



# **Morbidity and Mortality Weekly Report**

Weekly June 14, 2002 / Vol. 51 / No. 23

# West Nile Virus Activity — United States, 2001

In 2001, West Nile virus (WNV) activity was reported from 359 counties in 27 states and the District of Columbia (DC) to ArboNET, a web-based, surveillance data network maintained by 54 state and local public health agencies and CDC. This activity represented a marked increase from 2000, when WNV activity was reported from 138 counties in 12 states and DC (1). This report summarizes surveillance data for 2001, which indicate that 66 human illnesses were reported from 10 states and that widespread WNV activity in birds, horses, and mosquitoes extended into the midwestern United States and several southern states unaffected previously. The findings in this report underscore the need for public education, increased WNV surveillance aimed at early viral detection, and sustained, integrated mosquito-control activities.

In 2001, CDC conducted WNV surveillance with 54 ArboNET surveillance coordinators from health departments in the contiguous 48 states and six jurisdictions (Chicago, DC, Houston, Los Angeles, New York City, and Philadelphia). Local WNV surveillance networks collected and tested for WNV or antibodies specimens from human and veterinary patients, dead birds, captive sentinel animals (mostly chickens), wild-caught birds, and mosquitoes. Test results, including county and week of specimen collection or illness onset, were entered into local electronic databases, and standardized summaries were forwarded weekly to CDC's ArboNET database system. In addition, reports of human WNV cases and other reports of WNV activity were reported to CDC by telephone, facsimile, or e-mail.

In 2001, a total of 66 human cases of WNV disease (64 persons with central nervous system infections [WNV meningoencephalitis] and two persons with uncomplicated WNV fever) were reported from 39 counties in 10 states (Figure 1). New York (13 WNV meningoencephalitis cases; two WNV fever cases), New Jersey (12 WNV meningoencephalitis cases), and Florida (12 WNV meningoencephalitis cases) accounted

for 39 (59%) reported cases. Among 64 persons with WNV meningoencephalitis, the median age was 68 years (range: 9–90 years). Nine (14%) cases were fatal; the median age of these persons was 70 years (range: 44–90 years). The dates of human illness onset ranged from July 13 to December 7 (Figure 2). In 36 (92%) counties reporting human cases, the first case was preceded by at least one report of a WNV-infected bird, sentinel animal, horse, or mosquito pool; 320 counties detected enzootic WNV activity but no human infections.

Of the 359 counties reporting WNV activity, 328 (91%) counties in 27 states and DC reported 7,333 dead WNVinfected birds (5,154 crows from two Corvus species, 966 blue jays, and 1,213 birds from 71 other avian species). In 238 (66%) counties, dead crows were the first indicators of WNV activity. Of 9,679 crows tested for WNV, 5,154 (53%) were positive for WNV infection compared with 2,179 (9%) of 24,898 birds from other species. Dead infected birds were collected during April 4-December 26. A total of 55 seropositive wild-caught birds were reported from DC and five counties in three states (Florida, New York, and Ohio) and represented the first detection of WNV activity in two of these counties. A total of 218 seroconverting captive sentinel animals were reported from 26 counties in five states (Florida, New Jersey, New York, North Carolina, and Virginia). In four Florida counties, seroconverting sentinel chickens were the first sign of WNV activity.

# **INSIDE**

- 501 Disseminated Infection with Simiae-Avium Group Mycobacteria in Persons with AIDS — Thailand and Malawi, 1997
- 503 Update: Influenza Activity United States and Worldwide, 2001–02 Season, and Composition of the 2002–03 Influenza Vaccine

The MMWR series of publications is published by the Epidemiology Program Office, Centers for Disease Control and Prevention (CDC), U.S. Department of Health and Human Services, Atlanta, GA 30333.

### **SUGGESTED CITATION**

Centers for Disease Control and Prevention. [Article Title]. MMWR 2002;51:[inclusive page numbers].

# **Centers for Disease Control and Prevention**

David W. Fleming, M.D. *Acting Director* 

Julie L. Gerberding, M.D. Acting Deputy Director for Science and Public Health

> Dixie E. Snider, Jr., M.D., M.P.H. Associate Director for Science

# **Epidemiology Program Office**

Stephen B. Thacker, M.D., M.Sc. *Director* 

#### Office of Scientific and Health Communications

John W. Ward, M.D.

Director

Editor, MMWR Series

David C. Johnson
Acting Managing Editor, MMWR (Weekly)

Jude C. Rutledge Teresa F. Rutledge Jeffrey D. Sokolow, M.A. Writers/Editors, MMWR (Weekly)

Lynda G. Cupell Malbea A. Heilman Beverly J. Holland Visual Information Specialists

Quang M. Doan Michele D. Renshaw Erica R. Shaver Information Technology Specialists

# Division of Public Health Surveillance and Informatics

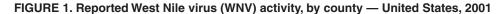
# Notifiable Disease Morbidity and 122 Cities Mortality Data

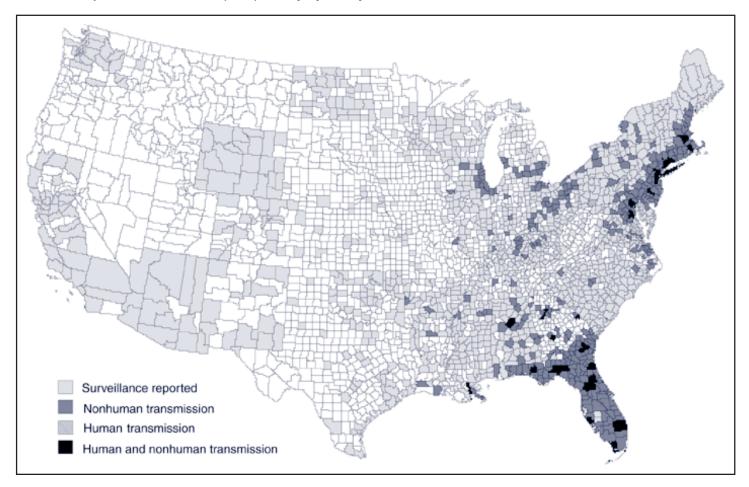
Robert F. Fagan Deborah A. Adams Felicia J. Connor Lateka Dammond Patsy A. Hall Pearl C. Sharp Horses were the only WNV-infected nonhuman mammals reported in 2001. A total of 733 equine cases were reported from 127 counties in 19 states (Alabama, Connecticut, Delaware, Florida, Georgia, Illinois, Indiana, Kentucky, Louisiana, Maryland, Massachusetts, Mississippi, New Hampshire, New Jersey, New York, North Carolina, Pennsylvania, Tennessee, and Virginia); this represented a 12-fold increase compared with 2000 (1). Florida reported 483 equine cases (66% of all reports) from 40 counties. The first equine illness preceded the first human illness; equine illness onset dates ranged from June 27 to December 18.

In 2001, a total of 564 counties conducted WNV testing on approximately 1.4 million mosquitoes from 91 species. WNV was detected in 919 mosquito pools (27 species) reported from 71 counties in 16 states (Connecticut, Delaware, Florida, Georgia, Illinois, Kentucky, Maryland, Massachusetts, Michigan, New Hampshire, New Jersey, New York, Ohio, Pennsylvania, Rhode Island, and Virginia) and DC. As in 2000, two enzootic vector species, *Culex pipiens* and *Cx. restuans*, collectively accounted for the majority (59%) of WNV-positive pools. WNV also was found for the first time in several additional species of potential public health importance, including *Anopheles quadrimaculatus*, *Coquilletidia perturbans*, *Cx. nigripalpus*, *Cx. quinquefasciatus*, *Ochlerotatus sollicitans*, *Oc. taeniorhynchus*, and *Psorophora columbiae*.

**Reported by:** DR O'Leary, DVM, RS Nasci, PhD, GL Campbell, MD, AA Marfin, MD, Div of Vector-Borne Infectious Diseases, National Center for Infectious Diseases, CDC.

**Editorial Note:** The 2001 surveillance data indicate that the geographic area with WNV activity has increased and that dead WNV-infected birds were reported in western Arkansas, southern Maine, and southern Florida. Seven states reported human cases for the first time. Canadian health authorities also detected WNV activity in dead birds from southwestern Ontario, a region of lower latitude than the northern limits of WNV detection in the United States (2). In 2001, a case of WNV encephalitis was diagnosed serologically in a resident of the Cayman Islands who had no recent travel history (CDC, unpublished data, 2001), suggesting that WNV has entered the Caribbean region. Human illness onsets on July 13 and December 7 in persons in Florida and Georgia, respectively, mark the earliest and latest reported human cases since the introduction of WNV to the United States. Extended seasonal activity in 2001 occurred in the northeast; two of five persons with illness onset on October 15 or later were from Massachusetts. The widespread occurrence of human cases and the occurrence of human cases outside of WNV's usual season (summer and early fall) suggest that 1) state and local health departments in the contiguous 48 states should, at a minimum, establish enhanced passive hospital-based surveillance



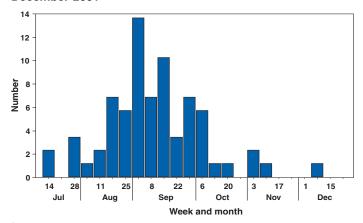


for human cases of encephalitis of unknown etiology and 2) this surveillance should extend beyond mid-October (3).

In 2001, infected birds, mosquitoes, or horses were detected in 16 states in which no previous WNV activity in animals had been reported. These findings demonstrate the dramatic spread of WNV westward and southward since 1999, when the virus was first recognized in North America. Although virus activity was detected for the first time in many southern states in 2001, the simultaneous appearance of two epizootic foci very early in 2001—one in the mid-Atlantic region and one in the southeast along the common borders of Florida, Georgia, and Alabama—suggests that WNV was introduced into the southern states by migrating birds in late 2000, but circulated at levels below the detection threshold of surveillance. Because many migratory bird species use well-established north-south flyways along the eastern seaboard, this movement of WNV from the mid-Atlantic region to the south Atlantic region and the Gulf states was expected; however, the reasons for WNV's rapid expansion into multiple foci in the central United States are less obvious. Possible mechanisms include carriage of the virus by the return of infected birds from wintering sites in southern states or by their incremental east-to-west local movements (4).

Surveillance of dead birds is essential in monitoring WNV activity. Infection in species within the family Corvidae (e.g., crows and jays) is a particularly important indicator of WNV activity. In 2001, the proportion of tested birds that were infected continued to be disproportionately higher in crows than all other birds (53% versus 9%). Although 83% of infected birds reported were either crows or blue jays, this might be attributed to greater emphasis placed by states on monitoring these species. State and local health department surveillance programs should continue to emphasize the collection and testing of dead corvids. However, because noncorvid birds were first indicators of WNV activity in 57 (16%) of 359 counties where the virus was detected, surveillance programs should include these other species wherever possible. In 2001, serosurveillance of sentinel chickens and wild-caught birds contributed additional information on WNV transmission and provided collectively the initial

FIGURE 2. Number\* of human West Nile virus disease cases, by week and month of illness onset — United States, July–December 2001



\* n=66.

signal of WNV activity in six counties. The limitations of these supplemental systems are documented (3,5) and their overall utility continues to be evaluated.

The 2001 equine WNV epizootic was unprecedented given its geographic span and the number of horses affected. In addition to a substantial epizootic in the northeast, an intense equine epizootic in Georgia and Florida accounted for 75% (551) of all reported equine cases. Scattered cases also were detected as far west as Louisiana, in the Ohio valley, and in northern Illinois. In August 2001, the U.S. Department of Agriculture granted conditional licensure of a commercial equine WNV vaccine because of the detrimental effect of these events on equine health and industry. Because WNV-infected horses are unlikely to develop viremias sufficient to infect feeding mosquitoes, they are unlikely to pose a risk to humans (6). However, equine epizootics reflect intense enzootic WNV activity in mosquitoes, which might place humans at increased risk.

In 2001, Culex mosquitoes (Cx. pipiens, Cx. restuans, and Cx. salinarius) were the most commonly identified mosquito vectors of WNV in the United States, and since 1999 these species have been found in close spatial and temporal proximity to the majority of human cases of WNV meningoencephalitis (1,7,8). Detection of WNV in several common human-feeding mosquito species (e.g., Cx. nigripalpus, Oc. sollicitans, Oc. taeniorhynchus, and Cq. perturbans) and recent studies demonstrating their ability to transmit this virus under laboratory conditions (9,10) raise concerns about increased human risk in areas where these species are common.

The data available to the ArboNET system likely underestimate actual geographic distribution and intensity of WNV virus transmission in the United States. Data provided by the 54 ArboNET coordinators are derived largely from local health

unit surveillance efforts, which vary according to capacity and ability. The 28 jurisdictions reporting activity probably support additional, undetected WNV transmission within their borders, and undetected foci of transmission probably exist in counties and states that have not reported transmission activity. In addition, some detected WNV infections might not have been entered into the ArboNET system.

In Florida, epizootic WNV activity has been reported since January 2002, indicating that year-round transmission is occurring in that state. In northern states, WNV activity has been reported since April. The extended seasonal activity, the broad vertebrate host and vector-mosquito range, and the establishment of multiple epizootic foci throughout the eastern United States demonstrate that WNV has established itself permanently in temperate North America and strongly suggest that it will spread further westward. This underscores the need for increased surveillance geared toward early viral detection and mosquito-control activities that weaken or break amplification cycles and decrease the risk for human and domestic animal infection with WNV. Prevention activities should continue to include 1) public education programs urging residential source reduction and personal protective measures to reduce mosquito exposure; 2) development of sustained, community-level integrated mosquito-surveillance and management programs (3); and 3) high-priority emphasis on the control of urban Culex mosquitoes.

## **Acknowledgments**

This report is based on data prepared by ArboNET surveillance coordinators in local and state health departments and ArboNET technical staff, Div of Vector-Borne Infectious Diseases, National Center for Infectious Diseases, CDC.

#### References

- Marfin AA, Petersen LR, Eidson ME, et al. Widespread West Nile virus activity, eastern United States, 2000. Emerg Infect Dis 2001;7:730-1.
- 2. Health Canada. West Nile virus surveillance 2002: Canada. Infectious Diseases News Brief. Centre for Infectious Disease Prevention and Control, Population and Public Health Branch, Health Canada. June 7, 2002. Available at http://www.hc-sc.gc.ca/pphb-dgspsp/bid-bmi/dsd-dsm/nb-ab/2002/nb2302\_e.html
- 3. CDC. Update: epidemic/epizootic West Nile virus in the United States: revised guidelines for surveillance, prevention, and control, 2001. Available at http://webdev.cdc.gov/ncidod/dvbid/westnile/resources/wnvguidelines-apr-2001.pdf.
- 4. Rappole JH, Derrickson SR, Hubalek Z. Migratory birds and spread of West Nile virus in the western hemisphere. Emerg Infect Dis 2000;6:319–28.
- Komar N. West Nile virus surveillance using sentinel birds. Ann NY Acad Sci 2001;951:58–73.
- Bunning ML, Bowen RA, Cropp CB, et al. Experimental infection of horses with West Nile virus. Emerg Infect Dis 2002;8:380–6.
- Kulasekera VL, Kramer L, Nasci R, et al. West Nile virus infection in mosquitoes, birds, horses, and humans, Staten Island, New York, 2000. Emerg Infect Dis 2001;7:722–5.

- 8. Nasci RS, White DJ, Stirling H, et al. West Nile virus isolates from mosquitoes in New York and New Jersey, 1999. Emerg Infect Dis 2001;7:626–9.
- Sardelis MR, Turrell MJ, Dohm DJ, et al. Vector competence of selected North American *Culex* and *Coquillettidia* mosquitoes for West Nile virus. Emerg Infect Dis 2001;7:1018–22.
- Turrell MJ, O'Guinn ML, Dohm JD, et al. Vector competence of North American mosquitoes (Diptera: Culicidae) for West Nile virus. J Med Entomol 2001;38:130–4.

# Disseminated Infection with Simiae-Avium Group Mycobacteria in Persons with AIDS — Thailand and Malawi, 1997

Persons with advanced human immunodeficiency virus (HIV)-1 infection are susceptible to disseminated mycobacterial infections. In the United States, most such infections are caused by Mycobacterium avium or M. intracellulare (i.e., M. avium complex [MAC]). In less developed countries, M. tuberculosis is equally or more prevalent than MAC in persons with HIV-1 infection (1). Other mycobacterial species have been reported to cause disseminated infection in HIVinfected persons, including Simiae-Avium (SAV) group mycobacteria. SAV group organisms share characteristics of M. avium and M. simiae (2). Although disseminated (i.e., the isolation of a mycobacterial species from the blood) infection with M. simiae has been reported in HIV-infected persons (3–6), another distinct species within the SAV group, M. triplex, was characterized in 1996 (7). Two cases of disseminated infection caused by M. triplex have been reported in HIV-1positive persons (8,9). This report describes four HIV-infected patients from Bangkok, Thailand, and Lilongwe, Malawi, who were infected with SAV group organisms. Because different mycobacterial species are not susceptible uniformly to antimycobacterial agents, accurate identification of mycobacterial species causing an infection is crucial for directing appropriate therapy.

These infections were detected during prospective blood culture studies of febrile, adult inpatients in these two countries (1,10). The Bangkok study was conducted at an infectious diseases hospital during February–March 1997 (10); the Lilongwe study was conducted at a general hospital during August–September 1997 (1). In both studies, adults (aged  $\geq 18$  years) admitted consecutively with fever (oral temperature  $\geq 100^{\circ}$  F [ $\geq 38^{\circ}$  C] in Bangkok and axillary temperature  $\geq 99^{\circ}$  F [ $\geq 37.5^{\circ}$  C] in Malawi) were recruited within 12 hours of hospital admission. After informed consent was obtained, patients gave a full medical history and underwent a comprehensive physical examination. Blood was drawn for HIV-1

testing and mycobacterial culture. All mycobacterial isolates were sent to Duke University Medical Center for confirmation and identification. M. tuberculosis complex and M. avium complex isolates were identified by using AccuPROBE (Gen-Probe, San Diego, California) DNA probes and biochemical tests. Isolates of uncommon Mycobacterium spp. (e.g., M. simiae) were sent to North Carolina State Public Health Laboratory and the Mycobacteria Reference Laboratory at CDC for further characterization and confirmation by high performance liquid chromatography analysis of mycolic acids. Personnel at both laboratories read all chromatograms visually. Susceptibilities of the isolates to antituberculous drugs were performed at CDC using methodology established for M. tuberculosis. Of 480 patients evaluated, four (two from Bangkok and two from Lilongwe) were found to have disseminated infection with SAV group mycobacteria, later identified as M. simiae.

# Bangkok, Thailand

Both patients had positive serology for HIV-1 antibody. Neither was receiving antiretroviral or antimycobacterial therapy. Patient 1, a man aged 32 years, presented with fever, cachexia, and diarrhea of 3 months' duration. Physical examination revealed oral candidiasis and lymphadenopathy. Patient 2, a man aged 36 years, presented with fever, cachexia, and cough and shortness of breath of 1 weeks' duration. Physical examination revealed lymphadenopathy. Additional laboratory studies on this patient revealed hematocrit 16% (normal: 39%–49%) and positive cerebrospinal fluid cryptococcal antigen. Both patients were treated with broad-spectrum antimicrobials for possible underlying bacterial infection and were discharged from the hospital.

# Lilongwe, Malawi

Both patients had positive serology for HIV-1 antibody. Neither was receiving antiretroviral or antimycobacterial therapy. Patient 3, a man aged 28 years, presented with chronic fever and cough of 7 months' duration. Physical examination revealed cachexia and skin lesions. No lymphadenopathy was noted. Patient 4, a man aged 36 years, presented with fever, chronic fever, and diarrhea of 5 months' duration. Physical examination revealed oral candidiasis. No lymphadenopathy was detected. Both patients were treated with penicillin and chloramphenicol for underlying bacterial infection and were discharged from the hospital.

# **Susceptibility testing**

All four isolates were available for susceptibility testing. These isolates were resistant to all first-line drugs (isoniazid,

rifampin, streptomycin, ethambutol, and pyrazinamide) used for treating *M. tuberculosis* infection and to alternative drugs (e.g., kanamycin and ciprofloxacin) used for treating atypical mycobacteria and multidrug-resistant tuberculosis (MDR-TB).

Reported by: LB Reller, Clinical Microbiology Laboratory, Duke Univ Medical Center, Durham, North Carolina. LK Archibald, MD, WR Jarvis, MD, Div of Healthcare Quality Promotion; Div of AIDS, STD, and TB Laboratory Research, National Center for Infectious Diseases; LA Grohskopf, MD, EIS Officer, CDC.

Editorial Note: Advances in laboratory methodology have enabled more rapid and reliable differentiation of mycobacterial species commonly associated with clinical illness (e.g., *M. tuberculosis* and MAC), and the identification of new or emerging species (e.g., *M. triplex*). However, ambiguities in determining specific mycobacteria species might occur in regions of the world where diagnostic resources are limited or not available. In addition, no standard susceptibility testing panel has been established for these organisms. These limitations might lead to difficulties in the clinical management of patients with disseminated mycobacterial infection.

The clinical manifestations of disseminated mycobacterial infection are nonspecific and are not indicative of the infecting species. Therefore, as with other mycobacterial infections, diagnosis and specific therapy should be guided by laboratory testing, including species identification and susceptibility testing whenever possible, rather than clinical findings alone.

The findings in this report are subject to at least three limitations. First, neither CD4 lymphocyte nor HIV-1 viral load data were obtained. However, because each patient had a marker of symptomatic HIV-1 infection (oral candidiasis, Kaposi's sarcoma, or positive cerebrospinal fluid cryptococcal antigen), all probably had clinical evidence of advanced immune deficiency. Second, because these patients had multiple conditions that could have produced their nonspecific symptoms and physical findings, it is unclear whether SAV mycobacteria were the cause of their symptoms. Further study and characterization of the SAV group of mycobacteria and of the clinical illness with which they are associated are required to better ascertain the prevalence and clinical significance of these mycobacterial infections. Finally, no information was available on treatment or postdischarge outcome for these patients.

Awareness of *M. simiae* and other SAV mycobacteria as potential causes of disseminated infection in patients with AIDS is important for several reasons. Because of the phenotypic similarity between SAV mycobacteria and other mycobacterial species, patients infected with SAV mycobacteria might go unrecognized and be presumed to be infected with other *Mycobacterium* species (e.g., *M. tuberculosis*), particularly

in resource-poor settings without access to adequate laboratory testing. This might lead to ineffective treatment, because not all species are susceptible to all agents. Also, if these isolates were assumed to be *M. tuberculosis*, they could be misclassified as MDR-TB.

Because of the lack of data and of clinical experience with *M. simiae* and other SAV group mycobacteria, the best treatment is unknown. Infections with other mycobacteria, particularly *M. tuberculosis*, require treatment for prolonged periods with multiple agents to which the organisms are susceptible; not adhering to these principles promotes the development of drug-resistant organisms. Additional investigation is needed to determine whether similar hazards exist when SAV mycobacteria are treated with ineffective agents or otherwise suboptimal therapy.

### **Acknowledgments**

This report is based on data contributed by S Tansuphasawadikul, B Eampokalap, A Chaovavanich, Bamrasnaradura Hospital, Nonthaburi; S Rheanpumikankit, Field Epidemiology Training Program, Ministry of Health, Thailand. P Kazembe, O Nwanyanwu, H Dobbie, Lilongwe Central Hospital, Lilongwe; Ministry of Health, Malawi. LF Turner, North Carolina Dept of Health and Human Svcs, State Laboratory of Public Health, Raleigh, North Carolina.

#### References

- Archibald LK, McDonald LC, Nwanyanwu O, et al. A hospital-based prevalence survey of bloodstream infections in febrile patients in Malawi: implications for diagnosis and therapy. J Infect Dis 2000;181:1414–20.
- Tortoli E, Piersimoni C, Kirschner P, et al. Characterization of mycobacterial isolates related to, but different from, *Mycobacterium simiae*. J Clin Microbiol 1997;35:697–702.
- Levy-Frebault V, Pangon B, Bure A, et al. Mycobacterium simiae and Mycobacterium avium—M. intracelluare mixed infection in acquired immune deficiency syndrome. J Clin Microbiol 1987;25:154–7.
- Torres RA, Nord J, Feldman R, et al. Disseminated mixed Mycobacterium simiae—Mycobacterium avium complex infection in acquired immune deficiency syndrome. J Infect Dis 1991;164:432–3.
- 5. Munier D, Dux S, Samra Z, et al. *Mycobacterium simiae* infection in Israeli patients infected with AIDS. Clin Infect Dis 1993;17:508–9.
- 6. Koeck JL, Debord T, Fabre M, et al. Disseminated *Mycobacterium simiae* infection in a patient with AIDS: clinical features and treatment. Clin Infect Dis 1996;23:832–3.
- Floyd MM, Guthertz LS, Silcox VA, et al. Characterization of an SAV organism and proposal of *Mycobacterium triplex* sp. nov. J Clin Microbiol 1996;34:2963–7.
- 8. Cingolani A, Sanguinetti M, Antinori A, et al. Disseminated mycobacteriosis caused by drug-resistant *Mycobacterium triplex* in a human immunodeficiency virus-infected patient during highly active antiretroviral therapy. Clin Infect Dis 2000;31:177–9.
- 9. Hoff E, Sholtis M, Procop G, et al. *Mycobacterium triplex* infection in a liver transplant patient. J Clin Microbiol 2001;2033–4.
- 10. Archibald LK, McDonald LC, Rheanpumikankit S, et al. Fever and human immunodeficiency virus infection as sentinels for emerging mycobacterial and fungal bloodstream infections in hospitalized patients ≥15 years old, Bangkok. J Infect Dis 1999;180:87–92.

# Update: Influenza Activity — United States and Worldwide, 2001–02 Season, and Composition of the 2002–03 Influenza Vaccine

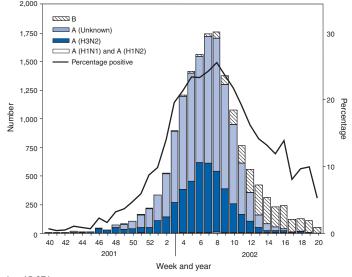
The 2001–02 influenza season in the United States was mild to moderate. Influenza A (H3N2) viruses predominated, but influenza B viruses were identified more frequently than influenza A viruses toward the end of the season. Worldwide, influenza A (H3N2) and B viruses predominated. This report summarizes influenza activity in the United States\* (September 30, 2001–May 18, 2002) and worldwide (October–May) during the 2001–02 influenza season and describes the composition of the 2002–03 influenza vaccine.

## **United States**

Influenza activity increased in mid-January and peaked during mid-to-late February. Influenza A (H3N2) viruses predominated; however, the number of influenza B viruses increased as the season progressed. Influenza B viruses were the most frequently identified influenza viruses from the week ending March 30 (week 13) through the week ending May 18 (week 20) (Figure).

During September 30, 2001–May 18, 2002, the World Health Organization and National Respiratory and Enteric

FIGURE. Number\* and percentage of respiratory specimens testing positive for influenza reported by World Health Organization and National Enteric and Respiratory Virus Surveillance System collaborating laboratories, by week and year — United States, 2001–02 season†



<sup>\*</sup> n=15,671.

Virus Surveillance System collaborating laboratories in the United States tested 100,815 respiratory specimens for influenza; 15,671 (16%) were positive (Figure). Of these, 13,706 (87%) were influenza type A and 1,965 (13%) were influenza type B. Of the 4,507 subtyped influenza A viruses, 4,420 (98%) were H3 viruses and 87 (2%) were H1 viruses. Influenza A viruses predominated in all nine surveillance regions. The proportion of specimens testing positive for influenza first increased to ≥10% during the week ending January 12 (week 2), peaked at 26% during the week ending February 23 (week 8), and declined to <10% during the week ending April 27 (week 17). The peak percentage of specimens testing positive for influenza during the previous three seasons ranged from 23% to 31%.

CDC has characterized antigenically 690 influenza viruses collected from U.S. laboratories since September 30: 393 influenza A (H3N2) viruses, 30 influenza A H1 viruses, and 267 influenza B viruses. All 393 influenza A (H3N2) viruses were similar to A/Panama/2007/99, the H3N2 component of the 2001–02 influenza vaccine. All 30 influenza A H1 viruses had an H1 protein similar antigenically to the A/New Caledonia/20/99, the H1N1 component of the 2001–02 influenza vaccine. Sixteen of the 30 H1 viruses were identified as influenza A (H1N2) viruses. These H1N2 viruses were collected in Hawaii, Massachusetts, New York, Pennsylvania, and Wisconsin. Two additional H1N2 viruses were identified from patient specimens collected during July and September in Texas and Nevada, respectively.

Influenza B viruses circulating currently can be divided into two antigenically distinct lineages: B/Yamagata/16/88-like and B/Victoria/2/87-like viruses. CDC has characterized antigenically 267 influenza B viruses collected from U.S. laboratories since September 30; 61 were of the B/Yamagata lineage and 206 of the B/Victoria lineage. Of the 61 B/Yamagata lineage viruses, 13 were similar to the vaccine strain B/Sichuan/379/99, and 48 demonstrated reduced titers to ferret antisera produced against B/Sichuan/379/99. Most of the viruses that demonstrated reduced titers to ferret antisera produced against B/Sichuan/379/99 were related closely to B/Shizuoka/15/01, a minor antigenic variant of B/Sichuan/379/99.

During the weeks ending December 29–January 5 (weeks 52–1) and the weeks ending January 19–March 16 (weeks 3–11), the weekly percentage of patient visits for influenza-like illness (ILI)<sup>†</sup> reported by U.S. sentinel physicians<sup>§</sup> exceeded

<sup>\*</sup>Data reported as of June 10, 2002. The four components of the influenza surveillance system have been described previously (1).

Data reported as of June 10, 2002.

 $<sup>^{\</sup>dagger}$  Temperature ≥100° F (≥37.8° C) and either cough or sore throat in the absence of a known cause.

The national baseline was calculated as the mean percentage of patient visits for ILI during noninfluenza weeks plus two standard deviations. Because of wide variability in regional-level data, calculating region-specific baselines is not possible, and the national baseline cannot be applied to regional level data.

baseline levels (0–1.9%). The peak percentage of patient visits for ILI was 3.2% during the week ending February 16 (week 7). During the previous three seasons, the peak percentage of patient visits for ILI ranged from 4% to 6%.

On the basis of data from state and territorial epidemiologists, influenza activity peaked during the week ending February 23 (week 8), when 40 states reported regional or widespread influenza activity. Regional influenza activity was reported by one or more states during all but 1 week from the week ending October 27 (week 43) through the week ending May 4 (week 18). Widespread activity was reported by one or more states during all but 1 week from the week ending December 1 (week 48) through the week ending March 30 (week 13). The peak number of states reporting regional or widespread activity during the previous three seasons ranged from 38 to 44.

As reported by the 122 Cities Mortality Reporting System, the percentage of deaths in the United States associated with pneumonia and influenza (P&I) exceeded the epidemic threshold\*\* for 5 consecutive weeks (weeks ending March 2 [week 9] to March 30 [week 13]). During the previous three seasons, the number of consecutive weeks during which the percentage of deaths attributed to P&I exceeded the epidemic threshold ranged from 0 to 13.

### Worldwide

During October 2001–May 2002, influenza A (H3N2) and B viruses circulated widely in Africa, the Americas, Asia, Europe, and Oceania, and influenza A (H1N1) and A (H1N2) viruses were reported sporadically. Influenza A (H3N2) viruses predominated in Africa (Egypt, Madagascar, Mauritus, Senegal, and Tunisia), the Americas (Argentina, Canada, French Guiana, and Mexico), Asia (China and Hong Kong), Europe (Croatia, Czech Republic, Denmark, Finland, France, Germany, Iceland, Ireland, Israel, Latvia, the Netherlands, Norway, Portugal, Romania, Spain, and the United Kingdom),

and Oceania (Australia). Influenza A (H3N2) viruses also were reported from Asia (Japan, the Philippines, the Republic of Korea, Singapore, Taiwan, and Thailand) and Europe (Bulgaria, Greece, Italy, Poland, the Russian Federation, and Switzerland), and Oceania (New Zealand).

Influenza B viruses were identified more frequently than influenza A viruses in Austria, Chile, Greece, Italy, India, Paraguay, the Russian Federation, Slovakia, Slovenia, and Switzerland. Many of the influenza B viruses from the Americas, Asia, Europe, and Oceania were B/Sichuan/379/99-like viruses that were identified in Argentina, Australia, Canada, Croatia, France, Greece, Hong Kong, India, Israel, Italy, Malaysia, Norway, the Philippines, the Russian Federation, Singapore, Slovakia, Thailand, Taiwan, and the Ukraine. However, the majority of influenza B isolates from Canada, Hong Kong, India, Oman, and the Philippines were B/Hong Kong/330/01-like viruses; these viruses also were identified in China, Israel, Italy, Malaysia, the Netherlands, Norway, the Philippines, Singapore, and Switzerland.

Influenza A (H1N1) viruses were reported from Africa (Egypt and South Africa) the Americas (Canada and Chile), Asia (China, Hong Kong, Iran, Japan, the Philippines, the Republic of Korea, Singapore, Taiwan, and Thailand), Europe (Bulgaria, Croatia, Finland, France, Ireland, Israel, Italy, Norway, Poland, Portugal, Romania, the Russian Federation, Spain, the Ukraine, and the United Kingdom), and Oceania (Australia, New Caledonia, and New Zealand). Since September 30, influenza A (H1N2) viruses have been identified from Canada, Egypt, Hong Kong, India, Israel, Malaysia, Romania, Singapore and the United Kingdom. Before September 30, H1N2 viruses also were collected from India and Oman. Influenza A (unsubtyped) viruses were reported from Belarus, Belgium, and Brazil.

# Composition of the 2002–03 Influenza Vaccine

The Food and Drug Administration's Vaccines and Related Biological Products Advisory Committee (VRBPAC) recommended that the 2002–03 trivalent influenza vaccine for the United States contain A/New Caledonia/20/99-like (H1N1), A/Moscow/10/99-like (H3N2), and B/Hong Kong/330/01-like viruses. This recommendation was based on antigenic analyses of influenza viruses isolated recently, epidemiologic data, and postvaccination serologic studies in humans.

Most influenza A (H3N2) viruses isolated worldwide during the 2001–02 season were similar to A/Panama/2007/99-like and A/Moscow/10/99-like (H3N2) viruses. Some influenza A (H3N2) viruses were distinguished antigenically from the reference strains but were heterogeneous antigenically

<sup>¶</sup> Levels of activity are 1) no activity; 2) sporadic—sporadically occurring ILI or culture-confirmed influenza with no outbreaks detected; 3) regional—outbreaks of ILI or culture-confirmed influenza in counties with a combined population of <50% of the state's population; and 4) widespread—outbreaks of ILI or culture-confirmed influenza in counties with a combined population of ≥50% of the state's population.</p>

<sup>\*\*</sup> The expected baseline proportion of P&I deaths reported by the 122 Cities Mortality Reporting System is projected by using a robust regression procedure in which a periodic regression model is applied to the observed percentage of deaths from P&I over the previous 5 years. The epidemic threshold is 1.654 standard deviations above the seasonal baseline. Before the 1999–2000 season, a new case definition for a P&I death was introduced. During summer 2000, the baseline and epidemic thresholds were adjusted manually to account for the changes in case definition. For the 2001–02 season, sufficient data have been collected by using the new case definition to allow projection of the baseline using the regression procedure employed before the 2000–01 season.

and genetically. Antibodies produced following vaccination with the 2001–02 vaccine containing the A/Panama/2007/99 (H3N2) virus reacted equally well with recent influenza A (H3N2) viruses and the vaccine strain (2); therefore, VRBPAC recommended that an influenza A/Moscow/10/99-like (H3N2) virus be retained in the 2002–03 vaccine. U.S. vaccine manufacturers will use the antigenically equivalent virus A/Panama/2007/99 because of its growth properties.

The hemagglutinin of most influenza A (H1N1) and A (H1N2) viruses were related antigenically to A/New Caledonia/20/99; therefore, VRBPAC recommended that an A/New Caledonia/20/99 (H1N1) virus be retained in the 2002–03 vaccine. Genetic analyses showed that the neuraminidase of influenza A (H1N2) viruses was related closely to the circulating H3N2 viruses. Current vaccines containing A/New Caledonia/20/99-like virus antigen induced antibodies to H1N2 strains, which were similar in titer and frequency to those of the vaccine strain.

Many influenza B isolates were from the B/Yamagata/16/88 lineage represented in the 2001–02 vaccine by B/Sichuan/379/99. However, B/Hong Kong/330/01-like viruses, which belong to the B/Victoria/2/87 lineage, have spread to countries in Asia, Europe, and North America. Current vaccines containing antigen of B/Sichuan/379/99-like viruses induced antibodies that reacted poorly with viruses related to B/Hong Kong/330/01. Consequently, VRBPAC recommended that the influenza B component be updated for the 2002–03 vaccine to an influenza B/Hong Kong/330/01-like virus. U.S. manufacturers will use one of the antigenically equivalent viruses: B/Hong Kong/330/01 or B/Hong Kong/1434/02.

**Reported by:** WHO Collaborating Center for Reference and Research on Influenza; A Postema, MPH, L Brammer, MPH, H Hall, A Klimov, PhD, K Fukuda, MD, N Cox, PhD, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases; P Terebuh, MD, EIS Officer, CDC.

Editorial Note: Overall, the 2001–02 influenza season was mild to moderate. Influenza A (H3N2) viruses predominated; however, influenza B viruses were isolated more frequently during the late spring in the United States and continue to circulate. Since 1990, viruses of the B/Yamagata lineage have circulated widely. The influenza B component of this season's influenza vaccine belonged to the B/Yamagata lineage. However, during 2001–02, the majority of influenza B viruses characterized in the United States and worldwide was from the B/Victoria/2/87 lineage. Until March 2001, viruses of the B/Victoria lineage had not been identified outside of Asia since 1991. Since March 2001, B/Victoria lineage viruses have been identified in Africa, Asia, Europe, and North America. The 2002–03 influenza vaccine will contain a virus from the B/Victoria lineage.

During the 2001-02 season, influenza A (H1N2) viruses were isolated from several countries, including the United States (3). These new A (H1N2) viruses resulted from gene reassortment among the circulating influenza A (H1N1) and A (H3N2) subtypes. Because hemagglutinin proteins of the A (H1N2) viruses were similar to those of this season's A (H1N1) viruses, and the neuraminidase proteins were similar to this season's A (H3N2) viruses, the 2001-02 vaccine should have provided protection against the A (H1N2) viruses (2). No information suggests that A (H1N2) viruses have been causing more severe illness than other influenza A viruses, and no unusual increases in influenza activity have been associated with these viruses. Influenza A (H1N2) viruses were isolated in China during the 1988-89 influenza season but were not reported in other parts of the world. Whether the new A (H1N2) viruses will persist is uncertain.

Influenza vaccine manufacturers project that approximately 92–97 million doses will be available for distribution during the 2002–03 influenza season. This estimate is based on early projections and could change as the season progresses. In comparison, approximately 70.4 million doses were distributed in 2000 when there were difficulties with growing and processing the influenza A (H3N2) vaccine strain and other manufacturing problems resulted in substantial distribution delays (4). In 2001, a less severe delay occurred. By December 2001, approximately 87.7 million doses of vaccine were produced, more than in any year except the 1976–1977 swine influenza vaccine campaign (5).

The 2002-03 recommendations of the Advisory Committee on Immunization Practices (ACIP) for the Prevention and Control of Influenza (6) contain important changes concerning the timing of vaccination and target groups for vaccination. The optimal time to receive influenza vaccine is during October-November. However, because of vaccine distribution delays during the previous 2 years and uncertainty about vaccine supply in future seasons, ACIP recommends that vaccination efforts during October focus on persons at greatest risk for influenza-related complications (e.g., persons aged ≥65 years and persons aged 6 months-64 years with certain medical conditions), household contacts of these high-risk persons, children aged 6 months to <9 years receiving vaccine for the first time, and health-care workers, and that vaccination of other groups begin in November. Vaccination efforts for all groups should continue into December and later, for as long as vaccine is available. Because young, healthy children are at increased risk for influenza-related hospitalization, vaccination of healthy children aged 6-23 months and close contacts of children aged 0-23 months is encouraged when feasible and should begin during October. Vaccination of children aged  $\geq 6$  months who have certain high-risk medical conditions continues to be recommended strongly (6).

Although influenza epidemics in the temperate regions of the Northern Hemisphere typically peak during December–March, sporadic cases and outbreaks can occur during the summer (7–9). U.S. health-care providers should consider influenza types A and B when diagnosing a febrile respiratory illness during the summer, particularly among persons who have traveled recently in the tropics or Southern Hemisphere, or with large international groups.

# **Acknowledgments**

This report is based on data contributed by participating state and territorial epidemiologists and state health laboratories. WHO collaborating laboratories. National Respiratory and Enteric Virus Surveillance System laboratories. Sentinel Physicians Influenza Surveillance System. Div of Public Health Surveillance and Informatics, Epidemiology Program Office; Div of Vital Statistics, National Center for Health Statistics, CDC. World Health Organization National Influenza Centers, Communicable Diseases, Surveillance and Response, World Health Organization, Geneva, Switzerland. A Hay, PhD, WHO Collaborating Center for Reference and Research on Influenza, National Institute for Medical Research, London, England. I Gust, MD, A Hampson, WHO Collaborating Center

for Reference and Research on Influenza, Parkville, Australia. M Tashiro, MD, WHO Collaborating Center for Reference and Research on Influenza, National Institute of Infectious Diseases, Tokyo, Japan.

#### **References**

- CDC. Influenza activity—United States, 1999–2000 season. MMWR 1999;48:1039–42.
- World Health Organization. Recommended composition of influenza virus vaccines for use in the 2002–2003 season. Wkly Epidemiol Rec 2002;77:62–6.
- 3. World Health Organization. Influenza A(H1N2) viruses. Wkly Epidemiol Rec 2002;77:77–80.
- CDC. Delayed supply of influenza vaccine and adjunct ACIP influenza vaccine recommendations for the 2000–01 influenza season. MMWR 2000;49:619–22.
- CDC. Influenza activity—United States, 2001–02 Season. MMWR 2001;50:1084–6.
- CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 2002;51(No. RR-3).
- CDC. Influenza A—Florida and Tennessee, July-August 1998, and virologic surveillance of influenza, May-August 1998. MMWR 1998;47:756-9.
- 8. CDC. Update: outbreak of influenza A infection—Alaska and the Yukon territory, July–August 1998. MMWR 1998;47:685–8.
- CDC. Outbreak of influenza A infection among travelers—Alaska and the Yukon Territory, May–June 1999. MMWR 1999;48:545–6, 555.

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending June 8, 2002, with historical data

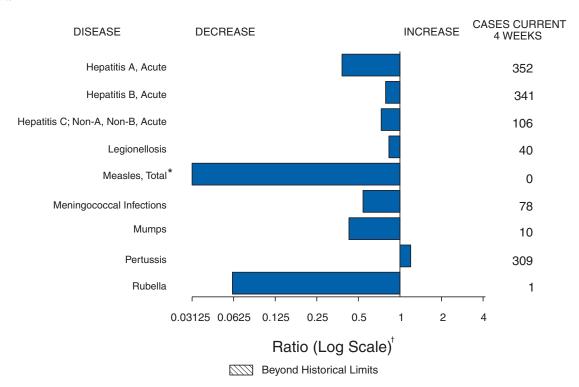


TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending June 8, 2002 (23rd Week)\*

		Cum. 2002	Cum. 2001		Cum. 2002	Cum. 2001
Anthrax		1	-	Encephalitis: West Nile†	1	-
Botulism:	foodborne	7	9	Hansen disease (leprosy)†	34	30
	infant	21	45	Hantavirus pulmonary syndrome†	4	3
	other (wound & unspecified)	8	5	Hemolytic uremic syndrome, postdiarrheal <sup>†</sup>	48	41
Brucellosis†	, , ,	33	46	HIV infection, pediatric <sup>†§</sup>	31	75
Chancroid		28	17	Plague	-	-
Cholera		2	2	Poliomyelitis, paralytic	-	-
Cyclosporiasi	s <sup>†</sup>	52	37	Psittacosis†	11	4
Diphtheria		-	1	Q fever <sup>†</sup>	14	5
Ehrlichiosis:	human granulocytic (HGE)†	56	28	Rabies, human	-	-
	human monocytic (HME)†	22	24	Streptococcal toxic-shock syndrome <sup>†</sup>	37	44
	other and unspecified	2	1	Tetanus	5	18
Encephalitis:	California serogroup viral†	5	1	Toxic-shock syndrome	49	61
·	eastern equine <sup>†</sup>	-	-	Trichinosis	5	5
	Powassan <sup>†</sup>	-	-	Tularemia <sup>†</sup>	15	28
	St. Louis <sup>†</sup>	-	-	Yellow fever	1	-
	western equine†	-	-			

<sup>-:</sup> No reported cases.

<sup>\*</sup> No measles cases were reported for the current 4-week period yielding a ratio for week 23 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.

<sup>\*</sup>Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

Not notifiable in all states.

<sup>\$</sup> Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP). Last update May 26, 2002.

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending June 8, 2002, and June 9, 2001 (23rd Week)\*

(23rd Week)*							Escherichia coli					
		IDS	Chlai	mydia <sup>†</sup>	Cryptos	poridiosis	015	7· <b>U</b> 7		in Positive, non-O157		
Reporting Area	Cum. 2002§	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001		
UNITED STATES	16,795	16,721	307,454	331,455	834	806	630	635	24	32		
NEW ENGLAND	637	575	11,116	9,595	39	35	51	61	5	14		
Maine	19	18	599	578	2	3	2	8	-	-		
N.H. /t.	17 6	14 10	701 300	584 259	10 8	12	4 1	8 2	-	2		
Mass.	318	325	4,636	3,701	9	13	26	29	2	4		
R.I. Conn.	50 227	42 166	1,135 3,745	1,206 3,267	5 5	3 4	4 14	4 10	3	- 8		
MID. ATLANTIC	3,498	4,575	32,097	34,741	92	114	46	50	-	-		
Jpstate N.Y.	259	668	6,969	5,540	27	32	33	31	-	-		
N.Y. City	1,838	2,617	12,826	12,934	43	52	1	4	-	-		
N.J. Pa.	668 733	712 578	2,019 10,283	5,560 10,707	6 16	3 27	9 N	15 N	-	-		
E.N. CENTRAL	1,779	1,155	48,720	61,792	216	276	168	156	1	2		
Ohio	316	190	9,361	16,177	60	49	29	37	1	1		
nd. II.	207 815	117 562	6,698 13,391	6,969 18,378	20 29	28 23	16 53	22 42	-	-		
n. Mich.	358	224	13,869	13,070	46	59	32	23	-	1		
Wis.	83	62	5,401	7,198	61	117	38	32	-	-		
W.N. CENTRAL	270	353	15,077	17,217	99	43	87	79	3	2		
Иinn. owa	56 42	65 40	4,051 629	3,560 2,056	42 9	18	32 19	34 11	3	-		
Лo.	117	161	5,682	5,997	16	12	16	13	-	-		
N. Dak.	2	1 9	469	474 806	5 5	2 4	- 5	1 6	-	- 1		
S. Dak. Nebr.	23	34	978 589	1,561	16	7	9	6	-	1		
Kans.	30	43	2,679	2,763	6	-	6	8	-	-		
S. ATLANTIC	5,478	4,854	61,922	64,375	147	139	65	60	10	9		
Del. Md.	96 822	83 591	1,218 6,384	1,284 6,616	1 5	1 26	1 2	3	-	-		
D.C.	266	357	1,330	1,585	3	9	-	-	-	-		
Va.	350	426	7,157	7,750	2	7	14	15	-	1		
W. Va. N.C.	41 418	33 189	1,035 10,271	1,041 10,353	1 18	14	2 9	1 24	-	-		
S.C.	433	327	5,881	7,139	2	1	-	2	-	-		
Ga. Fla.	922 2,130	575 2,273	12,079 16,567	13,036 15,571	77 38	54 27	29 8	10 5	6 4	6 2		
E.S. CENTRAL	768	813	22,186	21,650	53	16	34	29	7	2		
(y.	122	181	3,526	3,842	1	1	8	29 9	-	-		
Tenn.	341	227	7,264	6,335	27	3	19	12	-	-		
Ala. Miss.	144 161	182 223	6,955 4,441	6,030 5,443	21 4	5 7	3 4	6 2	-	-		
W.S. CENTRAL	1,834	1,587	46,423	47,044	9	22	6	43	_	_		
Ark.	123	89	2,545	3,348	4	2	1	2	-	-		
∟a. Okla.	442 95	392 90	8,396	7,759	2 3	7 3	- 5	2 9	-	-		
Tex.	1,174	1,016	4,398 31,084	4,584 31,353	-	10	-	30	-	-		
MOUNTAIN	565	634	19,311	19,403	56	49	58	66	3	1		
∕lont.	6	12	731	983	4	5	8	5	-	-		
daho Nyo.	10 2	14 1	1,053 387	784 353	16 5	5 1	7 2	8 2	- 1	-		
Colo.	108	139	4,669	5,276	14	15	16	26	i	1		
N. Mex.	34	53	2,600	2,624	6	8	4	5	1	-		
Ariz. Jtah	247 30	243 52	5,969 2,086	6,434 693	6 2	2 10	5 10	10 6	-	-		
Nev.	128	120	1,816	2,256	3	3	6	4	-	-		
PACIFIC	1,966	2,175	50,602	55,638	123	112	115	91	2	4		
Vash. Dreg.	235 181	241 102	6,173 2,842	6,158 3,127	24 17	U 11	15 35	17 18	2	4		
Calif.	1,509	1,800	38,420	43,449	81	99	43	49	-	-		
Alaska	9	10	1,511	1,187	-	-	4	1	-	-		
Hawaii	32	22	1,656	1,717	1	2	18 N	6 N	-	-		
Guam P.R.	2 503	8 533	1,583	182 1,305	-	-	N -	N -	-	-		
V.I.	57	2	30	80	-	-	-	<u>-</u>	<u>-</u>	-		
Amer. Samoa	U	U U	U	U U	U	U U	U	U U	U	U U		
C.N.M.I. N: Not notifiable	11: Unavailable		90		-	Ith of Northern	-		-	U		

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

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

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

§ Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention. Last update April 28, 2002.

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending June 8, 2002, and June 9, 2001 (23rd Week)\*

(23rd Week)*								s influenzae, sive	
	Shiga To	richia coli xin Positive, rogrouped	Giardiasis	Gono	orrhea		Ages, erotypes	Age <5 Serot	ype
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
Reporting Area UNITED STATES	<b>2002</b>	2001 4	<b>2002</b> 5,863	<b>2002</b> 130,634	2001	<b>2002</b> 746	<b>2001</b> 736	<b>2002</b> 10	2001 12
	0	1			148,863		736 34	10	
NEW ENGLAND Maine	-	-	611 65	3,304 41	2,579 58	55 1	1	-	1 -
N.H.	-	-	22	55	59	4	-	-	-
Vt. Mass.	-	1 -	48 288	41 1,473	37 1,120	3 25	1 24	-	1
R.I.	-	-	49	414	305	9	2	-	-
Conn.	-	-	139	1,280	1,000	13	6	-	-
MID. ATLANTIC Upstate N.Y.	-	-	1,302 450	14,735 3,610	15,934 3,368	135 59	101 29	1 1	2
N.Y. City	-	-	535	5,231	5,387	32	30	-	-
N.J. Pa.	-	- -	117 200	1,923 3,971	1,964 5,215	31 13	24 18	- -	2
E.N. CENTRAL	3	2	1,102	23,075	31,511	123	125	2	1
Ohio	3	2	346	5,082	8,589	47	40	-	1
lnd. III.	-	-	- 253	2,890 7,255	2,887 9,875	25 36	20 45	1	-
Mich.	-	-	339	6,133	7,638	9	7	1	-
Wis.	-	-	164	1,715	2,522	6	13	-	-
W.N. CENTRAL Minn.	-	-	711 260	6,183 1,201	7,013 1,117	23 15	28 15	-	1
lowa	-	-	96	170	524	1	-	-	-
Mo. N. Dak.	-	-	202 6	3,466 27	3,531	5	11	-	-
N. Dak. S. Dak.	-	-	28	27 107	16 124	-	-	-	-
Nebr.	-	-	52	137	538	-	1	-	1
Kans.	-	-	67	1,075	1,163	2	1	-	-
S. ATLANTIC Del.	-	-	997 19	36,029 728	38,691 705	191 -	191 -	1 -	1 -
Md.	-	-	40	3,469	3,796	45	47	1	-
D.C. Va.	-	-	19 84	1,124 4,631	1,305 3,851	13	- 15	-	-
W. Va.	-	-	13	409	254	2	4	-	1
N.C. S.C.	-	- -	27	6,964 3,418	7,554 5,381	20 10	28 4	- -	-
Ga.	-	-	399	6,559	7,070	61	52	-	-
Fla.	-	-	396	8,727	8,775	40	41	<del>-</del>	-
E.S. CENTRAL Ky.	-	1 1	130	12,555 1,377	13,861 1,504	24 2	49 2	1	-
Tenn.	-	-	60	4,063	4,167	14	22	-	-
Ala. Miss.	-	-	70	4,394 2,721	4,722 3,468	6 2	23 2	1	-
W.S. CENTRAL			54	20,110	22,640	29	28	2	1
Ark.	-	-	54	1,277	2,092	1	-	-	-
La. Okla.	-	-	-	5,140 1,903	5,315 2,079	2 24	5 22	-	-
Tex.	-	-	-	11,790	13,154	2	1	2	1
MOUNTAIN	3	-	541	4,073	4,563	99	92	2	2
Mont. Idaho	-	-	31 27	39 39	50 35	- 1	- 1	-	-
Wyo.	-	-	9	27	24	i	-	-	-
Colo. N. Mex.	3	-	183 65	1,397 493	1,390 417	19 15	26 13	-	-
Ariz.	-	-	78	1,409	1,780	49	40	1	1
Utah	-	-	92	165	61	10	4	- 1	-
Nev.	-	-	56	504	806	4	8	•	1
PACIFIC Wash.	-	-	415 166	10,570 1,222	12,071 1,312	67 2	88 1	1 1	3 -
Oreg.	-	-	168	354	516	35	29	-	-
Calif. Alaska	-	-	37	8,512 252	9,820 146	9 1	39 3	-	3 -
Hawaii	-	-	44	230	277	20	16	-	-
Guam	-	-	-	-	22	-	-	-	-
P.R. V.I.	-	-	1 -	235 17	300 13	-	1 -	-	-
Amer. Samoa	U	U	U	U	U	U	U	U	U
C.N.M.I.	-	U	-	7	U	-	U	-	U

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

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending June 8, 2002, and June 9, 2001 (23rd Week)\*

Reporting Area 2002 2001 2002 2001 2002 2001 2002 2001	C; Non-A, Non- Cum. 2002 336	-B Cum. 2001
Non-Serotype B   Unknown Serotype   A   B   Cum.   Cum.	<b>Cum.</b> <b>2002</b> 336 17	Cum.
Cum.         Cum. <th< th=""><th><b>Cum.</b> <b>2002</b> 336 17</th><th>Cum.</th></th<>	<b>Cum.</b> <b>2002</b> 336 17	Cum.
Reporting Area 2002 2001 2002 2001 2002 2001 2002 2001	<b>2002</b> 336 17	
	.336 17	2001
UNITED STATES 124 127 9 14 3.000 3.923 2.070 2.963 1	17	1,825
NEW ENGLAND 6 9 159 203 89 59		23
Maine 6 5 3 5		-
N.H 10 4 11 9 Vt 5 2 3	10	5
Mass. 3 7 73 70 46 11	7	18
R.I 20 8 14 9 Conn. 3 2 50 111 13 22	-	-
MID. ATLANTIC 19 16 1 2 468 506 617 579	594	494
Upstate N.Y. 7 4 - 1 86 111 64 55	26	15
N.Y. City 6 4 203 191 357 290 N.J. 4 2 51 121 111 100	- 558	445
Pa. 2 6 1 1 128 83 85 134	10	34
E.N. CENTRAL 17 21 - 1 489 478 365 337	53	98
Ohio 5 5 151 109 45 55 Ind. 6 4 - 1 26 37 13 15	5	5 1
III. 5 8 146 142 31 35	7	8
Mich 114 152 276 212 Wis. 1 4 52 38 - 20	41	84
W.N. CENTRAL 2 1 3 2 161 168 96 96	405	578
Minn. 2 1 1 - 23 14 6 9	-	1
lowa 41 17 10 9 Mo 2 2 37 34 57 57	1 398	573
N. Dak 1 - 1 -	-	-
S. Dak 3 1 - 1 Nebr 5 21 14 11	6	1
Kans 51 81 8 9	-	3
S.ATLANTIC 29 27 - 4 1,119 684 695 524	71	27
Del 8 4 7 10 Md. 1 4 131 97 61 59	3 9	1 3
D.C 44 20 8 4	-	-
Va. 2 4 41 60 97 59 W.Va 10 4 13 14	1	6
N.C. 3 1 - 4 120 55 98 98	13	8
S.C. 4 1 37 26 39 6 Ga. 13 13 276 369 227 171	4 16	3
Fla. 6 4 452 49 145 103	24	6
E.S. CENTRAL 7 10 - 2 72 158 80 187	82	114
Ky 1 25 29 18 24 Tenn. 5 5 67 - 80	2 17	5 29
Ala. 2 4 - 1 22 51 32 44	3	2
Miss 1 25 11 30 39	60	78
W.S. CENTRAL 7 4 50 470 171 392 Ark 22 28 51 47	12 1	400 4
La. 1 11 52 12 59	11	96
Okla. 6 4 16 77 1 47 Tex 1 313 107 239	-	3 297
MOUNTAIN 23 11 4 1 298 343 218 221	36	29
Mont 7 5 3 2	-	-
Idaho 19 28 3 7 Wyo 3 2 9 -	- 5	1 4
Colo. 2 49 33 43 51	17	5
N. Mex. 4 6 - 1 7 13 39 60 Ariz. 12 4 3 - 161 187 81 68	3	10 5
Utah 4 1 26 30 15 11	-	1
Nev. 1 - 1 - 26 45 25 22	11	3
PACIFIC 14 28 1 2 839 913 345 568 Wash. 1 1 81 46 29 45	66 11	62 14
Oreg. 4 5 41 61 69 71	10	10
Alaska 1 1 7 12 3 4	45 -	38
Hawaii 2 1 1 9 3 10	-	-
Guam	-	1
V.I.		-
Amer. Samoa         U         U         U         U         U         U         U         U         U           C.N.M.I.         -         U         -         U         -         U         26         U	U -	U U

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

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending June 8, 2002, and June 9, 2001 (23rd Week)\*

(23rd Week)*			ı							
	Legior	nellosis	Liste	riosis	Lyme	Disease	Mala	aria	Mea: To:	
Panarting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001
Reporting Area UNITED STATES	275	341	164	2001	2,051	2,281	440	523	9 <sup>†</sup>	74 <sup>§</sup>
NEW ENGLAND	14	16	19	20	119	532	27	38	-	5
Maine N.H.	2 2	1 3	2 2	-	25	7	1 5	3 2	-	-
Vt.	1	4	-	-	2	1	1	-	-	1
Mass. R.I.	5 -	3 1	12 1	11 1	70 22	218 35	10 1	17 3	-	3 -
Conn.	4	4	2	8		271	9	13	-	1
MID. ATLANTIC Upstate N.Y.	62 17	75 19	27 13	37 12	1,557 1,077	1,223 324	98 18	137 19	5	9
N.Y. City	13	6	7	8	64	35	60	83	5	1
N.J. Pa.	10 22	5 45	3 4	6 11	112 304	260 604	12 8	18 17	-	1 3
E.N. CENTRAL	73	90	22	31	22	172	53	70	-	10
Ohio Ind.	34 6	40 4	9 3	4 3	19 3	5 2	10 1	9 10	-	3 4
III.	-	10	1	9	-	14	15	27	-	3
Mich. Wis.	25 8	18 18	7 2	13 2	- U	1 150	20 7	16 8	-	-
W.N. CENTRAL	20	18	7	6	39	46	35	16	-	4
Minn. Iowa	2 4	1 5	- 1	-	21 5	25 9	12 2	6 1	-	2
Mo.	9	8	4	3	11	9	9	5	-	2
N. Dak. S. Dak.	1	-	1	-	-	-	1	-	-	-
Nebr.	4	3	-	1	-	1	5	2	-	-
Kans. S. ATLANTIC	- 59	1 45	1 25	2 26	2 245	2 207	6 135	2 101	1	4
Del.	5	-	-	1	30	27	1	1	-	-
Md. D.C.	7 2	10 2	4	2	130 9	128 7	36 5	41 4	- -	3
Va. W. Va.	5 N	7 N	2	5 4	14 3	35 1	10 2	21 1	-	-
N.C.	5	5	3	-	35	6	8	2	-	-
S.C. Ga.	5 8	1 6	3 7	2 6	2 1	2	4 49	4 18	-	1
Fla.	22	14	6	6	21	1	20	9	1	-
E.S. CENTRAL Ky.	8 5	29 7	8 2	8 2	12 5	14 5	6 1	11 2	-	2 2
Tenn.	-	10	3	3	3	5	2	5	-	-
Ala. Miss.	3 -	8 4	3 -	3	4	2 2	2 1	3 1	- -	<del>-</del> -
W.S. CENTRAL	3	14	3	18	2	46	3	37	-	1
Ark. La.	- 1	6	-	1 -	1	2	1 2	3 2	-	-
Okla.	2	2	3	-	-	-	-	1	-	-
Tex. MOUNTAIN	- 19	6 22	- 15	17 18	1 11	44 4	- 15	31 23	-	1
Mont.	1	-	-	-	-	-	-	2	-	-
Idaho Wyo.	3	1 2	1 -	1 1	2	2 1	-	2	- -	1 -
Colo.	4	8 1	2	4 3	3 1	-	7 1	12 1	-	-
N. Mex. Ariz.	1 3	6	2 8	3	1	-	2	2	-	-
Utah Nev.	6 1	2 2	2	1 5	3 1	- 1	2 3	2 2	-	-
PACIFIC	17	32	38	45	44	37	68	90	3	38
Wash. Oreg.	3 N	6 N	3 2	2	3	1	8	2 7	-	15 2
Calif.	14	21	29	38	41	32	51	74	3	15
Alaska Hawaii	-	1 4	4	1	- N	N	1 5	1 6	- -	6
Guam	-	-	-	-	-	-	-	-	-	-
P.R. V.I.	-	2	1 -	-	N -	N -	-	3 -	- -	- -
Amer. Samoa	U	U U	U	U	U	U	U	U	U	U
C.N.M.I.	-	U	-	U	-	U	-	U	-	U

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

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

† Of nine cases reported, three were indigenous and six were imported from another country.

§ Of 74 cases reported, 34 were indigenous and 40 were imported from another country.

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending June 8, 2002, and June 9, 2001 (23rd Week)\*

(23rd Week)*	Meningo						D. Live	A
	Disea Cum.	Cum.	Mun Cum.	Cum.	Cum.	ussis Cum.	Cum.	, Animal Cum.
Reporting Area UNITED STATES	<b>2002</b> 796	<b>2001</b> 1,339	<b>2002</b> 118	<b>2001</b> 100	2002 2,378	<b>2001</b> 2,215	2002 2,144	<b>2001</b> 3,028
NEW ENGLAND	790 56	63	5	-	2,376	212	335	258
Maine	4	1	-	-	3 5	 7	22	31
N.H. Vt.	5 4	6 4	3 -	-	5 44	23	11 57	6 34
Mass. R.I.	28 4	38 2	2	-	213 1	169 1	109 23	86 27
Conn.	11	12	-	-	5	12	113	74
MID. ATLANTIC	74	140	12	9	123	162	392	460
Upstate N.Y. N.Y. City	25 10	42 23	2 1	2 4	85 7	94 24	234 10	293 8
N.J. Pa.	11 28	23 52	1 8	3	3 28	8 36	59 89	68 91
E.N. CENTRAL	28 122	52 186	15	3 14	298	249	89 26	23
Ohio	47	54	3	1	178	132	5	4
Ind. III.	21 20	17 44	1 6	1 10	19 44	19 29	6 6	1 3
Mich.	22	42	5	2	33	22	9	10
Wis.	12	29	-	-	24	47	-	5
W.N. CENTRAL Minn.	78 19	88 13	10 2	5 2	240 70	102 31	166 10	153 17
Iowa Mo.	11 31	20 31	3	-	89 50	10 42	21 18	30 13
N. Dak.	-	3	1	-	-	-	8	20
S. Dak. Nebr.	2 10	4 8	-	- 1	5 4	3 2	20	23 1
Kans.	5	9	4	2	22	14	89	49
S. ATLANTIC	135	198	17	16	181	102	910	1,056
Del. Md.	6 5	- 27	3	4	2 18	- 16	9 137	21 218
D.C. Va.	20	23	3	2	1 83	1 12	226	- 195
W. Va.	-	6	-	-	6	1	77	59
N.C. S.C.	16 14	48 19	1 2	1 1	18 26	36 18	293 33	268 53
Ga.	21	31	4	7	14	10	132	150
Fla. E.S. CENTRAL	53 42	44 82	4 9	1 3	13 61	8 41	3 77	92 131
Ky.	6	13	4	3 1	18	11	15	10
Tenn. Ala.	18 11	31 29	2 2	-	34 9	17 10	46 16	106 15
Miss.	7	9	1	2	-	3	-	-
W.S. CENTRAL	45	219	10	8	491	149	46	657
Ark. La.	20 13	12 55	1	2	238 2	8 4	-	4
Okla. Tex.	11 1	18 134	9	6	27 224	3 134	46	39 614
MOUNTAIN	58	68	7	7	349	850	84	111
Mont.	2	2	, - 1	-	2	6	4	16
Idaho Wyo.	3 -	4	1 -	1	36 5	159 -	12	1 18
Cólo. N. Mex.	18 1	25 8	1	1 2	157 35	160 43	4	4
Ariz.	19	11	-	1	83	455	63	71
Utah Nev.	4 11	7 4	4 1	1 1	22 9	18 9	- 1	- 1
PACIFIC	186	295	33	38	364	348	108	179
Wash.	37	38	-	-	157	45	-	-
Oreg. Calif.	28 115	37 210	N 26	N 21	59 139	18 267	84	143
Alaska Hawaii	1 5	2 8	- 7	1 16	2 7	1 17	24	36
Guam	-	-	-	-	-	-	-	-
P.R.	1	3	-	-	1	-	34	52
V.I. Amer. Samoa	- U	U	U	U	U	Ū	U	U
C.N.M.I.	-	Ü	-	Ü	-	Ü	-	Ü

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

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending June 8, 2002, and June 9, 2001 (23rd Week)\*

(23rd Week)*								
		Mountain	Dut		ibella Cong		Calman	-ll-sis
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Salmon Cum. 2002	Cum. 2001
UNITED STATES	185	103	5	11	2	-	11,202	12,143
NEW ENGLAND	-	-	-	-	-	-	691	884
Maine	-	-	-	-	-	-	63	93
N.H. Vt.	-	-	-	-	-	-	41 28	57 33
Mass.	-	-	-	-	-	-	385	500
R.I. Conn.	-	-	-	-	-	-	30 144	42 159
MID. ATLANTIC	9	5	2	4	_	_	1,429	1,682
Upstate N.Y.	2	-	1	1	-	-	459	376
N.Y. City	1	1	-	2	-	-	520	450
N.J. Pa.	6	2 2	1 -	1 -	-	-	176 274	384 472
E.N. CENTRAL	2	7	-	2	-	-	1,924	1,666
Ohio	2	1	-	-	-	-	548	515
Ind. III.	-	6	-	2	-	-	154 587	146 470
Mich.	-	-	-	-	-	-	341	281
Wis.	-	-	-	-	-	-	294	254
W.N. CENTRAL	21	19	-	3	-	-	881	743
Minn. Iowa	- 1	1	-	1	-	-	199 131	244 114
Mo.	19	17	- -	1	-	-	354	173
N. Dak.	-	-	-	-	-	-	9	11
S. Dak. Nebr.	-	1	-	-	-	-	29 51	45 55
Kans.	1	-	-	1	-	-	108	101
S. ATLANTIC	129	37	1	1	=	-	2,697	2,599
Del.	1	-	-	-	-	-	15	29
Md. D.C.	17	6	1 -	-	-	-	271 31	260 29
Va.	3	1	-	-	-	-	300	412
W. Va.	-	-	-	-	-	-	39	37
N.C. S.C.	65 28	16 7	-	-	-	-	406 179	412 286
Ga.	14	4	-	-	-	-	655	442
Fla.	1	3	-	1	-	-	801	692
E.S. CENTRAL	19	24	-	-	1	-	688	663
Ky. Tenn.	14	1 20	-	-	1	-	111 195	116 175
Ala.	5	1	-	-	-	-	206	205
Miss.	-	2	-	-	-	-	176	167
W.S. CENTRAL Ark.	3	7 4	1	-	-	-	372 181	1,303 155
La.	-	1	-	-	-	-	75	256
Okla.	3	2	<del>.</del>	-	-	-	114	91
Tex.	-	-	1	-	-	-	2	801
MOUNTAIN Mont.	2	4 1	-	-	-	-	822 35	793 30
Idaho	-	i	-	-	-	-	55 55	45
Wyo.	1	1	-	-	-	-	21	26
Colo. N. Mex.	-	-	-	-	-	-	211 105	216 101
Ariz.	-	-	-	-	-	-	248	216
Utah Nev.	- 1	1	-	-	-	-	62 85	83 76
	1	-	-	-	-	-		
PACIFIC Wash.	- -	- -	1 -	1 -	1 -	- -	1,698 170	1,810 178
Oreg.	-	-	-	-	-	-	152	110
Calif.	-	-	1	-	-	-	1,248 26	1,360 20
Alaska Hawaii	-	-	-	1	1	-	102	142
Guam	-	-	-	-	-	-	-	3
P.R.	-	-	-	3	-	-	57	369
V.I. Amer. Samoa	- U	- U	- U	- U	- U	Ū.	- U	Ū
C.N.M.I.	-	Ü	-	U	-	U	17	U
		-		-		-	* *	-

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

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending June 8, 2002, and June 9, 2001 (23rd Week)\*

(23rd Week)*	Shige	ellosis	Streptococo Invasive,			s pneumoniae, tant, Invasive	Streptococcus Invasive (	
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001
UNITED STATES	5,323	5,997	2,066	2,143	1,319	1,644	111	283
NEW ENGLAND	102	104	106	145	6	76	11	63
Maine N.H.	3 4	4 1	14 23	10 9	- -	- -	-	-
Vt.	-	3	8	8	3	6	1	-
Mass. R.I.	72 4	71 7	53 8	44 6	3	-	10	36 1
Conn.	19	18	-	68	-	70	-	26
MID. ATLANTIC	300 66	636 214	363 179	334 147	66 62	91 89	38 38	63 63
Jpstate N.Y. N.Y. City	174	182	90	104	02 U	09 U	-	-
N.J.	21	121	67	54	-	-	-	-
Pa.	39	119	27	29	4	2	-	-
E.N. CENTRAL Ohio	578 308	888 333	306 126	476 122	109 3	114	32	72 -
Ind.	32	110	20	39	102	114	24	35
III. Mich.	141 61	215 136	4 156	160 112	2 2	-	8	25 12
Wis.	36	94	-	43	-	-	-	-
W.N. CENTRAL	499	582	141	205	333	78	23	25
Minn. Iowa	100 38	215 101	69 -	79 -	241	40	23	24
Mo.	58	118	31	47	5	9	-	-
N. Dak. S. Dak.	7 139	12 61	9	7 7	1 1	2	-	1 -
Nebr.	104	35	13	23	23	7	-	-
Kans.	53	40	19	42	62	17	-	-
S. ATLANTIC Del.	2,106 6	850 4	396 1	357 2	680 3	870 2	6	4
Md.	356	48	58	26	-	-	<del>.</del>	-
D.C. Va.	25 382	23 62	5 43	3 51	33	3	1 -	3
W. Va.	2	4	7	13	32	31	-	1
N.C. S.C.	125 33	162 100	77 25	85 6	114	- 187	5	-
Ga.	733	116	119	114	233	257	-	-
Fla.	444	331	61	57	265	390	-	-
E.S. CENTRAL Ky.	489 58	580 203	58 6	43 16	78 8	162 18	-	-
Tenn.	25	41	52	27	70	143	-	-
Ala. Miss.	232 174	115 221	-	-	-	1 -	-	-
W.S. CENTRAL	281	1,164	24	188	22	223	1	56
Ark.	84	273	4	-	5	12	-	-
La. Okla.	53 143	125 16	- 19	26	17 -	181 30	1 -	56 -
Tex.	1	750	1	162	-	-	-	-
MOUNTAIN Mont	240 1	328	377	213	25	29	-	-
Mont. Idaho	2	14	5	3	-	-	-	-
Wyo. Colo.	3 50	2 66	6 134	4 84	9	5	-	-
N. Mex.	48	53	59	46	16	22	-	-
Ariz. Utah	107 15	144 22	173	73 3	-	-	-	-
Nev.	14	27 27	-	-	- -	2	-	-
PACIFIC	728	865	295	182	-	1	-	-
Wash.	50 39	72 46	36	-	-	-	-	-
Oreg. Calif.	616	723	226	159	-	-	-	-
Alaska Hawaii	2 21	3 21	33	23	-	- 1	-	-
Guam	21	21 25	33	23 1	-	'	-	-
P.R.	1	25 8	-	- -	-	-	-	-
V.I. Amer. Samoa	- U	- U	- U	- U	-	-	- U	- U
C.N.M.I.	7	U	-	U	-	-	-	Ü

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

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

TABLE II. (*Continued*) Provisional cases of selected notifiable diseases, United States, weeks ending June 8, 2002, and June 9, 2001 (23rd Week)\*

		Syp	hilis			Typhoid		
	Primary & S			enital†	Tubero	ulosis		ver
Reporting Area	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001	Cum. 2002	Cum. 2001
UNITED STATES	2,533	2,444	49	207	4,530	5,367	99	128
NEW ENGLAND	47	18	-	4	146	198	10	7
Maine N.H.	3	- 1	-	-	5 6	7 8	-	1 1
Vt.	1	2	-	-	-	4	-	-
Mass. R.I.	31 2	9 1	-	2	86 15	104 29	8 -	4
Conn.	10	5	-	2	34	46	2	1
MID. ATLANTIC Upstate N.Y.	287 18	207 5	11 3	33 19	914 133	940 137	27 4	39 9
N.Y. City	172	121	-	-	475	474	13	13
N.J. Pa.	47 50	37 44	5 3	12 2	220 86	214 115	9 1	16 1
E.N. CENTRAL	454	415	-	32	474	543	11	17
Ohio	63	41	-	1	76	105	4	2
Ind. III.	27 114	78 132	-	5 24	49 239	38 273	1 1	1 9
Mich.	242	149	-	2	104	94	3	3
Wis.	8	15	-	-	6	33	2 4	2
W.N. CENTRAL Minn.	38 17	32 17	-	5	215 96	225 96	3	6 2
Iowa Mo.	- 11	1 6	-	3	14 67	18 52	- 1	4
N. Dak.	-	-	-	-	-	3	-	-
S. Dak. Nebr.	- 4	-	-	-	9 9	6 17	-	-
Kans.	6	8	-	2	20	33	-	-
S. ATLANTIC	646	881	7	51	911	1,007	10	18
Del. Md.	8 73	7 114	- 1	1	7 97	82	1	- 5
D.C.	36	14	- -	1	-	34	-	-
Va. W.Va.	32	50 -	-	1 -	72 9	101 13	-	4
N.C. S.C.	131 55	214	1	7 12	123	131 92	-	1
Ga.	100	128 139	-	11	68 130	184	6	6
Fla.	211	215	5	18	405	370	3	2
E.S. CENTRAL Ky.	249 37	258 18	1	9	305 48	341 43	2 2	-
Tenn.	101	144	-	4	110	122	-	-
Ala. Miss.	84 27	45 51	1	2 3	101 46	122 54	-	-
W.S. CENTRAL	357	314	25	38	574	829	-	7
Ark.	12	20	-	2	54	56	-	-
La. Okla.	56 28	60 34	- -	2	- 59	- 59	-	-
Tex.	261	200	25	34	461	714	-	7
MOUNTAIN Mont.	136	89	2	7	128 4	211	8	5 1
Idaho	5	-	-	-	-	3	-	-
Wyo. Colo.	10	14	- 1	-	2 21	1 57	4	-
N. Mex.	21	9	·	-	8	31	-	<del>-</del>
Ariz. Utah	91 6	57 6	1 -	7	79 12	76 8	3	1 -
Nev.	3	3	-	-	2	35	1	3
PACIFIC Wash	319 21	230	3	28	863 95	1,073 95	27	29 1
Wash. Oreg.	5	23 7	-	-	39	45	3 2	3
Calif. Alaska	288	196	3	28	640 25	846 18	22	23
Hawaii	5	4	-	-	64	69	-	2
Guam	<del>.</del>	2	-	, <del>-</del>	-	31	-	1
P.R. V.I.	101 -	118 -	-	11 -	8 -	47	-	-
Amer. Samoa	U 13	U U	U	U	U	U	U	U

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

\* Incidence data for reporting year 2001 and 2002 are provisional and cumulative (year-to-date).

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

TABLE III. Deaths in 122 U.S. cities,\* week ending June 8, 2002 (23rd Week)

TABLE III. Deaths	eaths in 122 U.S. cities,* week ending June 8, 2002 (23rd Week)  All Causes, By Age (Years)  All Causes, By Age								By Age (	Years)		_			
	All	<u> </u>		Jy Ago (I	ou. o,		P&I <sup>†</sup>		All		1		100.07	T	P&I <sup>†</sup>
Reporting Area	Ages	≥65	45-64	25-44	1-24	<1	Total	Reporting Area	Ages	≥65	45-64	25-44	1-24	<1	Total
NEW ENGLAND	439	309	87	26	10	7	37	S. ATLANTIC	1,068	642	234	131	35	26	78
Boston, Mass.	U 39	U 33	U 3	U 1	U 1	U 1	U 6	Atlanta, Ga.	153 235	85 127	38 67	23 28	5 10	2	5 31
Bridgeport, Conn. Cambridge, Mass.	18	13	4	1	-	-	1	Baltimore, Md. Charlotte, N.C.	109	67	13	23	3	3	11
Fall River, Mass.	20	18	2	-	-	-	2	Jacksonville, Fla.	Ü	Ü	Ü	Ü	ŭ	ŭ	Ü
Hartford, Conn.	79	53	16	7	3	-	10	Miami, Fla.	U	U	U	U	U	U	U
Lowell, Mass.	23	15	7	-	1	-	3	Norfolk, Va.	69	46	13	5	1	4	3
Lynn, Mass.	11 16	8 10	3 6	-	-	-	1 3	Richmond, Va.	78 66	36 41	26 12	12 9	3 3	1	10 6
New Bedford, Mass. New Haven, Conn.	48	33	9	1	1	4	4	Savannah, Ga. St. Petersburg, Fla.	64	43	9	8	2	2	5
Providence, R.I.	67	47	13	3	2	2	1	Tampa, Fla.	182	125	29	16	4	8	6
Somerville, Mass.	4	2	1	1	-	-	-	Washington, D.C.	100	66	21	7	4	2	1
Springfield, Mass.	41	23	10	6	2	-	-	Wilmington, Del.	12	6	6	-	-	-	-
Waterbury, Conn.	17	13	4	6	-	-	2	E.S. CENTRAL	615	429	133	27	18	8	50
Worcester, Mass.	56	41	9		-	-	4	Birmingham, Ala.	161	116	33	6	5	1	19
MID. ATLANTIC	2,283	1,533	498	167	49	36	124	Chattanooga, Tenn.	90	76	11	1	2	-	4
Albany, N.Y. Allentown, Pa.	33 12	22 10	8 2	1	-	2	5 1	Knoxville, Tenn. Lexington, Ky.	75 U	50 U	18 U	4 U	2 U	1 U	2 U
Buffalo, N.Y.	98	79	13	4	2	-	9	Memphis, Tenn.	Ü	Ü	Ü	Ü	Ü	Ü	Ü
Camden, N.J.	27	16	2	6	2	1	2	Mobile, Ala.	71	47	17	5	2	-	5
Elizabeth, N.J.	14	11	3	-	-	-	-	Montgomery, Ala.	26	18	7	-	-	1	5
Erie, Pa.	51	39	7	1	2	2	4	Nashville, Tenn.	192	122	47	11	7	5	15
Jersey City, N.J. New York City, N.Y.	39 1,150	28 758	7 255	2 105	20	2 12	48	W.S. CENTRAL	1,571	1,008	352	118	48	44	112
Newark, N.J.	56	24	233	7	1	1	40	Austin, Tex.	82	58	18	2	2	2	3
Paterson, N.J.	18	7	4	4	1	2	1	Baton Rouge, La.	53	41	11	-	-	1	1
Philadelphia, Pa.	428	274	107	28	15	4	21	Corpus Christi, Tex. Dallas, Tex.	57 208	37 117	14 54	2 20	1 10	3 7	4 12
Pittsburgh, Pa.§	35	28	4	1	1	1	3	El Paso, Tex.	74	49	14	9	10	1	6
Reading, Pa.	12 128	5 99	7 22	4	1	2	2	Ft. Worth, Tex.	128	78	36	11	3		10
Rochester, N.Y. Schenectady, N.Y.	23	99 17	22 5	4	1	-	13 6	Houston, Tex.	432	259	96	36	19	22	37
Scranton, Pa.	21	19	2	_		_	-	Little Rock, Ark.	86	54	19	10	1	2	7
Syracuse, N.Y.	47	30	9	1	1	6	2	New Orleans, La. San Antonio, Tex.	50 217	26 164	11 34	6 15	4 4	2	- 17
Trenton, N.J.	35	22	10	1	1	1	1	Shreveport, La.	45	30	10	2	4	3	6
Utica, N.Y.	27 29	23 22	4 4	2	1	-	1 1	Tulsa, Okla.	139	95	35	5	3	1	9
Yonkers, N.Y.								MOUNTAIN	934	644	183	48	24	15	69
E.N. CENTRAL Akron, Ohio	1,590 47	1,120 35	290 7	115 2	24	40 2	99	Albuquerque, N.M.	130	97	21	9	3	-	12
Canton, Ohio	33	22	7	3	1	-	2	Boise, Idaho	44	33	8	2	1	-	3
Chicago, III.	Ü	Ü	ύ	Ü	Ü	U	Ū	Colo. Springs, Colo.	78	49	7	1	-	1	4
Cincinnati, Ohio	U	U	U	U	U	U	U	Denver, Colo. Las Vegas, Nev.	106 214	69 132	28 60	2 11	4 5	3 6	10 9
Cleveland, Ohio	123	88	20	9	1	5	4	Ogden, Utah	27	18	6	1	2	-	4
Columbus, Ohio	243 145	167 111	46 21	19 8	4 3	7 2	20 8	Phoenix, Ariz.	U	U	Ü	U	U	U	U
Dayton, Ohio Detroit, Mich.	179	95	51	22	6	5	12	Pueblo, Colo.	29	23	3	2	-	1	3
Evansville, Ind.	62	48	9	4	-	1	5	Salt Lake City, Utah	120	86	22	8	3	1	11
Fort Wayne, Ind.	79	52	19	5	1	2	6	Tucson, Ariz.	186	137	28	12	6	3	13
Gary, Ind.	15	6	2	6	1	-	1	PACIFIC	1,418	1,013	250	99	34	22	104
Grand Rapids, Mich. Indianapolis, Ind.	35 170	21 133	7 24	3 7	1	3 6	- 12	Berkeley, Calif. Fresno, Calif.	17 169	12 109	2 37	1 13	7	2	7
Lansing, Mich.	32	20	8	4	_	-	4	Glendale, Calif.	31	25	5	-	-	1	-
Milwaukee, Wis.	122	86	24	9	-	3	8	Honolulu, Hawaii	83	60	17	5	1	-	5
Peoria, III.	46	36	4	3	1	2	3	Long Beach, Calif.	71	47	11	10	2	1	6
Rockford, III.	39	32	6	-	-	1	4	Los Angeles, Calif.	U	U	ñ	U	Ú	Ų	U
South Bend, Ind. Toledo, Ohio	37 111	27 81	6 19	4 5	5	1	2 8	Pasadena, Calif. Portland, Oreg.	28 155	21 118	5 26	7	1 1	1 3	3 18
Youngstown, Ohio	72	60	10	2	-	-	-	Sacramento, Calif.	203	143	29	16	10	5	29
W.N. CENTRAL	539	367	103	38	15	16	30	San Diego, Calif.	177	120	35	15	4	3	9
Des Moines, Iowa	49	38	7	3	-	1	1	San Francisco, Calif.	U	U	U	U	U	U	U
Duluth, Minn.	36	32	3	1	-	-	6	San Jose, Calif.	155	114	24	12	4	1	12
Kansas City, Kans.	22	12	6	4	-	-	3	Santa Cruz, Calif. Seattle, Wash.	39 135	31 101	6 22	2 9	2	1	1 8
Kansas City, Mo.	48	32	11	4	-	1	-	Spokane, Wash.	61	46	8	4	2	1	3
Lincoln, Nebr.	U 75	U 45	U 21	U 1	U 3	U 5	U 9	Tacoma, Wash.	94	66	23	5	-	-	3
Minneapolis, Minn. Omaha. Nebr.	75 73	45 56	12	3	1	5 1	9 5	TOTAL	10,457 <sup>¶</sup>	7,065	2,130	769	257	214	703
St. Louis, Mo.	116	69	20	16	6	5	-		. 0, 107	.,500	_,.00	. 50			. 50
St. Paul, Minn.	48	37	7	1	2	1	2								
Wichita, Kans.	72	46	16	5	3	2	4								

U: Unavailable. -: No reported cases.

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

† Pneumonia and influenza.

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

† Total includes unknown ages.

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

Use of trade names and commercial sources is for identification only and does not imply endorsement by the U.S. Department of Health and Human Services.

References to non-CDC sites on the Internet are provided as a service to MMWR readers and do not constitute or imply endorsement of these organizations or their programs by CDC or the U.S. Department of Health and Human Services. CDC is not responsible for the content of these sites. URL addresses listed in MMWR were current as of the date of publication.

The Morbidity and Mortality Weekly Report (MMWR) Series is prepared by the Centers for Disease Control and Prevention (CDC) and is available free of charge in electronic format and on a paid subscription basis for paper copy. To receive an electronic copy each week, send an e-mail message to listserv@listserv.cdc.gov. The body content should read SUBscribe mmwr-toc. Electronic copy also is available from CDC's World-Wide Web server at http://www.cdc.gov/mmwr or from CDC's file transfer protocol server at ftp://ftp.cdc.gov/pub/publications/mmwr. To subscribe for paper copy, contact Superintendent of Documents, U.S. Government Printing Office, Washington, DC 20402; telephone 202-512-1800.

Data in the weekly *MMWR* are provisional, based on weekly reports to CDC by state health departments. The reporting week concludes at close of business on Friday; compiled data on a national basis are officially released to the public on the following Friday. Address inquiries about the *MMWR* Series, including material to be considered for publication, to Editor, *MMWR* Series, Mailstop C-08, CDC, 1600 Clifton Rd., N.E., Atlanta, GA 30333; telephone 888-232-3228.

All material in the MMWR Series is in the public domain and may be used and reprinted without permission; citation as to source, however, is appreciated.

☆U.S. Government Printing Office: 2002-733-100/69035 Region IV