania	83	88	14 1	6.2			
Τοαο	53	77	117	10			
Uganda	61	68	123.5	79			
Zambia	84	85	98.7	24			
Zimbabwe	73	66	3.7	1.8			

* WHO and United Nations Children's Fund (UNICEF) estimates of routine measles vaccination coverage are based on reviews of surveys and national reports of administrative coverage. Administrative coverage is calculated by dividing the number of doses of vaccine administered through routine health services by the birth cohort of the previous year.

[†] Measles incidence is calculated using confirmed measles cases reported by member states to WHO and UNICEF through the Joint Reporting Form and population estimates from: World population prospects: the 2008 revision, United Nations Population Division, available at http://esa.un.org/ unpp.

[†] Angola, Benin, Burkina Faso, Cameroon, Democratic Republic of Congo, Equatorial Guinea, Ethiopia, Kenya, Mali, Niger, Nigeria, South Africa, Tanzania, and Uganda.

Country				Children reached in targeted age group				
Country	Year	Target age group	Type of SIA*	No.	Administrative coverage [†] (%)			
Algeria	NA§	NA	NA	NA	NA			
Angola	2003	9 mos–14 vrs	Catch-up	7.226.105	95			
3	2006	9–59 mos	Follow-up	3.210.160	97			
Benin	2001	9 mos–14 vrs	Catch-up	950.780	>100¶			
	2003	9 mos–14 vrs	Catch-up	2.299.583	>100			
	2005	9–59 mos	Follow-up	1,137,163	>100			
	2008	9–59 mos	Follow-up	1.272.621	>100			
Botswana	2005	9–59 mos	Follow-up	179,202	99			
Burkina Faso	2001	9 mos_14 vrs	Catch-up	4 943 115	96			
Barrana r abo	2004	9–59 mos	Follow-up	2 882 208	>100			
	2007	9–59 mos	Follow-up	3 145 255	>100			
Burundi	2002	9 mos - 14 yrs	Catch-up	2 767 054	90			
Baranar	2002	9_59 mos	Follow-up	1 226 689	>100			
Cameroon	2000	9 mos_14 vrs	Catch-up	2 789 542	2100			
Cameroon	2001	9 mos - 14 yrs	Catch-up	4 570 817	90			
	2002	0 50 mos	Eollow up	1,240,041	90			
	2000	9–59 mos	Follow up	1,249,041	99			
Cana Varda	2007	9–59 mos	Follow-up	1,703,107	91			
Cape Verde	2005	9–59 mos	Follow-up	40,009	93			
Central Alfican Republic	2005	9 mos 14 yrs	Catch-up	1,103,303	91			
	2006	9 1105–14 yrs	Calch-up	515,950	90			
	2008	9–59 mos	Follow-up	683,302	>100			
Chad	2005	9 mos-14 yrs	Catch-up	1,641,896	80			
	2006	9 mos–14 yrs	Catch-up	2,735,760	>100			
-	2008	9–59 mos	Follow-up	1,782,689	96			
Comoros	2005	6 mos–14 yrs	Catch-up	109,815	99			
	2007	6 mos–14 yrs	Catch-up	231,263	81			
Congo	2004	9 mos–14 yrs	Catch-up	1,356,625	78			
	2007	9–59 mos	Follow-up	677,390	95			
Côte d'Ivoire	2005	9 mos–14 yrs	Catch-up	7,894,327	88			
	2008	9–59 mos	Follow-up	3,082,438	95			
Democratic Republic of the Congo	2002	9 mos–14 yrs	Catch-up	5,554,824	96			
	2004	6 mos–14 yrs	Catch-up	8,604,754	86			
	2005	6 mos–14 yrs	Catch-up	6,957,653	89			
	2006	9 mos–14 yrs	Catch-up	6,970,229	**			
	2006	9–59 mos	Follow-up	5,723,858	99			
	2007	9–59 mos	Follow-up	3,768,794	>100			
	2008	9–59 mos	Follow-up	2,811,092	99			
Equatorial Guinea	2005	9 mos–14 yrs	Catch-up	119,462	44			
Eritrea	2003	9 mos–14 yrs	Catch-up	1,047,862	82			
	2006	9–59 mos	Follow-up	387,479	95			
Ethiopia	2003	9 mos–14 yrs	Catch-up	5,101,001	91			
•	2004	6 mos–14 yrs	Catch-up	7,422,074	84			
	2005	6 mos–14 yrs	Catch-up	136,935	69			
	2005	9 – 59 mos	Follow-up	987.221	92			
	2006	9–59 mos	Follow-up	10.169.187	87			
	2007	6–59 mos	Follow-up	1.072.701	98			
	2008	6–59 mos	Follow-up	10.848.474	92			
Gabon	2004	9 mos-14 vrs	Catch-up	502 959	80			
Gabon	2007	9-59 mos	Follow-up	190 035	83			
Gambia	2003	9 mos_14 vrs	Catch-up	677830	00 02			
Gampia	2000	0_50 moo	Follow up	0/1,000	06			
Chana	2007	9-09 1105 0 mon 14 uro	Catch up	241,214 700 700	90			
Gilalia	2001	9 1105-14 yrs	Catch up	190,190	99 - 100			
	2002	9 1105-14 YIS	Eallow up	1,021,000 2 004 050	> 100			
	2000	9-09 mos	ronow-up	3,994,052	19			

See Table 2 footnotes on page 1041.

TABLE 2. Measles supplementary immunization activities (SIAs), by type and country — World Health Organization (WHO) African Region, 2001–2008

				Childre targete	Children reached in targeted age group				
Country	Year	Target age group	Type of SIA*	No.	Administrative coverage [†] (%)				
Guinea	2003	9 mos–14 yrs	Catch-up	3,202,848	98				
	2006	9–59 mos	Follow-up	1,707,633	97				
Guinea-Bissau	2006	6 mos–14 yrs	Catch-up	590,602	85				
Kenya	2002	9 mos–14 yrs	Catch-up	13,302,991	98				
-	2006	9–59 mos	Follow-up	5,260,241	>100				
Lesotho	2003	9–59 mos	Follow-up	178,522	87				
	2007	9–59 mos	Follow-up	196,490	92				
Liberia	2004	_	_ '	· —	_				
	2007	9–59 mos	Follow-up	629,676	97				
Madagascar	2004	9 mos–14 vrs	Catch-up	8.900.657	99				
	2007	9–59 mos	Follow-up	3.053.702	100				
Malawi	2002	9–59 mos	Follow-up	1.906.985	>100				
	2005	9–59 mos	Follow-up	2.110.341	>100				
	2008	9–59 mos	Follow-up	2.087.375	100				
Mali	2001	9 mos-14 vrs	Catch-up	4,998,491	.00				
	2004	9–59 mos	Follow-up	2,426,497	>100				
	2007	9–59 mos	Follow-up	2,562,537	>100				
Mauritania	2004	9 mos-14 vrs	Catch-up	1 167307	>100				
Maanana	2008	9_59 mos	Follow-up	464 564	98				
Mauritius	NΔ	NA	ΝΔ	-10-1,50-1 ΝΔ	NA				
Mozambique	2005	9_59 mos	Catch-up	8 222 157	97				
Mozambique	2003	9_59 mos	Eollow-up	3 3/2 280	>100				
Namibia	2000	9–59 mos	Follow-up	318 2/0	2100				
Nambia	2005	9–59 mos	Follow-up	318 905	94				
Niger	2000	9 mos_14 vrs	Catch-up	5 071 1/0	00				
Niger	2004	9 mos 14 yrs	Catch-up	222 219	× 100				
	2005	9 1105–14 yrs	Eallow up	2 042 409	>100				
Nigoria	2006	9-59 mos	Catch up	2,942,490	100				
Nigeria	2005	9 mos 14 yrs	Catch-up	20,000,974	90				
	2000	9 11105–14 yrs	Calch-up	20,000,790	03 5 100				
Dwondo	2008	9–59 mos	Follow-up	28,303,479	>100				
Rwanua	2003	6 mos-14 yrs	Catch-up	3,082,383	>100				
On a Tana & Driveira	2006	9–59 mos	Follow-up	1,380,870	>100				
Sao Tome & Principe	2007	9 mos–14 yrs	Catch-up	64,487	>100				
Senegal	2003	9 mos–14 yrs	Catch-up	4,854,077	98				
Causahallaa	2006	9–59 mos	Follow-up	1,833,931	99				
Seyechelles	INA		NA Ostak uz	NA	INA 00				
Sierra Leone	2003	9 mos–14 yrs	Catch-up	2,404,882	93				
On with Africa	2006	9–59 mos	Follow-up	751,107	100				
South Africa	2004	9–59 mos	Follow-up	3,501,447					
Quantiland	2007	9–59 mos	Follow-up	3,784,440	87				
Swaziland	2002	9–59 mos	Follow-up	127,829	81				
- .	2006	9–59 mos	Follow-up	140,143	100				
Ianzania	2001	9 mos–14 yrs	Catch-up	3,687,390	>100				
	2002	7–14 yrs	Catch-up	6,739,197	97				
	2005	9–59 mos	Follow-up	6,036,865	99				
_	2008	6 mos–10 yrs	Catch-up	10,826,519	86				
Тодо	2001	9 mos–14 yrs	Catch-up	2,393,700	99				
	2004	9–59 mos	Follow-up	887,668	100				
Uganda	2001	9 mos–14 yrs	Catch-up	614,516	>100				
	2003	6 mos–14 yrs	Catch-up	13,457,127	>100				
	2006	9–59 mos	Follow-up	5,301,424	100				

Country Zambia				Childre targete	en reached in ed age group
	Year	Target age group	Type of SIA*	No.	Administrative coverage [†] (%)
	2002	6 mos-14 yrs	Catch-up	729,469	>100
	2003	6 mos–14 yrs	Catch-up	4,955,687	>100
	2007	9–59 mos	Follow-up	2,204,553	>100
Zimbabwe	2002	9–59 mos	Follow-up	1,537,263	85
	2006	9–59 mos	Follow-up	1,407,510	95
Total				397,625,156	

TABLE 2. Measles supplementary immunization activities (SIAs), by type and country — World Health Organization (WHO) African Region, 2001–2008

* SIAs include one-time catch-up vaccination campaigns targeting a wide age range with the aim to reduce susceptibility to measles in the population and periodic follow-up SIAs targeting children born since the last SIA, thus reducing the accumulation of susceptible children in new birth cohorts. SIAs provide an initial dose of measles vaccine for children who do not access routine services and a second dose for those previously vaccinated.

[†] Administrative coverage is calculated by dividing the number of doses of vaccine administered during the SIA by the targeted number of children. The number of targeted children is usually determined by using projections of available census data.

§ Not applicable; country did not conduct any SIAs.

¹ Administrative coverage >100% usually is attributed to either an underestimation of the number of children in the targeted age group (low denominator), or vaccination of children from nontargeted geographic areas or age groups (high numerator).

** Not available.

reported an aggregated number of clinically diagnosed measles cases; however, after implementation of measles case-based surveillance, by 2005, most countries had changed to reporting laboratory-confirmed measles cases (6). Second, although the case definition for suspected measles remained the same, the change in measles reporting practices might have led to either underreporting, because of the additional resources needed to complete individual case investigations and collect blood samples, or overreporting because of overall efforts to strengthen measles surveillance.

In light of progress made toward reducing measles deaths, a more advanced goal was proposed recently for the region with several recommendations to improve vaccination coverage and surveillance performance. The AFR measles technical advisory group met in May 2008 and recommended that AFR countries aim to meet the following targets by 2012: 1) reducing estimated measles deaths by 98%, compared with 2000 estimates; 2) reducing measles incidence to < 5 cases per 1 million population per year; 3) achieving $\geq 90\%$ routine MCV1 coverage nationwide and >80% in all districts; 4) achieving >95% SIA coverage in all districts; and 5) attaining two primary measles surveillance performance indicator targets (a nonmeasles febrile rash illness rate of >2 cases per 100,000 population per year and one or more suspected measles case investigated with blood specimen in >80% of districts per year); and 6) routine reporting from all districts (10). The group also recommended that AFR countries consider introduction of MCV2 in the routine vaccination schedule if MCV1 coverage of >80% has been achieved and maintained for ≥ 3 consecutive years and at least one of the two primary measles surveillance indicator targets has been achieved and maintained for at least 2 years. For countries adopting a 2-dose routine measles vaccination schedule, continued follow-up SIAs were recommended for all new birth cohorts every 3–5 years until national MCV2 coverage of \geq 90% is sustained for at least 2 years (*10*).

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Updated Recommendation from the Advisory Committee on Immunization Practices (ACIP) for Revaccination of Persons at Prolonged Increased Risk for Meningococcal Disease

The Advisory Committee on Immunization Practices (ACIP) recommends quadrivalent meningococcal conjugate vaccine, (MCV4) (Menactra, Sanofi Pasteur, Swiftwater, Pennsylvania) for all persons aged 11-18 years and for persons aged 2-55 years at increased risk for meningococcal disease (1-3). MCV4 is licensed as a single dose. Because of the high risk for meningococcal disease among certain groups and limited data on duration of protection, at its June 2009 meeting ACIP recommended that persons previously vaccinated with either MCV4 or MPSV4 (Menomune, Sanofi Pasteur) who are at prolonged increased risk for meningococcal disease should be revaccinated with MCV4. Persons who previously were vaccinated at age ≥ 7 years and are at prolonged increased risk should be revaccinated 5 years after their previous meningococcal vaccine, and persons who previously were vaccinated at ages 2-6 years and are at prolonged increased risk should be revaccinated 3 years after their previous meningococcal vaccine. Persons at prolonged increased risk for meningococcal disease include 1) persons with increased susceptibility such as persistent complement component deficiencies (e.g., C3, properdin, Factor D, and late complement component deficiencies), 2) persons with anatomic or functional asplenia, and 3) persons who have prolonged exposure (e.g., microbiologists routinely working with Neisseria meningitidis, or travelers to or residents of countries where meningococcal disease is hyperendemic or epidemic). This report provides the rationale for the new recommendation and updates and replaces previous recommendations for revaccination with MCV4.

ACIP's Meningococcal Vaccine Work Group reviewed data on the risk for meningococcal disease, antibody titer decline, and the safety and immunogenicity of revaccination with MCV4 at 3 years and 5 years after the first dose of MCV4 or MPSV4 (2,3). Persons with prolonged increased risk for meningococcal disease have increased susceptibility to the disease or ongoing increased risk for exposure to N. meningitidis, higher levels of serum bactericidal antibody (SBA) against N. meningitidis can provide these groups increased protection against disease. SBA is a measure of the ability of sera to kill a strain of N. meningitidis in the presence of complement. In clinical trials, a baby rabbit SBA titer of 1:128 was used as a conservative correlate of protection (1). Small subsets of subjects from the MCV4 prelicensure clinical trial were revaccinated 3 years (n = 76) and 5 years (n = 134) after receiving MCV4. Of 71 persons aged 11–18 years at primary vaccination who had been vaccinated with MCV4 3 years previously, 75% and 86% had SBA titers greater than 1:128 for serogroups C and Y, respectively, before revaccination. Of 108 persons aged 2-10 years at primary vaccination who had been vaccinated with MCV4 5 years previously, 55% and 94% had SBA titers greater than 1:128 for serogroups C and Y, respectively, before revaccination. All persons revaccinated with MCV4 in these studies achieved SBA titers greater than 1:128 for serogroups C and Y. Approximately 50%-70% of persons in both the previously vaccinated (n = 210) and vaccine naive groups (n = 323) reported mild to moderate local and systemic adverse events after revaccination (or initial vaccination) with MCV4. However, no serious adverse events were reported in either group (Sanofi Pasteur, unpublished data, 2009).

On the basis of these data, expert opinion of the workgroup members, and feedback from partner organizations, the workgroup proposed that persons at prolonged increased risk for meningococcal disease be revaccinated with MCV4. ACIP approved this proposal at its June 24, 2009, meeting. Persons who previously were vaccinated at age \geq 7 years and are at prolonged increased risk should be revaccinated 5 years after their previous meningococcal vaccine. Persons who previously were vaccinated at ages 2–6 years and are at prolonged increased risk should be revaccinated 3 years after their previous meningococcal vaccine. Persons who remain in one of these increased risk groups indefinitely should continue to be revaccinated at 5-year intervals.

Although the duration of protection from MCV4 is unknown, most entering college students will have received MCV4 within the preceding 4 years. Because of the limited period of increased risk, ACIP currently does not recommend that college freshmen living in dormitories who were previously vaccinated with MCV4 be revaccinated. However, college freshmen living in dormitories who were vaccinated with MPSV4 \geq 5 years previously are recommended to be vaccinated with MCV4. Information regarding MCV4 and other recommendations for persons aged 2–55 years (2,3), including a routine recommendation for vaccination with MCV4 in persons aged 11–18 years (4), has been published previously.

References

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- CDC. Prevention and control of meningococcal disease: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 2005;54(No. RR-7).

- CDC. Recommendation from the Advisory Committee on Immunization Practices (ACIP) for use of quadrivalent meningococcal conjugate vaccine (MCV4) in children aged 2–10 years at increased risk for invasive meningococcal disease. MMWR 2007;56:1265–6.
- CDC. Revised recommendations of the Advisory Committee on Immunization Practices to vaccinate all persons aged 11–18 years with meningococcal conjugate vaccine. MMWR 2007;56:794–5.

Announcement

World Heart Day – September 27, 2009

Each year, approximately 17 million persons die from cardiovascular disease, mainly heart disease and stroke, making it the world's leading cause of death (1). Controlling certain risk factors, such as high blood pressure, high cholesterol, diabetes, obesity, tobacco use, and physical inactivity, can help prevent heart disease and stroke.

In 2000, the World Heart Federation, a nongovernmental organization based in Geneva, Switzerland, created the annual World Heart Day campaign to increase public awareness of the threat of heart disease and stroke. The theme of the 2009 World Heart Day is "Work with Heart — A Workplace That Encourages Healthy Habits Can Reduce Heart Disease and Stroke." Promoting physical activity and healthful eating and discouraging tobacco use around the workplace are simple ways to foster health in the workplace. Activities organized by members and partners of the World Heart Federation will include public talks, concerts, and sporting events. The national member organizations in the United States are the American College of Cardiology and the American Heart Association.

CDC funds heart disease and stroke prevention programs in 41 states and the District of Columbia. Additional information about these programs is available at http://www.cdc.gov/dhdsp/ state_program/index.htm. Information about World Heart Day and the World Heart Federation is available at http://www. world-heart-federation.org/what-we-do/world-heart-day.

Reference

 World Health Organization. Preventing chronic diseases: a vital investment. Geneva, Switzerland: World Health Organization; 2005. Available at http://www.who.int/chp/chronic_disease_report.

Announcement

NHANES 50th Anniversary and Conference

The 50th anniversary of the National Health and Nutrition Examination Survey (NHANES) will be celebrated on September 29, 2009, at a conference at the National Center for Health Statistics in Hyattsville, Maryland. Collaborating agencies, data users, and program and field staff members will share their perspectives on the survey. NHANES began in 1959 as the National Health Examination Survey. NHANES data come from household interviews and standardized examinations and laboratory testing of a sample of the nation's civilian, noninstitutionalized population. NHANES has expanded since the survey's inception to include a nutritional component now conducted in collaboration with the U.S. Department of Agriculture and measures of environmental exposure with the National Center for Environmental Health.

NHANES has long been a primary source of data on the nation's health. NHANES findings were used to set the goals and track the progress in reducing cholesterol levels, the prevalence of high blood pressure, and the risks of blood lead exposure in the United States. NHANES documented the rise in obesity and diabetes and produced the first populationbased estimates of human immunodeficiency virus infection and osteoporosis. NHANES data also are used for the growth charts by which pediatricians and parents check children's growth and development.

A hallmark of NHANES is its partnerships with other CDC programs, the National Institutes of Health, other U.S. Department of Health and Human Services programs, and other government agencies to collect data needed for public health policies and practice. Additional information about the NHANES 50th anniversary is available at http://www.cdc.gov/ nchs/nhanes/nhanes50th.htm.

Announcement

Epidemiology in Action: Intermediate Analytic Methods Course

CDC and Emory University's Rollins School of Public Health will cosponsor the course Epidemiology in Action: Intermediate Analytic Methods, January 11–15, 2010, at Emory University's Rollins School of Public Health. The course is designed for practicing public health professionals who have had training and experience in basic applied epidemiology and would like training in additional quantitative skills related to analysis and interpretation of epidemiologic data.

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

The prerequisite is an introductory course in epidemiology, such as Epidemiology in Action or the International Course in Applied Epidemiology. Tuition will be charged. The application deadline is December 1, 2009, or until all slots have been filled. Additional information and applications are available by mail (Emory University, Hubert Global Health Dept [Attn: Pia], 1518 Clifton Rd. NE, Rm. 746, Atlanta, GA 30322); by telephone (404-727-3485); by fax (404-727-4590); online (http://www.sph.emory.edu/epicourses); or by e-mail (pvaleri@ sph.emory.edu).

Erratum: Vol. 58, No. 34

In the QuickStats on page 955, "Percentage of Adults Aged ≥18 Years Who Engaged in Leisure-Time Strengthening Activities, by Age Group and Sex — National Health Interview Survey, United States, 2008," an error occurred. The bar for males aged ≥18 years should show the value **30.9%**.



TABLE I. Provisional cases of infrequently reported notifiable diseases (<1,000 cases reported during the preceding year) — United States, week ending September 19, 2009 (37th)*

	Current	Cum	5-year weekly		Total c for pr	ases re evious	eported years	1	States reporting cases
Disease	week	2009	average [†]	2008	2007	2006	2005	2004	during current week (No.)
Anthrax	_	_	0	_	1	1	_		
Botulism:									
foodborne	_	12	0	17	32	20	19	16	
infant	1	35	2	109	85	97	85	87	WA (1)
other (wound and unspecified)	—	17	1	19	27	48	31	30	
Brucellosis	2	70	2	80	131	121	120	114	OH (1), OR (1)
Chancroid	1	21	0	25	23	33	17	30	PA (1)
Cholera	_	4	0	5	/	9	8	6	
Cyclosporiasis ³	1	106	2	139	93	137	543	160	FL (1)
Diprimena Demostia arbayiral diagogoo [§] 1:	_	_	_	_	_	_	_	_	
California serogroup	_	22	4	62	55	67	80	112	
eastern equine	_	2	0	4	4	8	21	6	
Powassan	_	1	0	2	7	1	1	1	
St. Louis	_	7	1	13	. 9	10	13	12	
western equine	_	_	_	_	_	_	_	_	
Ehrlichiosis/Anaplasmosis [§] ,**:									
Ehrlichia chaffeensis	12	516	16	1,137	828	578	506	338	NY (4), OH (1), MO (1), VA (2), FL (1), TN (2), OK (1)
Ehrlichia ewingii	_	6	0	9	_	_	_	_	
Anaplasma phagocytophilum	6	364	17	1,026	834	646	786	537	NY (6)
undetermined	2	81	4	180	337	231	112	59	TN (2)
Haemophilus influenzae,††									
invasive disease (age <5 yrs):							_		
serotype b	_	16	0	30	22	29	9	19	
nonserotype b	2	145	2	244	199	1/5	135	135	MN (1), OK (1)
	I	1/3	2	103	100	1/9	217	105	PA (1)
Hantavirus pulmopary syndrome [§]	_	45	2	18	101	40	26	24	
Hemolytic uremic syndrome nostdiarrheal§	2	134	8	330	292	288	221	200	MI (1) TN (1)
Hepatitis C viral acute	10	1 404	15	878	845	766	652	720	PA (1) FL (3) KY (1) TN (2) OK (2) CA (1)
HIV infection, pediatric (age <13 years)§§		.,	2	_			380	436	
Influenza-associated pediatric mortality [§] , [¶]	3	118	0	90	77	43	45	_	VA (1), TX (2)
Listeriosis	14	489	22	759	808	884	896	753	PA (3), OH (4), FL (3), AR (3), CA (1)
Measles***	_	55	1	140	43	55	66	37	
Meningococcal disease, invasive ^{†††} :									
A, C, Y, and W-135		185	4	330	325	318	297	—	
serogroup B	1	98	2	188	167	193	156	—	OK (1)
other serogroup	1	20	0	38	35	32	27	_	OK(1)
unknown serogroup	4	329	9	616	550	651	765		OH(2), GA(1), CA(1)
Numps	20	289	14	454	800	6,584	314	258	NYC (18), MO (1), NC (1)
Plaquo	_	333	0	2	4	17	0	2	
Polionvelitis paralytic	_	0	0	- 3	/	17	0	- 3	
Polio virus infection nonparalytic [§]	_	_		_	_	N	N	N	
Psittacosis§	_	7	0	8	12	21	16	12	
Q fever total [§] , ^{¶¶¶} :	2	60	3	124	171	169	136	70	
acute	1	50	1	110	_		_	_	CA (1)
chronic	1	10	0	14	_	_	_	_	NY (1)
Rabies, human	_	1	0	2	1	3	2	7	
Rubella****	—	4	0	16	12	11	11	10	
Rubella, congenital syndrome	—	1	_	_	—	1	1	—	
SARS-CoV [§] , ^{††††}	—	_	—	_	_	_	_	—	
Smallpox [®]	_					105			
Streptococcal toxic-shock syndromes	1	100	1	157	132	125	129	132	OH (1)
Syphilis, congenital (age <1 yr)	_	123	8	434	430	349	329	353	
Tetalius Toxic-shock syndrome (stanbylococcal)§	1	56	1	71	20	101	27	05	
Trichinellosis	_	12	2	7 I 20	52	101	90 16	90	
Tularemia	3	53	3	123	137	95	154	134	OK (3)
Typhoid fever	7	252	13	449	434	353	324	322	NC (1), FL (1), OK (1), CA (4)
Vancomycin-intermediate Staphylococcus aureus	· 1	54	1	63	37	6	2	_	NY (1)
Vancomycin-resistant Staphylococcus aureus§	_		_		2	1	3	1	
Vibriosis (noncholera Vibrio species infections)§	23	374	10	492	549	N	N	Ν	MN (1), FL (5), WA (8), CA (9)
Yellow fever	_	_	_	_	_	_	_	_	

See Table I footnotes on next page.

TABLE I. (Continued) Provisional cases of infrequently reported notifiable diseases (<1,000 cases reported during the preceding year) — United States, week ending September 19, 2009 (37th)*

- -: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts.
- * Incidence data for reporting year 2009 is provisional, whereas data for 2004 through 2008 are finalized.
- [†] Calculated by summing the incidence counts for the current week, the 2 weeks preceding the current week, and the 2 weeks following the current week, for a total of 5 preceding years. The total sum of incident cases is then divided by 25 weeks. Additional information is available at http://www.cdc.gov/epo/dphsi/phs/files/5yearweeklyaverage.pdf.
 [§] Not reportable in all states. Data from states where the condition is not reportable are excluded from this table, except starting in 2007 for the domestic arboviral diseases and
- influenza-associated pediatric mortality, and in 2003 for SARS-CoV. Reporting exceptions are available at http://www.cdc.gov/epo/dphsi/phs/infdis.htm. Includes both neuroinvasive and nonneuroinvasive. Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Zoonotic, Vector-Borne, and Enteric Diseases (ArboNET Surveillance). Data for West Nile virus are available in Table II.
- ** The names of the reporting categories changed in 2008 as result of revisions to the case definitions. Cases reported prior to 2008 were reported in the categories: Ehrlichiosis, human monocytic (analogous to *E. chaffeensis*); Ehrlichiosis, human granulocytic (analogous to *Anaplasma phagocytophilum*), and Ehrlichiosis, unspecified, or other agent (which included cases unable to be clearly placed in other categories, as well as possible cases of *E. ewingii*).
- ⁺⁺ Data for *H. influenzae* (all ages, all serotypes) are available in Table II.
- ^{§§} Updated monthly from reports to the Division of HIV/AIDS Prevention, National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention. Implementation of HIV reporting influences the number of cases reported. Updates of pediatric HIV data have been temporarily suspended until upgrading of the national HIV/AIDS surveillance data management system is completed. Data for HIV/AIDS, when available, are displayed in Table IV, which appears quarterly.
- ¹¹ Updated weekly from reports to the Influenza Division, National Center for Immunization and Respiratory Diseases. A total of 113 influenza-associated pediatric deaths occurring during the 2008–09 influenza season have been reported. Four influenza-associated pediatric death occurring during the 2009–10 influenza season beginning September 1, 2009, has been reported.
- *** No measles cases were reported for the current week.
- ttt Data for meningococcal disease (all serogroups) are available in Table II.
- §§§ CDC discontinued reporting of individual confirmed and probable cases of novel influenza A (H1N1) viruses infections on July 24, 2009. CDC will report the total number of novel influenza A (H1N1) hospitalizations and deaths weekly on the CDC H1N1 influenza website (http://www.cdc.gov/h1n1flu).
- In 2008, Q fever acute and chronic reporting categories were recognized as a result of revisions to the Q fever case definition. Prior to that time, case counts were not differentiated with respect to acute and chronic Q fever cases.
- **** No rubella cases were reported for the current week.
- titt Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Zoonotic, Vector-Borne, and Enteric Diseases.

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals September 19, 2009, with historical data



* No measles cases were reported for the current 4-week period yielding a ratio for week 37 of zero (0).

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



	Chlamydia [†] Coccidiodomycosis Cryptosporidio						diosis								
	Previous Current 52 weeks Cum			Cum	Current	Previ 52 we	ous eeks	Cum	Cum	Curront	Prev 52 v	vious veek	Cum	Cum	
Reporting area	week	Med	Max	2009	2008	week	Med	Max	2009	2008	week	Med	Max	2009	2008
United States	12,662	22,489	25,700	794,184	835,505	260	161	472	7,972	4,575	118	123	401	4,503	5,574
New England Connecticut Maine [§] Massachusetts New Hampshire Rhode Island [§] Vermont [§]	647 224 302 93 28	766 222 48 344 39 66 22	1,655 1,306 75 945 61 244 53	28,516 8,199 1,692 13,970 1,168 2,669 818	26,173 7,487 1,797 12,578 1,470 2,028 813	N N N	0 0 0 0 0 0	1 0 0 1 0 0	1 N N 1 	1 N N 1 N	1 — — — 1	5 0 2 1 0 1	30 23 4 11 4 3 5	246 23 22 111 45 4 41	323 41 36 139 47 7 53
Mid. Atlantic New Jersey New York (Upstate) New York City Pennsylvania	2,407 	2,924 406 579 1,146 835	6,734 838 4,563 3,130 1,072	108,566 14,296 22,063 42,269 29,938	103,671 15,896 19,309 39,605 28,861		0 0 0 0	0 0 0 0	N N N	N N N N	19 12 7	13 0 4 1 7	30 2 13 8 19	525 8 157 51 309	531 33 179 83 236
E.N. Central Illinois Indiana Michigan Ohio Wisconsin	1,260 1 328 878 53 —	3,484 1,090 428 854 767 340	4,072 1,369 713 1,332 1,231 494	119,862 36,330 16,604 32,711 22,434 11,783	137,130 41,469 15,270 32,197 32,956 15,238	N N N	0 0 0 0 0	4 0 3 2 0	23 N 12 11 N	37 N 28 9 N	9 	28 2 3 5 9 8	105 11 17 13 56 40	969 99 129 182 291 268	1,469 142 132 184 450 561
W.N. Central lowa Kansas Minnesota Missouri Nebraska [§] North Dakota South Dakota	454 — 346 38 7 63	1,317 192 144 257 509 105 24 56	1,666 256 549 342 646 219 60 80	45,571 6,730 5,312 8,342 18,506 3,756 809 2,116	47,332 6,246 6,506 10,228 17,336 3,709 1,278 2,029		0 0 0 0 0 0 0	1 0 0 1 0 0 0	7 N 7 N N N	1 N 1 N N N	11 3 6 2 	18 4 1 3 2 0 2	62 13 6 33 12 7 10 10	716 158 61 200 127 71 7 92	709 221 59 152 128 83 3 63
S. Atlantic Delaware District of Columbia Florida Georgia Maryland [§] North Carolina South Carolina [§] Virginia [§] West Virginia	2,273 86 569 9 396 580 572 61	4,082 87 127 1,420 746 423 0 540 616 69	5,453 180 226 1,597 1,909 772 1,193 1,422 926 101	139,130 3,371 4,737 51,803 21,473 15,049 17,710 22,386 2,601	170,793 2,559 4,920 50,520 29,708 16,513 23,829 18,403 22,084 2,257	z z z z z z		1 0 0 1 0 0 0 0	5 1 N 4 N N N N N N N	4 1 N N 3 N N N N N	32 23 	21 0 8 6 1 0 1 1	49 1 23 23 5 16 7 6 2	746 6 293 268 30 58 34 45 10	663 10 10 305 170 28 28 38 38 55 19
E.S. Central Alabama [§] Kentucky Mississippi Tennessee [§]	663 28 635	1,738 474 253 459 573	2,207 624 458 841 809	63,612 15,693 9,218 16,941 21,760	59,743 17,902 8,387 13,928 19,526	N N N N	0 0 0 0 0	0 0 0 0 0	N N N	N N N N	4 1 2 	3 1 1 0 1	10 4 4 3 5	141 40 41 11 49	119 53 23 13 30
W.S. Central Arkansas [§] Louisiana Oklahoma Texas [§]	2,430 402 183 402 1,443	2,892 273 414 174 1,986	5,339 417 1,134 2,732 2,521	107,194 10,194 14,901 10,010 72,089	105,016 10,145 15,076 9,560 70,235	N N N	0 0 0 0	1 0 1 0 0	1 N 1 N	3 N 3 N N	16 4 2 10	11 1 2 7	271 10 6 16 258	340 36 29 87 188	1,018 48 43 77 850
Mountain Arizona Colorado Idaho [§] Montana [§] Nevada [§] New Mexico [§] Utah Wyoming [§]	777 85 356 22 175 101 38 	1,466 460 384 67 56 166 179 95 34	2,145 735 727 313 88 456 540 251 97	51,036 15,802 12,882 2,437 2,079 7,115 6,182 3,203 1,336	52,391 17,481 12,437 2,784 2,172 6,896 5,350 4,226 1,045	218 216 N N 2 —	111 109 0 0 1 0 0 0 0	369 365 0 0 4 2 2 1	6,212 6,135 N N 46 9 22 —	3,077 2,998 N N 43 24 10 2	6 5 1 	9 1 2 1 0 2 0 0	22 4 10 7 4 4 7 3 2	346 25 109 59 27 14 78 19 15	443 65 85 45 38 12 151 31 16
Pacific Alaska California Hawaii Oregon [§] Washington	1,751 1,478 273	3,627 96 2,802 120 201 414	4,685 199 3,595 247 631 571	130,697 3,181 101,994 4,160 6,683 14,679	133,256 3,350 103,699 4,049 7,048 15,110	42 N 42 N N N	41 0 41 0 0 0	172 0 172 0 0 0	1,723 N 1,723 N N N	1,452 N 1,452 N N N	20 17 3	11 0 6 0 3 1	24 1 20 1 8 6	474 5 287 1 128 53	299 3 177 2 52 65
American Samoa C.N.M.I. Guam Puerto Rico U.S. Virgin Islands		0 3 130 9	0 8 332 17	 5,076 290	73 107 5,114 486	N 	0 0 0 0	0 0 0 0	N 	N N	N N	0 0 0	0 0 0 0	N N	N N

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending September 19, 2009, and September 13, 2008 (37th)*

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum. * Incidence data for reporting year 2009 is provisional. Data for HIV/AIDS, AIDS, and TB, when available, are displayed in Table IV, which appears quarterly. † Chlamydia refers to genital infections caused by *Chlamydia trachomatis*. § Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

		is				Gonorrhe	a	Ha	Haemophilus influenzae, invasive All ages, all serotypes [†]						
		Prev	/ious				Pre	vious				Prev	ious		
Reporting area	Current week	Med	Max	Cum 2009	Cum 2008	Current week	Med	Max	Cum 2009	Cum 2008	Current week	Med	Max	Cum 2009	Cum 2008
United States	233	324	499	11,799	12,586	3,072	5,295	7,135	188,088	236,414	25	60	124	2,195	2,029
New England	6	28	55	960	1,142	126	94	301	3,473	3,687	1	3	16	142	117
Maine [§]	_	э З	14	102	239 118	/5	40	275	1,607 96	69	_	0	2	42 14	28 9
Massachusetts	- 1	11	31	429	483	42	38	112	1,416	1,555		2	5	71	57
Rhode Island§	_	1	8	35	61	7	6	19	248	240	_	0	7	3	6
Vermont [§]	5	3	15	93	125	2	1	4	32	27	_	0	1	3	8
New Jersey	53	63 7	116 17	2,205 215	2,284 368	480	590 86	1,138 122	21,783 2,991	23,193 3,818	6	12 2	25 7	444 84	373
New York (Upstate)	44	25	81	895	759	149	106	664	4,099	4,344	3	3	20	106	108
Pennsylvania	6	15	30 46	540 555	597 560	257 74	210 190	267	7,905 6,788	7,201	3	2 4	10	84 170	136
E.N. Central	31	44	80	1,564	1,889	369	1,076	1,494	37,034	49,119	_	12	28	478	331
Illinois Indiana	N	9	23 11	297 N	516 N	130	336 149	453 252	11,181 5 411	14,494	_	3	9 22	122 50	105 56
Michigan	3	12	22	425	403	212	279	493	10,452	12,031	_	Ö	3	17	17
Ohio Wisconsin	28	16 8	27 19	581 261	607 363	27	239 91	431 140	6,999 2,991	11,881 4,508	_	2 3	6 20	76 213	104 49
W.N. Central	7	25	141	1,099	1,447	95	282	393	9,782	11,959	4	3	15	118	149
lowa Kansas	3	6	14 11	221 96	226 117	_	34 35	53 83	1,137	1,093	_	0	0	13	2 17
Minnesota		0	104	250	509		44	65	1,373	2,230	3	Ŏ	10	43	46
Missouri Nebraska [§]	4	8	29 9	343 118	348 143	79 9	129 23	178 54	4,653 957	5,732	1	1	4	38 19	54 21
North Dakota	_	Ö	16	9	10		2	7	46	84	—	Ő	4	5	- 9
South Dakota		2	100	62	94	670	1 105	20	256	231		10	0		
Delaware	45	0	3	2,582	2,013	21	1,165	2,042	40,014 676	781	9	0	1	536	519
District of Columbia	41	0 36	5 59	16 1 359	51 847	218	51 418	88 486	1,870 15 101	1,809	3	0	2 10	181	5 135
Georgia	—	13	67	661	486	3	247	876	7,284	11,043	_	3	9	116	106
Maryland ^s North Carolina	N	5 0	9	170 N	189 N	106	122 0	212 470	4,053	4,376	4	1	6 17	65 61	75 57
South Carolina§	1	2	8	69	87	180	169	412	5,588	6,758	2	1	5	43	47
Virginia ³ West Virginia	3	8 1	31	257 32	271 53	143	147 10	308 23	5,072 370	7,301 546	_	1 0	6	42 25	70 18
E.S. Central	2	7	20	249	330	186	510	714	18,340	21,643	2	3	9	122	111
Alabama [§] Kentucky	1 N	3	12	120 N	191 N	21	141 84	204 135	4,432 2,689	7,072	1	0	4	28 18	17
Mississippi	Ņ	Ŏ	Ő	N	N		145	252	5,302	5,075		Ö	1	4	12
l ennessees	1	4	13	129	139	165	162	2/3	5,917	6,215	1	2	6	/2	76
Arkansas [§]	2	9 2	8	96	299 96	669 107	857	1391	31,191	36,145	3	2	22	83 13	90 11
Louisiana		3	8	96	105	51	145	420	4,796	6,521		0	1	12	8
Texas [§]	Ň	õ	0	N	N	400	554	725	19,812	22,772	_	0	1	1	7
Mountain	23	27	51	1,024	1,119	111	174	313	5,948	8,259	—	5	11	179	226
Colorado	4 7	3	26	349	94 389	34	53 56	88 152	1,801	2,451 2,548	_	1	6	63 54	88 42
Idaho [§] Montana [§]	3	3	10	125	139	_	2	13	70 51	126	_	0	1	4	12
Nevada§	5	2	10	80	82	29	30	91	1,261	1,620	_	0	2	14	14
New Mexico [§] Utab	4	1	7 15	68 161	84 234	32	24 5	52 15	802 146	972 368	_	0	3	17 23	34 30
Wyoming§	_	1	4	31	30	—	1	7	52	90	—	Ö	1	3	3
Pacific	59	51	130	1,803	2,063	357	549	765	20,523	22,391	—	2	8	93	113
California	36	2 34	57	1,209	1,369	326	466	658	542 17,292	375 18,389	_	0	3	22	38
Hawaii		07	2	10 254	34	_	11	22	434	445 856	_	0	3	22	15
Washington	11	7	74	259	264	31	46	80	1,557	2,326	_	0	2	3	2
American Samoa	—	0	0	_	—	—	0	0		3	—	0	0	_	—
Guam	_	0	0	_	_	_	1	15	_	45	_	0	0	_	_
Puerto Rico	_	2	10	63	162	—	3	24	166	208		0	1	3	1
u.s. virgin Islands	_	U	U				2	/	80	96	IN	U	0	IN	IN

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 19, 2009, and September 13, 2008 (37th)*

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts. Med: Me * Incidence data for reporting year 2009 is provisional. † Data for *H. influenzae* (age <5 yrs for serotype b, nonserotype b, and unknown serotype) are available in Table I. § Contains data reported through the National Electronic Disease Surveillance System (NEDSS). Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.

MMWR

	Hepatitis (viral, acute), by type [†]																		
			Α					В			– Legionellosis								
	Previous Previous									Prev	vious								
Reporting area	Current	52 w	Max	Cum 2009	Cum 2008	Current	52 w	Max	Cum 2009	Cum 2008	Current	52 w	Max	Cum 2009	Cum 2008				
United States	21	36	89	1,299	1,899	37	64	197	2,185	2,683	48	51	127	1,982	2,116				
New England	_	2	8	67	92	_	1	4	27	60	1	3	18	104	137				
Connecticut Maine [§]	_	0	4	17	18	_	0	3	10	23	_	1	5	42	27				
Massachusetts	_	1	3	39	47	_	0	2	6	16	_	1	6	40	58				
New Hampshire	—	0	1	5	10	—	0	2	3	5	—	0	2	8	24				
Vermont§	_	0	1	2	2	_	0	1	_	2	1	0	14	4 6	5				
Mid. Atlantic	2	5	13	175	223	5	7	17	223	318	20	15	67	785	700				
New Jersey	_	1	5 4	33 37	58 44	2	1	6 11	54 40	93 44	16	2	14 29	119 261	88 218				
New York City	_	2	6	58	75	_	1	4	42	72	_	2	20	142	98				
Pennsylvania	2	1	4	47	46	3	3	8	87	109	4	6	25	263	296				
E.N. Central	1	5 1	17 12	178 77	254 94	2	8 1	21	269 .36	366 141	8	9 1	27	353 26	468 70				
Indiana	_	Ó	3	12	14	—	1	18	46	24		1	5	25	39				
Michigan Ohio	1	1	5 4	49 31	92 29	2	2	8 13	94 69	103 84	2	2	10 17	91 206	128 203				
Wisconsin	_	Ö	3	9	25	_	Ö	4	24	14	_	Ó	3	5	28				
W.N. Central	2	2	16	89	208	—	3	16	119	58	1	2	7	66	98				
lowa Kansas	_	0	2	25 7	100 14	_	1	3	24 5	14	_	0	2	16 3	15 1				
Minnesota	_	Ő	12	14	26	_	Õ	11	20	7		Ő	3	8	9				
Missouri Nebraska§	2	0	3	22 19	25 39	_	1	5	56 13	25 5	1	1	5	29 8	54 17				
North Dakota	_	Ő	2			_	õ	1		1	_	õ	3	1					
South Dakota	_	0	1	2	4	—	0	1	1		_	0	1	1	2				
S. Atlantic Delaware	11	0	14	294	287	16 U	18 0	32	652 U	653 U	9	9	20 5	333	344 9				
District of Columbia	U	0	0	Ŭ	Ű	Ŭ	0	0	Ŭ	Ű	_	0	2	4	12				
Georgia	9	4	8	141 45	106	6	6	11	219 105	231 124	1	3	7 5	121	100				
Maryland§	—	0	4	28	33	_	1	5	47	58	—	2	10	77	99				
North Carolina South Carolina [§]	_	0	4	25 27	48 12	5	2	19 4	135 35	51 52	_	0	6 1	39	23				
Virginia [§]	1	Ö	3	24	37	2	1	10	62	78	1	1	5	35	39				
West Virginia		0	1	1	5	1	0	19	49	59		0	2	6	24				
Alabama [§]	_	0	2	30	63 9	5	2	7	220 65	280	3	2	2	87	13				
Kentucky	—	0	1	7	23	—	2	7	58	67	2	1	3	39	43				
Tennessee§	_	0	2	8 8	4 27	3	2	6	18 79	33 98	1	1	8	37	34				
W.S. Central	_	3	43	103	180	4	10	99	338	528	1	1	21	45	59				
Arkansas§	_	0	1	4	6 10	_	1	5	37	42	1	0	2	4	10				
Oklahoma	_	0	6	3	7	4	2	17	75	78	_	0	6	3	3				
Texas§	_	3	37	93	157	—	6	76	193	341	—	1	19	34	38				
Arizona	2	3	7	116 56	170	2	3	7 4	96 36	146 56	1	2	8	77 35	61 14				
Colorado		Ō	5	34	31	—	ò	2	16	25	1	ò	2	8	7				
Idaho ^s Montana [§]	_	0	1	35	16 1	_	0	2	7	7	_	0	1	1	3				
Nevada§	1	Ő	3	8	.7	2	Ő	3	24	31	—	Ő	2	10	9				
New Mexico ^s	_	0	1	6 4	15 11	_	0	2	5	8 12	_	0	2	2 16	6 18				
Wyoming§	_	õ	Ó		3	—	ŏ	2	3	5	—	õ	1	1					
Pacific	3	7	17	247	422	3	6	36	241	274	4	3	12	132	158				
Alaska California	3	0 5	1 17	3 196	3 342	3	0 5	1 28	2 178	9 190	3	0	1	1 105	1 122				
Hawaii	_	0	1	5	16	_	0	1	4	6	_	0	1	1	6				
Uregon ^s Washington	_	0 1	2 4	12 31	23 38	_	0 1	4 8	26 31	33 36	1	0	2 4	10 15	14 15				
American Samoa	_	0	0	_	_	_	0	0	_	_	Ν	0	0	N	N				
C.N.M.I.	—			—	—	—			—	_	_			_	_				
Guam Puerto Rico	_	0	0	17	20	_	0	0 3	12	44	_	0	0	_	_				
U.S. Virgin Islands	_	0	0	_	_	_	0	0	_	_	_	0	0	_	_				

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 19, 2009, and September 13, 2008 (37th)*

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

U: Unavailable. —: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum. * Incidence data for reporting year 2009 is provisional. * Data for acute hepatitis C, viral are available in Table I. * Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

		L	yme disea	ise				Malaria		Meningococcal disease, invasive [†] All groups						
		Pre	vious				Prev	vious				Prev	ious			
Reporting area	Current week	Med	Max	Cum 2009	Cum 2008	Current week	Med	Max	Cum 2009	Cum 2008	Current week	5∠ w Med	Max	Cum 2009	Cum 2008	
United States	247	480	1,637	19,847	24,672	11	23	46	804	852	6	17	48	632	884	
New England	1	90	327	3,378	9,161	_	1	5	30	43	_	0	4	21	24	
Maine§	_	8	73	467	3,163	_	0	4	5 1	10	_	0	1	2	4	
Massachusetts	_	28	213	1,881	3,895	_	0	3	19	23	—	0	3	12	16	
Rhode Island [§]	_	0	72	765 54	1,325	_	0	1	1	2	_	0	1	2	1	
Vermont§	1	4	36	211	298	_	0	1	2	4	—	0	1	1	_	
Mid. Atlantic New Jersev	205	240 35	1,401 264	11,970 2.629	9,936 2.893	1	5 0	17 3	187	234 56	_	2 0	5 2	71 8	96 13	
New York (Upstate)	105	86	1,368	3,069	3,355	1	1	10	37	25	—	0	2	18	25	
Pennsylvania	100	4 53	618	6,124	3,071	_	3	4	39	30	_	1	2	33	39	
E.N. Central	2	19	179	1,555	1,941	_	3	8	111	117	2	3	8	103	153	
Illinois Indiana	_	1	11 4	83 33	98 33	_	1	4	46 12	62 5	_	1	6 3	27 24	55 22	
Michigan	_	1	11	76	64	—	Õ	3	18	13	_	Ö	5	18	26	
Ohio Wisconsin	2	1 14	3 165	36 1.327	34 1.712	_	1 0	6 1	31 4	22 15	2	0	3 1	28 6	32 18	
W.N. Central	_	5	336	172	532	_	1	7	41	51	_	1	9	50	77	
lowa	—	1	12	72	92	_	0	2	9	8	—	0	1	6	16	
Minnesota	_	0	326	67	418	_	0	7	13	20	_	0	4	10	21	
Missouri Nebraska§	_	0	2	4 13	4	_	0	2	9	10 8	_	0	3 1	18 5	23 10	
North Dakota	—	Ő	10	-		—	Ő	ò		_	—	Ő	3	1	1	
South Dakota		0	1	1	3	_	0	1	1		-	0	1	2	2	
Delaware	30	63 12	63	2,523 746	2,857	<u> </u>	6 0	1	247	210	_	2	9 1	2	126	
District of Columbia		0	5	18	54 50	6	0	2	5	2	—	0	0	<u> </u>		
Georgia	_	0	6	39	31		1	5	54	46	1	0	2	22	14	
Maryland [§] North Carolina	_	27 1	130 14	1,140 56	1,416 16	_	1	8	52 21	55 22	_	0	1 5	7 18	13 11	
South Carolina§		Ó	3	19	_18	—	Õ	1	2	8	—	Ö	1	10	20	
Virginia® West Virginia	19	11 0	61 27	342 100	544 108	_	1 0	4	32	36	_	0	2	9 5	17 5	
E.S. Central	1	0	2	20	39	_	1	3	24	13	_	0	3	21	40	
Alabama [§]	_	0	1	2	9	_	0	3	7	3	_	0	1	5	5	
Mississippi	_	Ö	Ó	-	1	_	Ö	1	1	1	_	Ö	1	2	9	
Tennessee	1	0	2	17	25	_	0	3	8	5	_	0	1	10	19	
W.S. Central Arkansas§	_	1 0	21 0	37	/8	_	1 0	8 1	34 3	57		1 0	12 2	60 5	95 13	
Louisiana	_	0	0	_	3	_	0	1	3	3		0	3	11	19	
Texas [§]	_	1	21	37	75	_	1	7	26	52		1	9	36	51	
Mountain	—	1	13	37	45	—	0	5	24	22	—	1	4	50	47	
Arizona Colorado	_	0	2	4 4	8	_	0	2	8	10	_	0	2	13 16	6 9	
Idaho [§]	—	0	2	9	7	—	0	1	1	1	—	0	1	5	4	
Nevada§	_	0	2	12	4 11	_	0	1	4	4	_	0	2	4	47	
New Mexico [§]	_	0	1	1	8	_	0	1		2	_	0	1	3	8	
Wyoming§	_	0	1	1	2	_	0	0	_		_	0	2	4	2	
Pacific	8	4	13	155	83	4	3	10	106	105	1	3	14	142	226	
California	7	3	11	133	5 44	3	2	8	2 80	4 75	1	2	2	5 95	167	
Hawaii Oragan [§]	Ν	0	0	N	N	_	0	1	1	2	—	0	1	3	4	
Washington	1	0	12	12	27 7	1	0	∠ 3	9 14	4 20	_	0	6	20 13	20 23	
American Samoa	Ν	0	0	Ν	Ν	_	0	0	_	_	_	0	0	_	_	
G.N.M.I. Guam	_	0	0	_	_	_	0	2	_	1	_	0	0	_	_	
Puerto Rico	N	0	0	N	N	—	0	1	2	2	—	0	1	—	2	
U.S. Virgin Islands	N	0	0	N	N	_	0	0	_	_	_	0	0	_		

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 19, 2009, and September 13, 2008 (37th)*

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum. * Incidence data for reporting year 2009 is provisional. † Data for meningococcal disease, invasive caused by serogroups A, C, Y, and W-135; serogroup B; other serogroup; and unknown serogroup are available in Table I. § Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

(01)	Pertussis						Ra	bies, anin	nal	Rocky Mountain spotted fever						
		Pre 52 v	vious				Previous 52 weeks					Prev 52 w	ious			
Reporting area	Current week	Med	Max	Cum 2009	Cum 2008	Current week	Med	Мах	Cum 2009	Cum 2008	Current week	Med	Max	Cum 2009	Cum 2008	
United States	114	279	1,697	9,309	6,523	111	68	138	2,658	3,117	8	29	179	1,061	1,707	
New England	—	14	27	437	735	1	7	14	226	293	—	0	2	9	4	
Maine [†]	_	1	10	64	42 26	_	1	5	36	36	_	0	2	4	1	
Massachusetts	—	8	21	266	570	—	0	0			—	0	1	4	1	
Rhode Island [†]	_	0	5	57 11	23 63	_	0	3	24 27	26	_	0	2	_	1	
Vermont [†]		0	1	8	11	1	1	4	38	54	—	0	1	1	_	
Mid. Atlantic New Jersev	13	22 3	64 12	794 128	774 162	8	14 0	27 0	447	676	_	1 0	29 2	54	107 73	
New York (Upstate)	10	5	41	155	293	8	8	20	328	366	—	0	29	10	12	
New York City Pennsylvania	3	0 12	21 33	53 458	50 269	_	0 4	2 17	1 118	14 296	_	0	4 2	24 20	11 11	
E.N. Central	52	54	238	1,922	1,081	9	2	19	197	205	_	1	6	62	127	
Illinois Indiana	_	11 4	45 158	284 181	229 42	3	1	9	80 17	85 7	_	1	6	39 4	94 6	
Michigan	15	11	30	522	175	4	1	6	57	66	_	õ	2	5	3	
Ohio Wisconsin	37	20 3	57 12	829 106	526 109	2 N	0	7	43 N	47 N	_	0	4	14	24	
W.N. Central	1	35	872	1,304	538	4	5	17	214	230	3	4	26	237	367	
lowa Kansas	_	6 4	21 12	139 143	84 42	_	0	5	24 60	17 52	_	0	2	4	7	
Minnesota		Ö	808	165	156	1	Ó	11	45	44	_	Ő	1	2		
Nissouri Nebraska†	1	20 4	51 32	706	171 62	3	1	5	54	51 31	3	4 0	25	218	341 16	
North Dakota	—	0	24	17	1	—	0	9	4	17		0	1	—		
S Atlantic	27	28	71	24 1 174	643	84	25	4	1 215	1 279	2	13	42		589	
Delaware	_	0	2	10	11	_	0	0			_	0	3	16	26	
Florida	16	9	2 32	2 426	4 194	_	0	0 95	131	138	_	0	0	5	6 9	
Georgia	1	3	11	115	63	72	0	71	334	290	—	0	6	37	68	
North Carolina	9	20	9 65	213	93 79	N	2	4	264 N	328 N	2	6	36	227	263	
South Carolina [†]	1	4	17	175	86	10	0	0	200	456	_	0	9	16	31	
West Virginia	_	0	5	24	8	2	2	6	87	67	_	0	1	47	8	
E.S. Central	3	15	33	573	228	_	2	7	71	140	3	4	19	193	249	
Kentucky	_	4	19	178	30 59	_	1	0 4	37	35	3	0	6	46	67	
Mississippi		1	4	41	78	—	0	2		102	_	0	1	120	10	
W S Central		56	389	1 872	1 031	_	0	13	45	75	_	1	161	106	225	
Arkansas [†]	_	4	38	176	68	—	ŏ	5	23	41	_	Ö	61	47	44	
Oklahoma	_	2	8 45	90 37	64 32	_	0	0 13	21	32	_	0	1 98	2 44	5 142	
Texas [†]	—	46	304	1,569	867	—	0	1	1	2	_	0	6	13	34	
Mountain Arizona	6	17 3	31 10	638 152	623 172	N	1	9 0	57 N	73 N	_	0	3	19 4	36 10	
Colorado	5	5	12	205	116	_	0	0	_		_	Ö	0		1	
Idano ¹ Montana [†]	1	1	5 4	60 12	24 76	_	0	2 4	16	9	_	0	1	1	1	
Nevada [†]	—	0	3	10	26	—	0	1	4	10	—	0	1	1	2	
Utah	_	4	19	152	163	_	0	6	4	24 7	_	0	1	1	4 5	
Wyoming [†]		0	5	8	13	_	0	4	17	15	—	0	1	3	10	
Alaska	12	18 1	98 21	595 33	870 118	5	5 0	12 2	186 11	146 12	N	0	1 0	2 N	3 N	
California	_	5	19	143	394	5	4	12	160	127		0	1	2		
Oregon [†]	1	3	16	186	130	_	0	3	15	7		0	0		3	
Washington	10	6	76	210	218		0	0				0	0			
American Samoa C.N.M.I.	_	0	0	_	_	N	0	0	N	N	N	0	0	N	N	
Guam	_	0	Q	_	_	_	0	0			N	0	0	N	N	
U.S. Virgin Islands	_	0	0	í —	_	N	0	0	∠8 N	47 N	N	0	0	N	N	

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending September 19, 2009, and September 13, 2008 (37th)*

C.N.M.I.: Commonwealth of Northern Mariana Islands. U: Unavailable. —: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum. * Incidence data for reporting year 2009 is provisional. † Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

		s	almonello	sis		Shig	ga toxin-p	roducing	E. coli (ST	Shigellosis						
	Current	Pre 52 v	vious weeks	Cum	Cum	Current	Prev 52 w	/ious /eeks	Cum	Cum	Current	Pre 52 v	vious weeks	Cum	Cum	
Reporting area	week	Med	Max	2009	2008	week	Med	Max	2009	2008	week	Med	Max	2009	2008	
United States	762	907	2,323	29,893	32,838	71	86	255	2,804	3,470	165	313	1,268	10,584	13,838	
New England Connecticut Maine [§] Massachusetts New Hampshire Rhode Island [§]	1 	32 0 22 3 2	324 298 7 38 42 11	1,522 298 83 805 206 87	1,739 491 107 887 111 73	 	3 0 1 1 0	50 50 3 6 3 1	159 50 14 58 24 —	184 47 15 86 15 7	 	3 0 3 0 0	33 28 1 27 4 1	239 28 2 183 13 8	185 40 18 110 4 10	
Vermont§	1	1	5	43	70	—	0	6	13	14	_	0	2	5	3	
Mid. Atlantic New Jersey New York (Upstate) New York City Pennsylvania	50 — 40 3 7	87 9 24 19 29	182 32 66 49 66	3,203 237 937 813 1,216	4,129 982 949 928 1,270	5 4 1	7 1 3 1 1	19 5 9 5 6	247 31 100 39 77	352 104 118 39 91	24 6 18	56 13 5 9 24	79 35 23 23 61	2,030 416 166 308 1,140	1,752 604 457 552 139	
E.N. Central Illinois Indiana Michigan Ohio Wisconsin	32 — 3 29 —	91 26 7 18 28 11	142 50 50 29 52 29	3,340 892 245 688 1,085 430	3,711 1,087 426 697 929 572	3 3	12 1 3 3 3	74 10 6 43 15 10	437 66 39 106 104 122	562 96 71 98 132 165	8 1 	62 12 1 5 35 10	132 25 21 24 80 42	1,885 384 38 167 940 356	2,691 749 504 92 1,049 297	
W.N. Central lowa Kansas Minnesota Missouri Nebraska [§] North Dakota South Dakota	38 8 6 24 	51 7 13 12 5 0 3	109 15 18 51 29 41 30 22	1,969 313 270 458 460 272 40 156	2,099 324 346 530 576 178 31 114	4 2 1 1 	12 3 1 2 2 1 0 0	39 14 7 18 10 6 28 12	531 131 33 155 86 66 3 57	606 158 35 124 124 126 1 38	22 — 1 21 — —	16 2 3 2 4 0 0 0	49 12 11 14 40 3 9 1	662 49 159 64 364 19 3 4	672 117 32 234 176 5 33 75	
S. Atlantic Delaware District of Columbia Florida Georgia Maryland [§] North Carolina South Carolina [§] Virginia [§] West Virrinia	336 1 229 59 10 14 23	262 2 0 115 39 15 22 15 20 4	440 8 5 197 96 26 104 54 88 23	8,188 80 20 3,912 1,542 502 788 530 655 159	8,026 115 49 3,258 1,576 615 795 757 707 154	8 6 2	12 0 3 1 2 0 3 0	30 2 1 7 4 6 21 3 16 3	446 11 120 52 60 74 21 88 19	614 10 69 103 71 32 188 32	24 — 13 6 2 1 2	46 1 9 13 6 5 3 5 0	85 8 24 30 14 27 14 59 3	1,626 76 6 326 469 257 253 90 143 6	2,289 7 16 630 833 73 139 433 130 28	
E.S. Central Alabama [§] Kentucky Mississippi Tennessee [§]	24 5 9 1 9	56 15 10 14 14	124 38 18 47 62	1,924 467 351 578 528	2,382 679 320 806 577	9 2 7	4 1 1 0 2	12 4 7 1 5	160 36 55 6 63	201 49 66 4 82	5 1 2 	18 3 2 1 11	58 11 25 4 48	586 99 145 32 310	1,392 330 210 277 575	
W.S. Central Arkansas [§] Louisiana Oklahoma Texas [§]	89 24 20 45	110 12 14 14 56	1,333 34 43 102 1,204	3,185 435 599 457 1,694	4,558 541 795 542 2,680	4 4	4 0 1 2	139 4 1 82 55	125 26 — 21 78	253 42 7 22 182	33 7 11 15	55 8 4 5 41	967 20 17 61 889	1,850 243 108 208 1,291	3,035 402 508 103 2,022	
Mountain Arizona Colorado Idaho [§] Montana [§] Nevada [§] New Mexico [§] Utah Wvoming [§]	42 15 17 2 	57 20 13 2 4 5 6 1	121 47 34 10 7 13 26 15 6	2,122 743 488 135 73 185 228 227 43	2,423 767 532 128 87 171 430 252 56	15 1 2 2	10 1 2 0 0 1 2 0	40 4 18 15 3 4 2 7 2	364 55 114 60 15 22 23 70 5	407 51 118 82 29 14 42 61 10	27 16 6 1 	24 17 2 0 1 2 0 0	54 42 11 2 5 11 12 3 1	856 636 73 8 13 52 59 15 	691 335 76 10 6 162 73 26 3	
Pacific Alaska California Hawaii Oregon [§]	150 99 6 	126 1 95 5 8	537 6 516 13 15	4,440 56 3,355 184 296 540	3,771 42 2,735 197 327 470	23 4 10	10 0 5 0 1	31 15 1 6	335 169 3 47	291 5 136 11 51	22 — 19 —	27 0 20 0 1	75 1 65 4 10	850 2 686 27 29	1,131 1 981 35 55	
American Samoa C.N.M.I. Guam Puerto Rico U.S. Virgin Islands	43 — — 2 —	0	00 1 2 40 0	549 — 251 	470 2 11 521		0 0 0 0 0	0 0 0 0		00 — — — —	3 — — — —	3 1 0 0 0	1 2 1 2 0	3 — 7	59 1 14 24	

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 19, 2009, and September 13, 2008 (37th)*

C.N.M.I.: Commonwealth of Northern Mariana Islands.
 U: Unavailable. —: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.
 * Incidence data for reporting year 2009 is provisional.
 † Includes *E. coli* O157:H7; Shiga toxin-positive, serogroup non-O157; and Shiga toxin-positive, not serogrouped.
 § Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

		Streptococcal	diseases, inv	asive, group A		Streptococcus pneumoniae, invasive disease, nondrug resistant [†] Age <5 years							
		Prev	vious				Previ	ious					
Reporting area	Current week	Med	Max	Cum 2009	Cum 2008	Current week	Med	Max	Cum 2009	Cum 2008			
United States	35	101	239	3,928	4,211	11	36	122	1,205	1,260			
New England	_	5	28	229	301	_	1	12	43	61			
Connecticut	_	0	21	63	86	_	0	11					
Maines	_	3	10	97	20 140	_	1	4	30	45			
New Hampshire	—	1	4	34	20	—	0	2	8	8			
Rhode Island [§]	_	0	2	9 13	22	_	0	2	2	7			
Mid Atlantic	1	19	43	788	861	1	5	33	182	158			
New Jersey	_	3	6	104	155	_	1	4	31	47			
New York (Upstate)	_	7	25	262	270	_	2	17	85	69			
Pennsylvania	1	6	12	272	278	N	0	2	N	42 N			
E.N. Central	1	17	42	742	798	2	6	18	182	228			
Illinois	—	5	12	207	212	_	1	5	23	64			
Indiana Michigan	_	3	23	119 121	106	1	0	13	26 49	26 58			
Ohio	1	4	13	186	219	1	1	6	54	42			
Wisconsin	—	2	11	109	122	—	1	4	30	38			
W.N. Central	1	6	37	321	314	1	2	11	108	68			
Kansas	_	1	5	37	32	N	0	1	N	N			
Minnesota	—	0	34	146	150	1	0	10	61	19			
Missouri Nebraska§	_	2	8	71	74	_	0	4	29	30			
North Dakota	_	0	4	11	8	_	0	3	4	6			
South Dakota	1	0	3	21	19	—	0	2	6	6			
S. Atlantic	10	22	48	897	862	4	6	16	226	246			
Delaware District of Columbia	_	0	3	10	6 12	N	0	0	N	N			
Florida	4	6	12	221	196	1	1	6	53	46			
Georgia Maryland [§]	3	5	13	213	190 148	2	2	6	58 51	67 46			
North Carolina	2	2	12	83	110	N	Ó	0	N	Ň			
South Carolina§	_	1	5	57	55	1	1	6	34	43			
Virginia ^s West Virginia		3	9	34	33	_	0	4	18	38			
E.S. Central	2	3	10	151	149	_	2	7	66	65			
Alabama§	Ň	Õ	Ő	N	Ň	N	ō	Ó	Ň	Ň			
Kentucky Mississioni	1 N	1	5	29 N	32 N	N	0	0	N 14	N 8			
Tennessee§	1	3	9	122	117	_	1	6	52	57			
W.S. Central	19	9	79	343	380	1	6	46	204	198			
Arkansas§	_	0	2	14	8	1	0	4	22	11			
Oklahoma	3	3	20	111	88	_	1	7	43	49			
Texas [§]	16	5	59	207	269	—	3	34	126	127			
Mountain	1	10	22	341	435	2	4	16	171	198			
Colorado	_	3	9	110	152		2	10	90 32	45			
Idaho§	1	Õ	2	8	12	—	Ó	2	7	3			
Montana [§]	N	0	0	N	N	N	0	0	N	N			
New Mexico§	_	2	7	59	103	_	0	4	15	27			
Utah	—	1	6	41	43	—	0	5	27	28			
vvyoming ^s	_	0	1	1	6	_	0	1	_	1			
Alaska	_	3	9	116 22	111 28	_	0	4	23 17	38 24			
California	Ν	0	õ	N	Ň	Ν	õ	õ	N	N			
Hawaii Orogon [§]		3	8	94	83		0	2	6	14			
Washington	N	0	0	N	N	N	0	0	N	N			
American Samoa	_	0	0	_	30	Ν	0	0	Ν	Ν			
C.N.M.I.	—			—	_	_			—	—			
Guam Puerto Rico	N	0	0	N	N	N	0	0	N	N			
U.S. Virgin Islands		õ	Ő		_	N	0	Ő	N	N			

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 19, 2009, and September 13, 2008 (37th)*

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

 U: Unavailable. —: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.
 * Incidence data for reporting year 2009 is provisional.
 * Includes cases of invasive pneumococcal disease, in children aged <5 years, caused by *S. pneumoniae*, which is susceptible or for which susceptibility testing is not available. (NNDSS event code 11717). § Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

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<u>·</u>		S	treptococ	cus pneur	<i>noniae</i> , ir	vasive dise	ease, dru	g resistan	t†								
			All ages				Aç	ged <5 yea	ars		Syphilis, primary and secondary						
	Current	Prev 52 w	/ious /eeks	C	C 111	Current	Prev 52 w	/ious /eeks	C	C	Current	Prev 52 w	vious veeks	C	C		
Reporting area	week	Med	Max	2009	2008	week	Med	Max	2009	2008	week	Med	Max	2009	2008		
United States	16	60	276	2,031	2,264	_	9	21	313	348	135	264	452	9,274	9,035		
New England	_	1	48	35	53	_	0	5	3	7	4	5	15	233	222		
Connecticut Maine [§]	_	0	48 2	9	/ 15	_	0	5	1	1	1	1	5 1	43	23 9		
Massachusetts	—	Ö	1	3		_	Ö	1	2	_	3	4	11	164	156		
New Hampshire	_	0	3	5	19	_	0	0	_		_	0	2	13	13		
Vermont§	_	0	2	11	13	_	0	0	_	2	_	0	2		7		
Mid. Atlantic	3	3	14	121	231	_	0	3	20	21	31	35	51	1,315	1,183		
New Jersey	2	0	0	54		_	0	0	10		2	4	13	157	155		
New York City		ò	4	3	93	_	0	2		1	23	23	40	825	740		
Pennsylvania	1	1	8	64	90	_	0	2	10	14	6	6	12	245	192		
E.N. Central	3	11	41	456 N	479 N		1	7	64	64	7	23	44	769	838		
Indiana		3	32	162	164		0	6	22	20	1	2	10	120	102		
Michigan	_	0	2	19	17	_	0	1	2	2	5	4	18	180	130		
Ohio Wisconsin	3	0	18	275	298	_	1	4	40	42	1	6 1	1/	215	225 41		
W.N. Central	_	2	161	95	159	_	0	3	20	32	_	6	11	218	304		
lowa	—	ō	0	_		—	ŏ	Ő		_	_	ŏ	2	17	14		
Kansas Minnesota	_	1	5 156	39	59	_	0	2	13	23	_	0	3	22 40	24		
Missouri	_	1	5	44	69	_	0	1	5	20	_	3	7	121	179		
Nebraska§	—	0	0		_	_	0	0	_	_	—	0	3	14	10		
South Dakota	_	0	3	2	2	_	0	2	2	3	_	0	1	3	_		
S. Atlantic	6	26	53	965	929	_	4	14	144	153	24	64	262	2,298	1,975		
Delaware		0	2	15	3		0	0			—	0	3	23	10		
Florida	N 4	15	36	563	527	IN	2	13	N 89	IN 98	1	20	32	697	95 735		
Georgia	1	8	25	295	313	_	1	5	48	47		14	227	541	444		
Maryland ⁹		0	1	4 N	4 N	N	0	0	N	1 N	3 17	6	16 21	221	244		
South Carolina [§]		ŏ	Ö				ŏ	Ő			3	2	6	86	62		
Virginia [§] West Virginia	N	0	0	N	N	Ν	0	0	N	N	_	7	15	224	183		
FS Control	2	2	13	109	240		1	3	20	15		22	2	790	0 771		
Alabama§	Ň	0	23	N	240 N	N	Ó	0	29 N	43 N	-	8	17	288	319		
Kentucky	1	1	5	56	59	_	0	2	7	10	—	1	10	47	61		
Tennessee§	1	3	23	139	152	_	0	3	20	8 27	4	4	18	296	282		
W.S. Central	2	1	6	74	76	_	0	3	14	12	41	48	80	1.751	1.542		
Arkansas§	2	1	5	42	13	—	0	3	9	3	14	4	35	167	113		
Louisiana Oklahoma	N	1	5	32 N	63 N	N	0	1	5 N	9 N	2	11	40 7	303	425		
Texas§	_	Õ	ŏ	_	_	_	Õ	Ő	_	_	25	32	50	1,236	948		
Mountain	_	2	7	84	95	_	0	3	17	12	13	9	18	314	454		
Arizona Colorado	_	0	0	_	_	_	0	0	_	_	_	4	9 4	132 64	232 109		
Idaho§	Ν	Õ	1	Ν	N	Ν	Õ	1	Ν	Ν	—	Ó	2	3	3		
Montana [§]	_	0	1	33		_	0	0	7		10	0	7	76	60		
New Mexico§	_	ò	4 0			_	0	0			3	1	5	37	31		
Utah	_	1	6	42	50	_	0	3	9	7	—	0	2		16		
wyoming ^s	_	0	2	9	1	_	0	1	1			0	1	1 5 0 7	1 740		
Alaska	_	0	0			_	0	0				44	0	1,587	1,740		
California	N	0	0	N	N	N	0	0	N	N	8	40	60	1,447	1,579		
Hawaii Oregon§	N	0	1	3 N	2 N	N	0	1	2 N	2 N	_	0	3	21 32	16 13		
Washington	N	ŏ	Ő	N	N	N	ŏ	õ	N	N	3	2	7	87	137		
American Samoa	Ν	0	0	Ν	Ν	Ν	0	0	Ν	Ν	_	0	0	_	_		
C.N.M.I. Guam	_				_				_	_	_			_	_		
Puerto Rico	_	0	0	_	_	_	0	0	_	_	_	3	16	142	108		
U.S. Virgin Islands	_	0	0	_	_	_	0	0	_	_	_	0	0	_	_		

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 19, 2009, and September 13, 2008 (37th)*

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

Christian Commonwealth of Normer Mariana Islands.
 U: Unavailable. —: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum.
 * Incidence data for reporting year 2009 is provisional.
 † Includes cases of invasive pneumococcal disease caused by drug-resistant *S. pneumoniae* (DRSP) (NNDSS event code 11720).
 § Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

<u> </u>						West Nile virus disease [†]										
			Ne	euroinvasi	ve		Nonneuroinvasive [§]									
	Previous						Prev	ious				Prev	ious			
Reporting area	Current	52 v	Max	Cum	Cum	Current	52 w	Max	Cum	Cum	Current	52 w	Max	Cum	Cum	
United States	101	455	1 035	12 879	21 268		1	43	178	550		0	34	166	575	
New England		-00	46	199	1.193	_	0	1		5	_	0	0		3	
Connecticut	_	Ő	21		617	_	ŏ	Ó	_	5	_	Õ	Ő	_	3	
Maine ¹ Massachusetts	_	0	11	5	178	_	0	0	_	_	_	0	0	_	_	
New Hampshire	_	4	11	145	191	_	ŏ	Ó	_	_	_	ŏ	Ő	_	_	
Rhode Island [¶]	—	0	1	4		—	0	1	—	—	_	0	0		—	
Vermont [®]	10	2	17	43	1 694	_	0	0			_	0	0	-	17	
New Jersey	N	0	0	1,090 N	1,004 N	_	Ő	2		2	_	Ő	0	_	4	
New York (Upstate)	N	0	0	Ν	Ν	—	0	3	1	17	—	0	1	—	6	
New York City Pennsylvania	19	39	0 58	1 090	1 684	_	0	0	1	8	_	0	1	1	5	
E.N. Central	34	161	254	4,583	5,143	_	0	6	3	30	_	0	3	3	17	
Illinois	5	38	73	1,126	783	—	0	4	1	6	—	0	0	_	8	
Indiana Michigan	9	2 48	24 90	250 1.342	2 127	_	0	1	2	2	_	0	1	1	1	
Ohio	20	42	91	1,475	1,643	_	ŏ	2	_	11	_	ŏ	2	2	_	
Wisconsin		13	55	390	590	_	0	2		2	_	0	0		4	
W.N. Central	13 N	21	114	705 N	882 N	_	0	3	12	41	_	0	5	32	118	
Kansas		5	22	183	330	_	ŏ	2	_	10	_	ŏ	2	4	13	
Minnesota		0	0			—	0	0	_	2	—	0	1	1	8	
Missouri Nebraska¶	13 N	10	51	465 N	516 N	_	0	2	1	8	_	0	0	15	31	
North Dakota	_	õ	108	57	_	_	ŏ	ō	_	2	_	ŏ	1	1	35	
South Dakota	_	0	4		36	—	0	3	5	11	_	0	2	9	26	
S. Atlantic Delaware	20	56 0	146 4	1,472	3,495 32	_	0	2	5	17	_	0	3	_	16 1	
District of Columbia	_	Õ	3	8	18	_	Õ	Õ	_	4	_	Õ	1	_	2	
Florida	9 N	28	67	947 N	1,217	—	0	0		3	—	0	0	—		
Maryland	N	Ő	0	N	N	_	0	2		4	_	Ő	2	_	6	
North Carolina	N	0	0	N	N	—	0	0	_	2	—	0	0	—	1	
South Carolina [®]	_	2	54 119	154 28	1 060	_	0	2	3	_	_	0	0	_	1	
West Virginia	11	9	32	327	531	_	ŏ	õ	_	1	_	ŏ	Ő	_		
E.S. Central	_	11	28	358	899	_	0	5	25	42	_	0	5	15	50	
Alabama ¹ Kentucky	N	11	28	356 N	888 N	_	0	0	2	11	_	0	2	_	5	
Mississippi	_	õ	1	2	11	_	ŏ	5	22	19	_	ŏ	4	14	38	
Tennessee ¹	N	0	0	N	N	_	0	1	1	11	—	0	1	1	7	
W.S. Central Arkansas ¹	_	97	747 47	3,421 96	6,321 528	_	0	12	56 1	58	_	0	5	17	48	
Louisiana	_	1	7	76	58	_	õ	3	7	13	_	õ	5	6	19	
Oklahoma	N	0	0	N 2 240	N	—	0	1	4	2	_	0	0		5	
Mountain	15	00 32	83	3,249 971	5,735 1,557	_	0	8	44 41	57 74	_	0	12	61	163	
Arizona		0	0			_	ŏ	5	11	39	_	ŏ	7	4	35	
Colorado	15	12	44	403 N	637 N	_	0	4	13	15	—	0	11	38	53	
Montana [¶]		2	20	105	233	_	0	1	2		_	0	1	1	5	
Nevada	N	0	0	N	N	—	0	2	7	8	—	0	1	5	7	
New Mexico ¹	_	12	20	134	171 506	_	0	2	4	4	_	0	1	2	10	
Wyoming [¶]	_	0	1	525	10	_	ŏ	1	2		_	Ő	2	5	8	
Pacific	—	2	7	80	94	—	0	19	34	247	—	0	10	37	143	
Alaska California	_	1	6	50	46	_	0	0 19	25	242	_	0	0	22	129	
Hawaii	_	1	4	30	48	_	ŏ	0			_	ŏ	0			
Oregon [¶]	N	0	0	N	N	—	0	1	1	3	—	0	3	6	13	
American Samoa	N N	0	0	IN N	IN N	_	0	3	8	2	_	0	4	9	-	
C.N.M.I.						_			_	_	_			_	_	
Guam	10	2	3		55	—	0	0	_	—	—	0	0	—	—	
US Virgin Islands	13	0	23	332	450	_	0	0	_	_	_	0	0	_	_	
			0				~	0					<u> </u>			

TABLE II. (Continued) Provisional cases of selected notifiable diseases, United States, weeks ending September 19, 2009, and September 13, 2008 (37th)*

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

U: Unavailable. —: No reported cases. N: Not reportable. Cum: Cumulative year-to-date counts. Med: Median. Max: Maximum. * Incidence data for reporting year 2009 is provisional. Data for HIV/AIDS, AIDS, and TB, when available, are displayed in Table IV, which appears quarterly. † Updated weekly from reports to the Division of Vector-Borne Infectious Diseases, National Center for Zoonotic, Vector-Borne, and Enteric Diseases (ArboNET Surveillance).

Data for California serogroup, eastern equine, Powassan, St. Louis, and western equine diseases are available in Table I.

[§] Not reportable in all states. Data from states where the condition is not reportable are excluded from this table, except starting in 2007 for the domestic arboviral diseases and influenza-associated pediatric mortality, and in 2003 for SARS-CoV. Reporting exceptions are available at http://www.cdc.gov/epo/dphsi/phs/infdis.htm. ¹ Contains data reported through the National Electronic Disease Surveillance System (NEDSS).

TABLE III. Deaths in 122 U.S. cities,* week ending September 19, 2009 (37th)

	All causes, by age (years)								All causes, by age (years)						
Reporting area	All Ages	≥65	45–64	25–44	1–24	<1	P&I [†] Total	Reporting area	All Ages	≥65	45–64	25–44	1–24	<1	P&l [†] Total
New England	507	352	109	27	13	6	36	S. Atlantic	1,289	783	341	95	38	32	72
Boston, MA	144	89	38	8	6	3	9	Atlanta, GA	179	97	56	14	11	1	7
Bridgeport, CT	26	20	6	-	_	_	2	Baltimore, MD	136	64 71	50	13	2	7	12
Fall River MA	20	12	2	1	_	1	2		153	0/	29	9 17	1	3	12
Hartford CT	51	32	15	4	_	_	4	Miami Fl	62	42	15	4	1	_	5
Lowell, MA	25	17	6	1	1	_	3	Norfolk, VA	48	33	7	4	2	2	_
Lynn, MA	11	7	3	1	_	_	1	Richmond, VA	65	37	20	2	4	2	3
New Bedford, MA	22	20	1	1	_	_	1	Savannah, GA	63	47	9	2	2	3	2
New Haven, CT	22	17	3	_	2	_	3	St. Petersburg, FL	53	37	10	3	1	2	8
Providence, RI	57	38	15	2	2	—	6	Tampa, FL	228	151	52	17	4	4	14
Somerville, MA	2	1	_	1	—	—	_	Washington, D.C.	168	102	51	8	5	2	4
Springfield, MA	22	16	4	1		1	1	Wilmington, DE	15	8	4	2		1	1
Waterbury, CT	30	20	6	2	2	_	2	E.S. Central	802	507	202	65	21	7	69
Worcester, MA	60	48	/	4		1	2	Birmingham, AL	170	99	52	15	3	1	/
Mid. Atlantic	2,119	1,450	481	106	38	37	92	Chattanooga, TN	100	39	8	4	2	-	10
Albariy, NT Allentown PA	52 24	18	9	1	_	0	_	Levington KV	54	30	24 12	2	4		12
Buffalo NY	61	41	17	2	1	_	1	Memphis TN	190	118	45	21	5	1	23
Camden, NJ	19	9	3	2	2	3	1	Mobile, AL	73	53	14	4	1	1	3
Elizabeth, NJ	16	11	3	1	_	1	_	Montgomery, AL	34	22	6	4	1	1	2
Erie, PA	54	45	5	4	_	_	7	Nashville, TN	126	70	41	9	4	2	9
Jersey City, NJ	29	18	9	2	—	—	1	W.S. Central	1,434	865	383	109	50	27	66
New York City, NY	1,020	704	236	48	19	12	51	Austin, TX	73	48	19	3	1	2	4
Newark, NJ	40	22	14	3	—	1	4	Baton Rouge, LA	73	46	15	10		2	_
Paterson, NJ	5	5		_	_	_	_	Corpus Christi, TX	58	32	21	3	1	1	3
Philadelphia, PA	396	250	104	24	9	9	4	Dallas, IX	202	104	66	18	11	3	10
Pittsburgh, PA ³	50	30	13	6	_	1	3	El Paso, TX	103	/2	27	3			3
Reading, PA	122	31	26	5	2	2	10	Houston TY	200	229	107	26	17	11	10
Schenectady NY	26	22	20	2		_	1	Little Bock AB	72	220	23	7	3	<u> </u>	19
Scranton, PA	22	18	3	1	_	_	2	New Orleans, LA	Ű	Ŭ	Ŭ	Ú	ŭ	U	Ŭ
Syracuse, NY	80	61	18		1	_	5	San Antonio, TX	224	150	58	5	8	3	17
Trenton, NJ	18	13	4	_	1	_	_	Shreveport, LA	71	43	14	7	2	5	3
Utica, NY	24	21	3	_	_	—	2	Tulsa, OK	159	103	33	17	6	_	4
Yonkers, NY	12	9	1	1	1	—	_	Mountain	1,122	706	284	84	22	26	58
E.N. Central	1,727	1,126	405	97	36	63	99	Albuquerque, NM	136	99	24	9	2	2	12
Akron, OH	44	28	11	2	1	2	3	Boise, ID	51	30	15	2	1	3	2
Canton, OH	40	34	5			1	3	Colorado Springs, CO	101	57	30	6	4	4	2
Chicago, IL	0	U 40	10	U	0	U	U	Denver, CO	79	38	28	6	3	4	3
	211	49	10	14	3	2	10	Cadon LIT	200	103	73	20 1	э	_	10
Columbus OH	229	137	56	14	5	13	12	Phoenix AZ	157	29 76	58	17	_	6	6
Davton, OH	135	98	28	5	1	3	8	Pueblo, CO	26	15	6	1	4	_	1
Detroit, MI	161	84	46	15	6	10	7	Salt Lake City, UT	92	69	11	5	3	4	6
Evansville, IN	51	30	21	_	_	_	4	Tucson, AZ	158	110	33	12	_	3	5
Fort Wayne, IN	68	44	15	5	2	2	1	Pacific	1,627	1,093	385	90	38	21	156
Gary, IN	13	5	6	2	—	—	_	Berkeley, CA	13	9	4	—	_	—	3
Grand Rapids, MI	50	37	7	4	1	1	5	Fresno, CA	113	67	37	7	1	1	8
Indianapolis, IN	188	118	53	8	4	5	11	Glendale, CA	25	18	6	1		_	5
Lansing, Mi	34	26	5	2	1		1	Honolulu, HI	68	48	14	2	3	1	6
Nilwaukee, WI	104	20	28	3	∠ 1	4	0	Long Beach, CA	242	171	10	0	11	2	42
Rockford II	43 56	29 41	13		1	1	6	Pasadena CA	243	16	49	2			42
South Bend IN	56	38	8	4	4	2	4	Portland OB	97	61	25	7	1	3	10
Toledo, OH	98	68	23	3	1	3	6	Sacramento, CA	163	112	38	8	4	1	16
Youngstown, OH	62	53	7	1	_	1	4	San Diego, CA	240	154	62	16	7	1	18
W.N. Central	623	387	159	45	14	17	35	San Francisco, CA	109	66	29	7	4	3	11
Des Moines, IA	63	44	13	3	1	2	6	San Jose, CA	222	167	41	8	2	4	18
Duluth, MN	34	24	8	2	_	—	3	Santa Cruz, CA	30	20	6	4	—	_	1
Kansas City, KS	30	15	14	1	—	—	3	Seattle, WA	122	76	28	12	3	3	9
Kansas City, MO	82	50	18	5	4	5	6	Spokane, WA	57	41	15	_	_	1	4
Lincoln, NE	33	30	3	_	_	—	1	Tacoma, WA	103	67	27	7	2		3
Minneapolis, MN	54	34	15	3	2	_	2	Iotal™	11,250	7,275	2,749	718	270	236	683
Omana, NE St. Louis MO	105	64	23	13	3	2	5	1							
St. Louis, MO	00 65	41	20 22	2	3	3	2	1							
Wichita. KS	77	48	17	9	1	1	2								

U: Unavailable. —:No reported cases. * Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of >100,000. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included. * Pneumonia and influenza.

⁵ Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks. ¹ Total includes unknown ages.

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