

MORBIDITY AND MORTALITY

WEEKLY REPORT

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Use of Medical Care, Police Assistance, and Restraining Orders by Women Reporting Intimate Partner Violence — Massachusetts, 1996–1997

Approximately 1.5 million women in the United States are physically or sexually assaulted by an intimate partner (IP) each year (1). The Woman Abuse Tracking in Clinics and Hospitals (WATCH) Project at the Massachusetts Department of Public Health analyzed data from the 1996 and 1997 Behavioral Risk Factor Surveillance System (BRFSS) in Massachusetts to 1) estimate the percentage of women aged 18–59 years experiencing intimate partner violence (IPV) who used medical care, police assistance, and restraining orders during the preceding 5 years, 2) determine where women experiencing IPV went for medical care, and 3) examine the overlap in use of these three services. This report describes the results of these analyses, which indicate that a higher percentage of women aged 18–59 years use police assistance rather than obtain a restraining order or seek medical care.

BRFSS is an ongoing, state-based, random-digit-dialed telephone survey of the U.S. civilian, noninstitutionalized population aged \geq 18 years. Questions on IPV developed by the WATCH Project were added to the Massachusetts BRFSS in 1996 and 1997. During the 2 years, 2940 women aged 18–59 years responded to the survey (response rate: 64.5%). Of these, 129 (5.5%) were excluded from analysis because they either refused or responded "don't know/not sure" to the initial questions about whether they had ever been physically or sexually hurt, and if so, if this was by an IP*. Women aged \geq 60 years also were excluded from the analyses because of low levels of reporting recent IPV. Data were aggregated across the 2 years and weighted to reflect the probability of selection and the demographic distribution of the Massachusetts adult population. Estimated proportions and standard errors were calculated using SUDAAN (2).

Survey respondents were asked whether they had ever been physically or sexually hurt[†] by an IP and when this violence last occurred. Respondents who reported IPV during the preceding 5 years also were asked the following questions about service use: 1) "Did you see a doctor or nurse as a result of being hurt by any of these people in the past five years?"; 2) "In the past five years, were the police called about any of these incidents?"; and 3) "In the past five years, have you gotten a restraining order at a court

^{*}Same or opposite sex, current or ex-husband/wife, partner, boyfriend, girlfriend, or date.

[†] Being physically or sexually hurt included being shoved, slapped, hit with an object, or forced into any sexual activity.

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against a current or ex-(husband/wife), partner, boyfriend, girlfriend, or date?"[§] Respondents who reported having seen a doctor or nurse were asked where they sought care most recently, and those who reported police assistance were asked how many times the police had come for incidents of IPV during the preceding 5 years.

Among women aged 18–59 years, 18.0% reported ever having experienced IPV, 6.6% reported IPV during the preceding 5 years, and 2.1% reported IPV during the preceding 12 months (Table 1). Among women reporting IPV during the preceding 5 years, 39.0% received police assistance, 33.8% obtained a restraining order, and 28.7% sought medical care as a result of IPV. Most women who received police assistance also reported obtaining a restraining order: 69.7%¹ of women who received police assistance for IPV also obtained a restraining order against an IP. Among women reporting IPV, 11.1% sought medical care as a result of IPV but did not obtain police assistance or a restraining order. Approximately half (55.9%) of women reporting IPV received one or more of the three services.

Most women reporting IPV during the preceding 5 years were aged 18–29 years (64.0%), employed (69.8%), had some college education (60.3%), and had children in the household (52.5%). Half (50.1%) of women had never been married, 28.6% were divorced or separated, and 21.3% were married or cohabitating.

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Editorial Note: Federal, state, and local efforts are under way to establish surveillance systems for IPV. The WATCH Project, along with projects in Michigan and Rhode Island, have been funded by CDC to establish statewide tracking systems for IPV against women. IPV surveillance systems are frequently based on service provider data; however, these data represent only persons accessing that particular service. Service provider data are unable to provide estimates of the total number of women experiencing IPV in a population or the extent to which the same women may be represented in different service provider data sets. Surveillance data from the WATCH Project provide state-based estimates of the percentage of women experiencing IPV using three key types of services and the degree of overlap in service use.

Other population-based studies report similar findings regarding the frequency at which women experiencing IPV use services. Police assistance for IPV is received by 35%-56% of women reporting IPV (3-5). Of women physically abused by their partners, 22% seek restraining orders against an IP (4). Among women reporting IPV, 10%-21% receive medical care as a result of IPV, and approximately 70% of these women seek care at an emergency department (3,4,6). Finally, 16% of persons who experience family violence or IPV identified through police incident reports have violence-related contact with a regional hospital (7).

[§]Questions on medical care and restraining orders were revised during 1996–1997 for clarification. The question on medical care was reworded from "after being hurt" to "as a result of being hurt" and the question on restraining orders was reworded from "have you been to court to get a restraining order" to "have you gotten a restraining order at a court." Response frequencies for women aged 18–59 years did not vary significantly for each version of the question.

¹Calculated as the percentage of women who used police and restraining order and the percentage who used police, restraining order, and medical care divided by the percentage who used police with or without other services.

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TABLE 1. Number and percentage of women aged 18–59 years who reported intimate partner violence (IPV) and use of medical care, police assistance, or re-

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Category	No.†	(% [§])	(95% CI [¶])
Incidence of IPV			
Ever	578	(18.0)	(16.0–19.9)
During preceding 5 years	227	(6.6)	(5.3– 7.8)
During preceding 12 months	70	(2.1)	(1.3– 2.8)
IPV not reported	2233	(82.0)	(80.0-84.0)
Services used for IPV			
during preceding 5 years**			
Medical care only	16	(11.1)	(3.7–18.4)
Police only	21	(7.4)	(3.1–11.6)
Restraining order only	16	(3.4)	(1.1– 5.7)
Medical care and police	9	(4.2)	(0.0- 8.3)
Medical care and restraining order	6	(2.8)	(0.1- 5.5)
Police and restraining order	49	(16.5)	(9.5–23.4)
All three services	33	(10.7)	(5.1–16.3)
None of three services	75	(44.1)	(34.0–54.1)
Where medical care for IPV was received			
during preceding 5 years ^{tt}			
Hospital emergency department	44	(60.6)	(41.4–79.8)
Private doctor's office	12	(27.9)	(9.3–46.4)
Hospital walk-in clinic	6	(5.1)	(0–10.5)
Other	2	(6.5)	(0–16.0)
Number of times police came for IPV			
during preceding 5 years ^{ss}			
1 time	45	(47.2)	(32.6–61.8)
2–3 times	41	(29.9)	(17.3–42.5)
4–5 times	14	(19.3)	(6.2–32.4)
6–9 times	5	(2.3)	(0- 5.0)
≥10 times	4	(1.2)	(0- 2.6)

* n=2811; missing=129. [†] Unweighted data.

⁵ Percentages calculated based on weighted data and may not total 100% because of rounding.

[¶] Confidence interval.

** n=227; missing=2.

[#] n=64; missing=0.

^{§§} n=113; missing=4.

The findings in this report are subject to at least three limitations. First, BRFSS is a retrospective self-report survey and may be subject to recall bias. Second, women experiencing IPV who were not eligible to be included in the phone survey, declined participation, or did not disclose IPV may have a different pattern of service use than respondents. Persons who were ineligible to participate included those who were homeless, lived in group housing, did not have a phone, or did not speak English, Spanish, or Portuguese. Finally, IPV may not have been reported because of mistrust, fear of reprisals, and feelings of shame and/or denial.

Intimate Partner Violence — Continued

These findings have implications for both IPV surveillance and medical practice. For surveillance, these results suggest that police data may capture a larger portion of women aged 18–59 years experiencing IPV than a medical care-based surveillance system. In Massachusetts, where police are directed to inform women reporting IPV about the availability of restraining orders, police and restraining order data appear to capture a similar demographic group. However, a medical care-based tracking system may capture a sizeable portion of women experiencing IPV who do not receive police or restraining order assistance. Emergency departments appear to provide the most efficient location within the medical system for tracking IPV-related injuries because most women who seek medical care following incidents of IPV are seen in emergency departments. However, a surveillance system designed to include police, restraining order, and medical care data may miss nearly half of women experiencing IPV.

Medical practitioners, particularly those in emergency departments, need to be prepared to identify and provide support, safety planning, and resources to those experiencing IPV (8). Because many women experiencing IPV do not disclose partner violence unless directly asked, some groups believe women patients whose conditions may be injury-related should be screened systematically for IPV (9, 10). Because 38.7% of women who received medical care for IPV had not received police or restraining order assistance, medical practitioners may be a critical source of support and intervention to many women.

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Unexplained Illness and Death Among Injecting-Drug Users — Glasgow, Scotland; Dublin, Ireland; and England, April–June 2000

Since April 19, 2000, 30 injecting-drug users (IDUs) died or were hospitalized with unexplained severe illness in Glasgow, Scotland. Illness was characterized by extensive local inflammation at a subcutaneous or intramuscular injection site often followed by hypotension and circulatory collapse. Since April 24, 2000, 15 IDUs in Dublin, Ireland, and 14 IDUs in England with similar illnesses have been identified. Despite debridement and broad spectrum antibiotics, 30 (51%) of the 59 patients in all three countries have died. This report further describes the clinical syndrome and key epidemiologic features of the illness as characterized by a preliminary investigation by health authorities in Scotland, Ireland, England, and the United States (1).

A case of unexplained illness was defined as soft tissue inflammation (i.e., abscess, cellulitis, fasciitis, or myositis) at an injection site, and either 1) severe systemic toxicity (i.e., sustained systolic blood pressure <90 mm Hg despite fluid resuscitation and total peripheral white blood cell count [WBC] >30,000 cells/mm³), or 2) postmortem evidence of a diffuse toxic or infectious process including pleural effusions and soft tissue edema or necrosis, in an IDU admitted to a hospital or found dead since April 1, 2000. As of June 5, in Glasgow, 16 (53%) of 30 IDUs evaluated had illnesses that met the case definition (Figure 1). In Dublin, eight (53%) of 15 IDUs, and in England, six (42%) of 14 IDUs had illnesses that met the case definition (Figure 1). Demographic information, peripheral WBC, and case-fatality among IDUs were similar in all three countries (Table 1). Most cases had progressive symptoms with a median of 3 days (range: 0–14 days) between illness onset and hospitalization. Among persons who died while hospitalized, the median

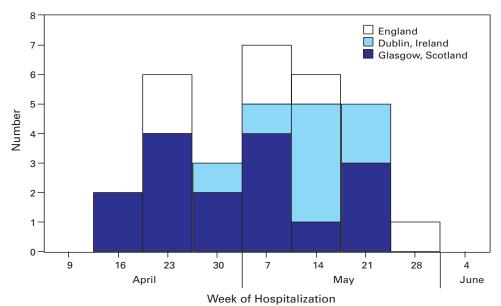


FIGURE 1. Number of cases of unexplained severe illness and death among injectingdrug users — Glasgow, Scotland; Dublin, Ireland; and England, April–June 2000

Characteristic	Glasgow (n=16)	Dublin (n=8)	England (n=6)
Median age, yrs	29	34	34
(Range)	(20-48)	(22–51)	(30-48)
Women	56%	25%	33%
Median WBC, cells/mm ³	76,600	60,000	51,900
(Range)	(39,200–153,000)	(8,200*-96,500)	(39,700-82,000)
Case-fatality	94%	100%	83%

TABLE 1. Demographic characteristics, peripheral white blood cell count (WBC), and percentage case-fatality among injecting-drug users who had illnesses that met the case definition for unexplained severe illness and death — Glasgow, Sectiond: Dublin, Iroland: and England, April, June 2000

* One patient from Dublin with a WBC of 8,200 on admission to a hospital died 6 days later and had an illness that met the case definition based on findings at postmortem examination.

time from admission to death was 2 days (range: 0–13 days). Pleural effusion and extensive edema at an injection site were prominent features at postmortem examination.

Cultures of blood and tissue yielded multiple organisms from several patients including group A streptococcus, *Staphylococcus aureus*, *Clostridium* species, and *Bacillus* species. However, the variable and polymicrobial results and potential postmortem contamination complicate the interpretation of these findings and fail to reveal a definitive etiologic agent. Clinical and drug specimens are being evaluated at CDC, the Public Health Laboratory Service in England, and local laboratories in Glasgow and Dublin. Culture, serologic, molecular, immunopathologic, and histopathologic evaluation of blood and tissue from case-patients have revealed no evidence of *Bacillus anthracis*. *B. anthracis* was isolated from the cerebrospinal fluid of an IDU residing in Oslo, Norway, hospitalized in early April 2000 with a localized abscess, elevated WBC (45,000 cells/ mm³), and hemorrhagic meningitis resulting in death (*2*).

Investigations continue to characterize further the 29 reported unexplained illnesses among IDUs whose illnesses failed to meet the case definition but may be linked to this outbreak. Surveillance activities have been initiated in all hospitals in Scotland, Ireland, England, and Wales to identify additional cases. Information regarding these illnesses is being disseminated to medical practitioners and IDUs, and a case-control study is under way to identify risk factors for disease and to develop prevention strategies. As of June 5, no similar illnesses have been reported in the United States to CDC through the Council of State and Territorial Epidemiologists.

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Editorial Note: The localized inflammatory process affecting skin or muscle combined with systemic toxicity characterized by a leukemoid reaction suggests the role of a toxin-mediated cause of illness among IDUs in Scotland, Ireland, and England. Despite extensive microbiologic evaluation for several of these cases, no specific causative pathogen has been identified. Although the initial symptoms of anthrax can be nondescript before the onset of circulatory collapse and death (3), the absence of *B. anthracis* bacteremia or histologic or molecular evidence for *B. anthracis* suggests that anthraxassociated toxemia is not a cause of illness among these IDUs. Streptococcal toxic shock syndrome and staphylococcal toxic shock syndrome are both characterized by the sudden onset of shock and organ failure, often associated with skin and soft tissue damage (4,5). However, most cases in Scotland, Ireland, and England have not had group A streptococcus isolated (a required feature of streptococcal toxic shock syndrome). and none developed a rash or desquamation of the palms and soles (diagnostic criteria of staphylococcal toxic shock syndrome). Fastidious, anaerobic bacteria, such as *Clostridium* species, have caused a distinctive, toxin-mediated, often fatal infection characterized by sudden onset of shock with unrelenting hypotension, myonecrosis, generalized tissue edema, and a profound leukemoid reaction in the absence of high fever and rash (6)—a clinical course resembling that seen among cases in Scotland, Ireland, and England. Laboratory procedures have been enhanced for the identification of anaerobic bacteria and noninfectious toxins.

The emergence of a new illness among IDUs is possible because the injection of nonsterilized drugs into skin and soft tissue can provide a suitable environment for contaminating pathogens and their toxins or noninfectious toxins alone. Up to 32% of IDUs, particularly those who inject drugs subcutaneously or intramuscularly, have soft tissue abscesses or cellulitis at any given time (7,8). Unusual infections have been linked to subcutaneous or intramuscular drug use, including tetanus and wound botulism among heroin and black tar heroin users, respectively, in California (9, 10), and group A streptococcal infections among cocaine users in Switzerland (11). Microbial or chemical contamination can occur at one of many steps, including production, mixing, dilution, or preparation of the drugs or at the time of injection through contaminated paraphernalia or skin.

Because the source of contamination remains unknown and may be common in these countries, this investigation highlights the importance of enhanced surveillance for syndrome-based illness across national boundaries. Health-care providers and public health personnel are encouraged to report persons with illnesses meeting the case definition to their designated public health authorities.

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Illnesses Associated With Use of Automatic Insecticide Dispenser Units — Selected States and United States, 1986–1999

To control indoor flying insects, restaurants and other businesses commonly use pyrethrin and pyrethroid insecticides sprayed from automatic dispensing units. Usually placed near entrances, these units are designed to kill flying insects in food service or work areas. On May 18, 1999, the Florida Department of Health (FDH) was notified by the Florida Department of Business and Professional Regulation (DBPR) that during May 12– 17, three persons developed pesticide-related illnesses associated with improperly placed automatic insecticide dispensers. After FDH conducted a follow-up investigation and notified CDC's National Institute for Occupational Safety and Health (NIOSH) of this event, surveillance data were reviewed to identify additional cases of pesticide-related illnesses associated with automatic insecticide dispensers. Data were provided by the Toxic Exposure Surveillance System (TESS), the California Department of Pesticide Regulation (CDPR), the Montana Department of Agriculture (MDA), the National Pesticide Telecommunications Network (NPTN), and the Washington State Department of Health (WSDH)*. This report describes cases, summarizes surveillance data for pesticide-

^{*}The data from TESS, NPTN, and MDA were provided by the U.S. Environmental Protection Agency (EPA). EPA and several state health departments collaborate with NIOSH and CDC's National Center for Environmental Health to conduct surveillance of acute pesticide-related illness and injury.

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related illnesses associated with automatic insecticide dispensers, and provides recommendations for safe dispenser use.

Case Reports

Cases 1–3. A 42-year-old cook working at a Florida restaurant developed a sore throat, dyspnea, headache, and dizziness on May 12, 1999, after a several-hour exposure to mist released from insecticide dispensers in the food preparation area. The insecticide dispensers had been installed on May 10, but it is unknown on what day the cook was first exposed. The cook removed the dispensers on May 12 and noted relief of his symptoms. However, the restaurant management reinstalled the dispensers on May 14, and on May 15, a 40-year-old male customer developed headache and shortness of breath within 1 hour of entering the restaurant. These symptoms lasted approximately 4 hours. On May 17, approximately 45 minutes after leaving this restaurant, a 47-year-old male customer experienced a sharp burning sensation in his left eye and noted swelling, redness, and irritation of the eyelid that persisted approximately 24 hours. The implicated pesticide dispenser was within 6 feet of the booth where this customer had been sitting, and it faced his left eye. This person reported his symptoms to DBPR on May 18. None of the three persons sought medical attention for their symptoms. The active ingredients released by these dispensers were pyrethrin and piperonyl butoxide.

Case 4. On August 20, 1995, a 17-year-old male restaurant employee in California was changing the cartridge of an automatic insecticide dispenser. When he closed the dispenser panel, the firing mechanism was activated and discharged a pyrethrincontaining mist into his right eye. The employee immediately experienced burning in the eye and promptly sought medical attention at the emergency department of a local hospital. He was diagnosed with chemical conjunctivitis and treated symptomatically.

Surveillance Data

TESS is maintained by the American Association of Poison Control Centers and collects poisoning reports submitted by approximately 85% of U.S. poison control centers (1). A review of TESS data from 1993 through 1996, the most recent years for which data are available, identified 54 cases of pesticide-related illnesses associated with automatic insecticide dispensers; suicides and intentional misuse/abuse were excluded. Among the 42 cases for which specific age information was available, the median age was 22.5 years (range: 3–73 years). Among the 53 cases for which sex was known, 27 (50%) were male. Twenty (37%) cases were work-related. In all cases, pyrethrin/piperonyl butoxide was the responsible insecticide.

During 1986–1999, 43 cases of acute pesticide-related illnesses associated with automatic insecticide dispensers were reported to CDPR (32 cases), MDA (four cases), FDH (three cases), NPTN (two cases), and WSDH (two cases). Age, sex, and state of occurrence for these cases were compared with those from the TESS database, and no overlap with TESS data was found. Thirty-five (81%) of these cases were in persons exposed while at work, including seven whose exposure occurred during dispenser cartridge replacement or attempts to service faulty dispensers. Seven (16%) cases were in persons exposed while they were customers in restaurants, and one was a movie theater customer. For the 27 with age data available, the median age was 40 years (range: 17–68 years); for the 38 with information on sex, 23 (61%) were women. Resmethrin, a pyrethroid insecticide, was implicated in three cases; the remaining

[†]Comparable information on the circumstances of incidents is not available in the TESS data.

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40 were exposed to pyrethrin/piperonyl butoxide. Most insecticide dispenser-related illnesses identified in the non-TESS data⁺ occurred when the dispensers were improperly placed too close (i.e., <12 feet) to food handling, dining, or work areas; were placed where ventilation currents entrained the mist to such areas; and/or were serviced by persons unfamiliar with proper maintenance of these units.

Among the 94 pyrethrin/piperonyl butoxide-exposed cases in the combined surveillance data, signs and symptoms for 36 (38%) involved the eye; 34 (36%), the neurologic system; 26 (28%), the respiratory system; 23 (24%), the gastrointestinal system; 20 (21%), the nose and throat; 10 (11%), the skin; and eight (9%), the cardiovascular system. Some persons experienced signs and symptoms in more than one system. Among the three resmethrin-exposed cases, reported signs and symptoms included pruritus, throat irritation, nausea, vomiting, diarrhea, headache, burning sensation in the lungs, and cough.

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Editorial Note: This report is the first to document pesticide-related illnesses attributable to automatic insecticide dispensers. Automatic insecticide dispensers are registered by the U.S. Environmental Protection Agency (EPA) for use in the restaurant industry and in other public settings, including schools, hotels, offices, supermarkets, hospitals, day-care centers, and long-term–care facilities (e.g., nursing homes). When used properly, automatic insecticide dispensers reduce the number of flying insects. However, given the dispensers' widespread use and potential for malfunction and/or improper use or maintenance, these units may pose a public health hazard.

Insecticide dispensers of the type described in this report are typically calibrated to spray automatically a fine mist of 50–100 mg of insecticide (consisting of approximately 0.5%–1.85% pyrethrin or resmethrin, along with other active and inert ingredients) every 15 minutes, 24 hours per day. Pyrethrins are insecticides derived from the oleoresin extract of dried chrysanthemum flowers (pyrethrum) (2). Piperonyl butoxide (either alone or combined with n-octyl bicycloheptene dicarboximide) often is added to pyrethrin products to inhibit microsomal enzymes that detoxify pyrethrins (2). Although pyrethrins (classified by EPA as acute toxicity category III compounds[§]) have little systemic toxicity in mammals, they possess irritant and/or sensitizing properties that can induce contact dermatitis, conjunctivitis, and asthma (2,3). Anaphylactic reactions (2) and gastrointestinal symptoms (4) related to inhalation of and cutaneous exposure to pyrethrin also have been reported; however, no previously published reports were identified associating pyrethrin exposure with reported cardiovascular (i.e., tachycardia, chest pain, and palpitations) or neurologic (i.e., headache, dizziness, malaise, altered taste, and lip numbness/burning) signs and symptoms. Resmethrin is a pyrethroid, a class of synthetic insecticides chemically similar to natural pyrethrins (2) and is classified in acute toxicity category III. Pyrethroids are reported to induce abnormal skin sensation, dizziness, salivation, headache, fatigue, vomiting, diarrhea, irritability to sound and touch, and

[§] EPA classifies all pesticides into one of four acute toxicity categories based on established criteria (40 CFR Part 156). Pesticides with the greatest toxicity are in category I, and those with the least are in category IV.

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other central nervous system effects (2,5).

The findings in this report are subject to at least two limitations. First, the surveillance systems that identified cases are passive and may have missed some acute pesticide-related illnesses. Second, lack of detailed information on incidents recorded in the surveillance data may have precluded identification of additional risk factors for insecticide dispenser-related illnesses.

Effective flying insect control can be achieved through nonchemical integrated pest management practices (e.g., proper sanitation practices by employees and installation of air curtains and screens). However, if automatic insecticide dispensers are used, they should be installed according to manufacturer labeling instructions. Warning stickers on dispensers should be considered, installation near supplied-air ducts should be avoided, and timers should be set to dispense insecticide during nonbusiness hours (6). Dispensers used in locations frequented by the public should be installed and serviced by commercial pest control operators. Although they are not required by EPA, persons servicing these devices should use personal protective equipment (i.e., chemical-resistant gloves and goggles designed to provide splash protection).

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Probable Locally Acquired Mosquito-Transmitted *Plasmodium vivax* Infection — Suffolk County, New York, 1999

In the United States, malaria transmission was eliminated in the 1940s, and malaria eradication was certified in 1970 (1). Since then, 60 small localized outbreaks of probable mosquito-transmitted malaria have been reported to CDC (2–6). Before 1995, the number of imported malaria cases reported to the Suffolk County (New York) Department of Health Services ranged from zero to eight per year. Since 1995, seven to 17 cases per year have been reported. In all of these cases, a history of residing in or traveling to an area with endemic malaria outside the United States was confirmed. This report describes the investigation of two cases of *Plasmodium vivax* malaria that occurred in Suffolk County in August 1999; the patients had no history of travel outside of the United States.

Case Reports

Case 1. On August 18, an 11-year-old boy residing in Suffolk County was seen by his physician with a 5-day history of fever, rigors, abdominal pain, arthralgias, and vomiting.

Acquired Mosquito-Transmitted Infection — Continued

Intracellular parasites consistent with *P. vivax* were noted on a complete blood count. The patient was admitted to a local hospital on August 21 with a temperature of 102.0 F (38.9 C), hepatosplenomegaly, and several healing maculopapular bite lesions. Initial laboratory examinations revealed leukopenia (white blood cell count: 2,800/mm³ [normal: 4,500–13,500/mm³]), anemia (hemoglobin: 9.8 g/dL [normal: 11.5–15.5 g/dL]), and severe thrombocytopenia (platelet count: 21,000/mm³ [normal: 150,000–400,000/mm³]). Serology was negative for Lyme disease and babesiosis. Serum electrolytes and chest radiograph were normal. Urinalysis demonstrated a slightly elevated urobilinogen. Examination of peripheral thick and thin blood smears at the New York State Department of Health (NYSDH) and CDC confirmed *P. vivax* infection. The patient was treated with chloroquine phosphate, quinine, clindamycin, and primaquine and was discharged from the hospital on August 25.

The patient's parents reported he had never traveled to a malarious area or had a history of a blood transfusion or organ transplantation. During August 1–7, the patient spent 1 week at a summer camp 20 miles from his hometown. He slept in a tent and went swimming in the camp pond. After his return home on August 7, the patient attended another camp in Massachusetts for 2 days.

Case 2. On August 22, an 11-year-old boy residing in Suffolk County was seen by his physician for a 12-day history of vomiting, diarrhea, fever, chills, and fatigue. On August 27, a complete blood count showed malarial ring forms; the boy was admitted to a hospital the following day. Physical examination on admission revealed a temperature of 100.0 F (37.8 C), no splenomegaly, and multiple healing maculopapular bite lesions. Initial laboratory examinations revealed leukopenia (white blood cell count: 4,300/mm³), severe anemia (hemoglobin: 8 g/dL), and thrombocytopenia (platelet count: 134,000/mm³). Routine blood and urine cultures were negative. Serology was negative for babesiosis. Urinalysis and chest radiograph were normal. Examination of peripheral thick and thin blood smears at NYSDH and CDC revealed intracellular parasites consistent with *P. vivax* (<1% parasitemia). The patient was treated with chloroquine phosphate and primaquine and was discharged from the hospital on August 29.

His parents reported he had never traveled to a malarious area or had a history of a blood transfusion or organ transplantation. The boy spent the same week at the same summer camp as case 1, which is 15 miles from his hometown. During the week he slept in a tent and participated in numerous outdoor activities. On August 10, he began having fevers ranging from 101.0 F to 104.0 F (38.3 C to 40.0 C) with rigors and sweats.

Epidemiologic Investigation

No other unexplained cases of malaria were reported to NYSDH during July 1– August 31, 1999. To identify potential unreported cases, a field investigation was conducted that included 1155 telephone interviews with boys who attended the camp, members of their families and the camp staff, and interviews with residents living within 1 mile of the camp. Sixty-three of 375 boys who attended the camp and members of their families who were interviewed reported having a fever during the defined time period. Fourteen of these persons had unexplained fevers; however, no malaria parasites were shown on peripheral blood smears on any of these persons. Two of the approximately 150 residents who lived within a 1-mile radius of the camp who were interviewed reported a fever during the specified time period. No malaria parasites were shown on their peripheral blood smears. Of 52 farm workers interviewed who had immigrated from Mexico, Guatemala, Honduras, El Salvador, and Bangladesh and who resided in

Acquired Mosquito-Transmitted Infection - Continued

three farms near the camp, three reported a recent history of fever; their blood smears did not reveal parasites.

Entomologic and Environmental Investigation

Routine mosquito trapping by the Suffolk County health department for eastern equine encephalitis during early August (the time these cases would have been transmitted) from sites 7 miles from the summer camp yielded *Anopheles quadrimaculatus* and *An. punctipennis.* Trapping from the campsite in eastern Long Island from August 24 to 31 yielded primarily *An. quadrimaculatus* and a few *An. punctipennis.* No mosquitoes (222 of 248 were tested) from the campsite or the boys' hometowns tested positive for *Plasmodium* species. Mosquito control measures to kill larvae and adults were performed at the camp. The adjacent state park was closed temporarily by the health department until surveillance indicated low numbers of mosquitoes.

Reported by: CB Bradley, MD, MH Zaki, MD, DG Graham, MD, M Mayer, MD, V DiPalma, MSN, SR Campbell, PhD, S Kennedy, Suffolk County Dept of Health Svcs, Hauppauge, New York. MA Persi, DO, Dept of Preventive Medicine, State Univ of New York at Stony Brook, Stony Brook, New York. A Szlakowicz, MA, P Kurpiel, J Keithly, PhD, J Ennis, P Smith, MD, State Epidemiologist, New York State Dept of Health. O Szlakowicz, Mayo School of Medicine, Rochester, Minnesota. Malaria Epidemiology Br (proposed), Entomology Br, Biology and Diagnostics Br, Div of Parasitic Diseases, National Center for Infectious Diseases; and an EIS officer, CDC.

Editorial Note: The two cases presented in this report represent the third episode of possible mosquito-borne malaria in New York during the preceding 7 years (4,5,7) and the 24th episode in the United States since 1985. The possibility of autochthonous (i.e., locally acquired) mosquito-borne malaria transmission in the United States remains a concern because of the frequency of international travel, the presence of gametocytemic persons (i.e., persons with malaria parasites in the blood stream that can infect mosquitoes) in the United States, the presence of competent mosquito vectors, and the occurrence of environmental conditions that favor transmission. This investigation confirmed two epidemiologically linked cases of *P. vivax* infection in children residing and camping in Suffolk County, who probably acquired their infections in eastern Long Island through the bite of one or more locally infected *Anopheles* mosquitoes, a competent vector for malaria.

Neither patient had risk factors for the acquisition of malaria infection, such as travel to a disease-endemic area or history of intravenous drug use. Neither had ever had a blood transfusion or organ transplantation. Other potential sources of infective mosquitoes, such as international airports, were too distant from the presumed site of infection. However, *Anopheles* mosquitoes were identified in the recreational area that both patients had visited during the month of August 1999. In addition, potentially gametocytemic persons were living near this recreational area, and environmental conditions were suitable for the development of the parasite in the mosquito (sporogonic cycle) and larvae into adult mosquitoes. Although case finding and contact tracing activities did not identify persons with malaria who might have been the source of the infection, this does not preclude local transmission, which may have occurred weeks before the investigation.

Suffolk County is one of the most heavily mosquito-infested areas in the northeast. In 1999, the northeastern United States experienced one of the warmest and driest summers in history (8). However, heavy rainfall shortly before the two boys arrived at the

Acquired Mosquito-Transmitted Infection - Continued

camp may have resulted in a large population of adult female mosquitoes. Dry weather followed by heavy rains, in addition to resulting in conditions conducive for mosquito breeding, could have reduced the mosquito predator population.

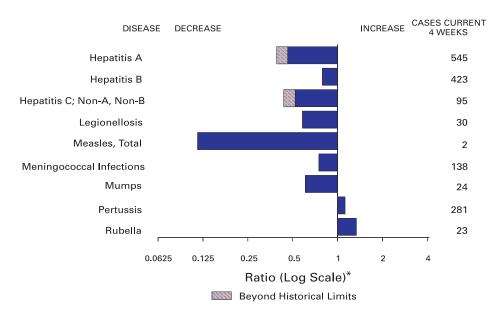
Gametocytemic persons still may be present in the community and constitute a potential reservoir for future episodes of mosquito-borne malaria. Thousands of travelers return to the United States each year from areas where malaria is endemic, and many fail to take adequate chemoprophylaxis. Reintroducing malaria transmission on a small scale in selected areas in the United States is possible. This cluster underscores the need for ongoing surveillance for vector-borne diseases, including malaria. Prompt recognition and adequate treatment of malaria, including improved access to diagnosis and treatment for migrant populations, rapid reporting of malaria cases to public health authorities, and implementation of appropriate control measures, are indicated. Finally, malaria should be considered in the differential diagnosis of illness in any patient with unexplained fevers, regardless of travel history.

During the summer months, persons should follow personal protective measures that reduce contact with potentially infective mosquitoes. These include the use of protective clothing and insect repellants, and sleeping in screened or air-conditioned enclosures. Repellant products containing N,N-diethylmetatoluamide (DEET) are more effective than other compounds.

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FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending June 3, 2000, with historical data



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

		Cum. 2000		Cum. 2000
Anthrax		-	HIV infection, pediatric*§	85
Brucellosis*		16	Plague	3
Cholera		-	Poliomyelitis, paralytic	_
Congenital ru	bella syndrome	4	Psittacosis*	6
Cvclosporiasis	s* ,	7	Rabies, human	-
Diphtheria		1	Rocky Mountain spotted fever (RMSF)	63
Encephalitis:	California serogroup viral*	2	Streptococcal disease, invasive, group A	1,394
•	eastern equine*	-	Streptococcal toxic-shock syndrome*	46
	St. Louis*	-	Syphilis, congenital ¹	45
	western equine*	-	Tetanus	11
Ehrlichiosis	human granulocytic (HGE)*	31	Toxic-shock syndrome	62
	human monocytic (HME)*	7	Trichinosis	4
Hansen diseas	se (leprosy)*	17	Typhoid fever	118
Hantavirus pu	Ilmonary syndrome**	4	Yellow fever	-
Hemolytic ure	mic syndrome, postdiarrheal*	34		

TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending June 3, 2000 (22nd Week)

-: No reported cases.

*Not notifiable in all states.

¹Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases (NCID). ¹Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP). Last update April 30, 2000.

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

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending June 3, 2000, and June 5, 1999 (22nd Week)

		5 chain		, 2000,		10 0, 10	Escherichia coli 0157:H7					
	All	-	Chlan	<u></u>		poridiosis	NET	SS	PH	LIS		
Reporting Area	Cum. 2000 ^s	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999		
UNITED STATES	13,355	18,500	235,842	289,298	486	682	685	557	409	492		
NEW ENGLAND Maine N.H. Vt.	802 14 11 2	940 22 25 6	8,783 563 425 223	8,852 366 437 214	27 6 2 11	36 7 5 6	79 6 6 3	88 5 10 8	63 6 4 2	81 - 10 2		
Mass. R.I. Conn.	535 34 206	614 61 212	4,242 1,009 2,321	3,745 997 3,093	6 2	15 - 3	34 3 27	40 4 21	28 - 23	39 6 24		
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	3,280 186 1,943 703 448	4,449 529 2,109 957 854	14,453 N 2,978 2,511 8,964	32,768 N 15,862 5,272 11,634	45 32 6 1 6	154 45 89 12 8	85 78 4 3 N	36 26 2 8 N	57 38 3 8 8	31 - 3 28 -		
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	1,310 194 100 809 153 54	1,280 211 167 590 248 64	39,037 9,306 4,893 11,273 9,791 3,774	50,430 12,884 4,991 13,421 9,452 9,682	93 21 9 7 16 40	117 16 8 18 17 58	118 24 21 32 24 17	104 40 15 29 20 N	43 13 9 - 14 7	81 24 12 19 16 10		
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr.	299 55 26 139 - 3 20	389 69 46 155 4 11 32	13,817 2,614 1,786 5,022 196 731 1,326	15,950 3,248 1,746 5,832 369 679 1,502	45 10 12 8 3 5 5	38 13 8 4 4 2 6	126 40 19 39 7 2 11	91 23 12 9 3 3 33	79 30 8 21 5 2 9	98 28 6 12 2 6 44		
Kans. S. ATLANTIC Del. Md. D.C. Va. W. Va.	56 3,641 65 392 264 278 21	72 5,168 72 561 207 263 25	2,142 48,871 1,259 4,943 1,407 6,607 753	2,574 59,438 1,201 5,545 N 6,085 769	2 98 2 7 2 4 3	1 130 - 6 5 8	8 - 8 - 13 3	8 68 3 4 - 19 3	4 34 - U 10 2	- 44 - U 17 1		
N.C. S.C. Ga. Fla. E.S. CENTRAL	195 294 357 1,775 639	358 482 827 2,373 840	8,676 3,694 8,695 12,837	9,662 8,102 15,047 13,027 18,472	9 - 53 18 20	3 74 34 8	3 9 3 7 15 33	15 7 5 12 41	2 3 2 7 9 22	12 5 U 9 31		
Ky. Tenn. Ala. Miss.	80 287 169 103	128 337 212 163	20,116 3,370 5,965 6,554 4,227	3,324 6,083 3,893 5,172	1 4 9 6	2 4 1 1	12 14 1 6	11 14 11 5	9 11 - 2	8 12 10 1		
W.S. CENTRAL Ark. La. Okla. Tex.	1,128 69 232 65 762	2,077 70 409 55 1,543	39,147 2,066 7,879 3,434 25,768	37,730 2,430 6,463 3,339 25,498	21 1 5 2 13	49 - 21 1 27	23 4 - 7 12	27 5 4 6 12	44 3 13 3 25	34 4 5 5 20		
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah	477 6 9 2 99 50 165 52	717 4 11 3 143 37 352 70	13,326 601 765 316 2,488 1,688 5,302 1,080	20,136 559 709 330 3,682 2,091 10,567 864	34 4 3 2 9 2 3 9	31 4 2 - 4 12 7 N	64 9 3 21 4 16 1	40 3 1 3 15 2 7 7	25 - 2 7 2 13 1	30 - 3 4 9 1 4 7		
Nev. PACIFIC Wash. Oreg. Calif. Alaska Hawaii	94 1,779 202 47 1,476 5 49	97 2,640 151 63 2,378 6 42	1,086 38,292 5,442 2,230 28,899 1,052 669	1,334 45,522 5,129 2,674 35,576 807 1,336	2 103 N 3 100	2 119 N 11 108	1 99 23 14 55 1 6	2 62 20 14 27 - 1	42 22 14 - 6	2 62 26 12 23 - 1		
Guam P.R. VI. Amer. Samoa C.N.M.I.	13 284 18 -	1 627 13 -	142	196 U U U U	- - - -	- - U U U	N 2 - -	N 10 U U		U U U U		

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

IN: NOV HOLINGUIE. U: Unavailable. -: No reported cases. C.N.M.I: Commonwealth of Northern Mariana Islands.
 Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).
 Chlamydia refers to genital infections caused by *C. trachomatis.* Totals reported to the Division of STD Prevention, NCHSTP.
 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 30, 2000.

Heporting AreaGonorrheaHeporting Cum. 2000Lynn-A, Non-BReporting AreaCum. 2000Cum. 1999Cum. 2000Cum. 1999UNITED STATES123,123149,7291,0611,617NEW ENGLAND Maine2,3462,698239N.H.4034N.H.4034Vt.262533Mass.1,0871,048182R.I.25824423Conn.9011,325MID. ATLANTIC9,40117,4772460Upstate N.Y.2,5072,5302430N.Y. City1,1306,561N.J.1,3473,050Pa.4,3975,336-30E.N. CENTRAL24,10428,926101924Ohio5,3696,9853-Ind.2,2222,6981-Ind.8,0298,941724Mich.6,9366,45290326Wis.1,5483,850-574W.N. CENTRAL5,8866,65227067Minn.1,0651,20142Iowa3754001-Mo.2,9623,22724263N. Dak.637S. Dak	Cum. 2000 270 19 2 2 1 9 2 3 53 21 2 30 68 33 13 6	nellosis 2017 1999 353 22 3 3 3 5 2 6 95 25 12 8 50 108 31 12 15 15 15 15 15 15 15 15 15 15	Dis Cum. 2000 1,532 245 30 1 96 981 417 4 114 446 19 15	yme sease 2,413 555 1 555 1 1 555 1 1 55 6 382 1,334 504 34 263 553 119
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TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending June 3, 2000, and June 5, 1999 (22nd Week)

	weekse	Finding Ju	ane 3, 20	oo, anu J	June 5, 1999 (22nd Week) Salmonellosis*					
	Mal	aria	Rabie	s, Animal	NF	TSS		HLIS		
Deposting Arr	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.		
Reporting Area	2000 376	1999 481	2,100	1999 2,440	2000 10.439	1999 11,478	2000 7.034	1999 10.392		
NEW ENGLAND	17	16	2,100	370	647	663	618	688		
Maine	3	1	61	67	55	45	33	33		
N.H. Vt.	1 2	- 1	4 24	25 56	48 48	35 24	45 49	37 27		
Mass.	6	6	93	80	363	391	340	388		
R.I. Conn.	3 2	- 8	6 81	45 97	25 108	32 136	36 115	53 150		
MID. ATLANTIC	58	138	399	452	1,357	1,568	1,427	1,324		
Upstate N.Y. N.Y. City	19 21	30 62	280 U	307 U	369 313	350 449	378 455	387 469		
N.J.	7	31	67	88	348	367	215	356		
Pa.	11	15	52	57	327	402	379	112		
E.N. CENTRAL Ohio	36 5	57 8	17 4	25 8	1,539 407	1,729 324	906 307	1,520 303		
Ind. III.	3 15	8 28	-	-	178 458	152 561	150	140 556		
Mich.	11	9	13	17	322	369	1 348	354		
Wis.	2	4	-	-	174	323	100	167		
W.N. CENTRAL Minn.	18 7	19 5	205 33	328 38	676 115	689 188	729 200	783 244		
lowa	-	5	31	53	104	70	76	63		
Mo. N. Dak.	1 2	8	5 57	11 71	246 15	217 15	264 25	272 22		
S. Dak. Nebr.	2	-	40	97 2	32 53	31 68	30 44	45 62		
Kans.	6	1	39	56	111	100	90	75		
S. ATLANTIC	103	119	922	877	2,002	2,067	1,159	1,920		
Del. Md.	2 38	1 36	18 166	26 194	32 286	48 277	30 223	53 309		
D.C. Va.	1 25	9 22	235	216	19 267	36 260	U 202	U 319		
W. Va.	-	1	54	49	56	37	42	35		
N.C. S.C.	10 1	10 1	231 51	185 64	281 154	348 111	171 116	379 124		
Ga. Fla.	4 22	12 27	109 58	73 70	346 561	361 589	329 46	504 197		
E.S. CENTRAL	16	10	50 75	115	501	585 614	368	412		
Ky.	3	2	10	20	119	143	76	100		
Tenn. Ala.	5 7	4 3	41 24	40 55	129 157	156 182	165 111	166 125		
Miss.	1	1	-	-	96	133	16	21		
W.S. CENTRAL Ark.	4	11 2	30	52	838 120	1,307 119	775 66	857 76		
La.	2	7	-	-	105	163	118	187		
Okla. Tex.	1	1 1	30	52	106 507	124 901	73 518	90 504		
MOUNTAIN	19	21	89	77	1,015	976	661	928		
Mont. Idaho	1	3 1	24 1	29	42 52	21 36	-	1 37		
Wyo.	-	1	24	27	19	14	14	17		
Colo. N. Mex.	10	8 2	-7	1 2	308 82	313 118	246 59	326 114		
Ariz. Utah	2 3	3	32 1	18	267 146	266 141	217	231 149		
Nev.	3	2	-	-	146 99	67	125	149 53		
PACIFIC	105	90	94	144	1,864	1,865	391	1,960		
Wash. Oreg.	8 21	5 11	-	- 1	171 132	166 151	157 157	273 191		
Calif.	74	69	75 19	137 6	1,469 24	1,408 17	18	1,372 8		
Alaska Hawaii	2	5	-	6 -	24 68	123	18 59	116		
Guam	-	-				20	U	U		
P.R. V.I.	-	Ū	19	36 U	68	203 U	U U	U U		
Amer. Samoa C.N.M.I.	-	Ŭ U	-	Ŭ U	-	Ŭ U	Ŭ	Ŭ U		
N: Not notifiable.	- U: Unav	-	-: No repo	-	-	U	U	U		

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending June 3, 2000, and June 5, 1999 (22nd Week)

N: Not notifiable. U: Unavailable. -: No reported cases. * Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

	WEEKS C	Shige		JU, aliu Ji		99 (22110 V	VEEK/	
	NET			HLIS		philis & Secondary)	Tube	rculosis
Reporting Area	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999 [†]
UNITED STATES	6,310	5,487	2,989	3,019	2,489	2,883	4,018	5,926
NEW ENGLAND Maine N.H. Vt. Mass.	115 5 1 1 78	136 2 7 4 86	94 4 62	121 6 3 76	31	27 - 1 17	144 2 2 95	157 6 1 86
R.I. Conn.	10 20	12 25	8 20	9 27	1 3	1 8	15 30	17 47
MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	835 364 329 75 67	386 87 129 109 61	570 137 296 61 76	203 28 92 79 4	86 7 28 15 36	119 11 47 29 32	895 96 515 210 74	945 119 479 190 157
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	1,135 101 347 296 321 70	894 235 34 342 132 151	404 58 33 2 283 28	453 47 12 286 91 17	482 30 196 117 119 20	480 38 144 180 95 23	474 108 25 255 52 34	599 81 43 320 117 38
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	589 103 159 255 2 2 2 19 49	403 51 6 294 2 8 23 19	403 103 112 151 1 9 26	289 64 9 184 2 5 12 13	33 2 10 16 - - 2 3	61 7 4 42 - 4 4	199 63 30 - 9 6 15	200 78 19 74 2 3 9 15
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	859 5 38 91 3 51 27 107 529	879 7 52 25 32 4 81 39 89 550	206 4 10 53 2 22 34 32 49	228 2 13 U 11 2 51 16 31 102	840 4 126 24 54 1 250 84 142 155	952 4 195 67 2 224 119 179 143	805 99 1 57 15 127 30 178 298	1,174 12 105 20 104 19 158 139 238 379
E.S. CENTRAL Ky. Tenn. Ala. Miss.	313 69 167 16 61	483 61 332 51 39	226 36 176 11 3	333 51 257 24 1	386 42 246 46 52	509 45 271 123 70	288 47 114 127	372 70 109 132 61
W.S. CENTRAL Ark. La. Okla. Tex.	797 83 69 25 620	1,201 42 73 249 837	628 24 53 8 543	387 21 48 74 244	360 44 83 68 165	428 27 113 95 193	126 78 1 47	880 70 U 48 762
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	400 3 29 1 70 41 157 33 66	277 6 4 2 47 37 142 19 20	168 - 2 30 20 81 35 -	176 3 1 36 22 86 22 6	93 - 1 2 11 77 - 2	165 - 1 6 154 2 2	169 6 5 1 15 19 75 20 28	172 5 1 U 21 95 18 32
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	1,267 295 89 860 7 16	828 39 31 737 - 21	290 222 54 3 11	829 50 28 732 19	178 23 3 152 -	142 28 2 110 1 1	918 89 8 736 37 48	1,427 67 40 1,227 29 64
Guam P.R. V.I. Amer. Samoa C.N.M.I. N: Not notifiable.	- 1 - - - U: Una	7 33 U U U	U U U U	U U U U U vrted cases.	- 56 - -	80 U U U		73 U U U

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending June 3, 2000, and June 5, 1999 (22nd Week)

*Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

⁺Cumulative reports of provisional tuberculosis cases for 1999 are unavailable ("U") for some areas using the Tuberculosis Information System (TIMS).

			and	June	5, 1999	(22nd	nd Week)					
	H. influ		He	epatitis (Vi	ral), By Ty	ре			Meas	es (Rubeo	la)	
	Inva		A	_	B		Indiger		Impo		Total	
Reporting Area	Cum. 2000†	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	2000	Cum. 2000	2000	Cum. 2000	Cum. 2000	Cum. 1999
UNITED STATES	525	526	4,685	8,133	2,404	2,804	1	14	-	5	19	52
NEW ENGLAND	36	40	101	90	23	62	-	-	-	-	-	9
Maine N.H.	1 6	4 7	7 11	2 7	5 9	- 6	-	-	-	-	-	- 1
Vt.	2	4	3	1	3	1	-	-	-	-	-	-
Mass. R.I.	20 1	17	46 1	30 9	4 2	27 11	-	-	-	-		6
Conn.	6	8	33	41	-	17	U	-	U	-	-	2
MID. ATLANTIC Upstate N.Y.	77 34	84 32	201 95	520 101	231 54	416 87	-	-	-	-	-	2 2
N.Y. City	18	27	106	136	177	128	-	-	-	-	-	-
N.J. Pa.	19 6	23 2	-	67 216	-	61 140	-	-	-	-	-	-
E.N. CENTRAL	65	84	582	1,431	270	259	-	3	-	-	3	1
Ohio Ind.	28 10	31 12	131 23	331 52	48 20	43 23	-	2	-	-	2	- 1
III.	22	34	206	287	43	-	-	-	-	-	-	-
Mich. Wis.	5	7	209 13	720 41	158 1	173 20	-	1	-	-	1	-
W.N. CENTRAL	30	23	535	322	216	123	1	2	-	-	2	-
Minn. Iowa	15	12	115 44	25 68	15 20	16 21	- 1	- 1	-	-	- 1	-
Mo.	5	2 2	260	191	139	71	-	-	-	-	-	
N. Dak. S. Dak.	1	- 1	-	1 8	2	- 1	1	-	-	-	-	-
Nebr. Kans.	3 6	3 3	17 99	21 8	18 22	11 3	-	- 1	-	-	- 1	-
S. ATLANTIC	145	115	554	730	482	420	-		-	-		4
Del.	-	-	-	2	-	-	-	-	-	-	-	-
Md. D.C.	33	30 3	69 3	142 33	54 5	82 11		-	-	-	-	-
Va. W. Va.	28 5	10 4	65 39	63 14	66 4	41 11	-	-	-	-	-	3
N.C.	13	21	85	51	115	100	-	-		-	-	-
S.C. Ga.	6 40	2 26	16 74	16 224	3 81	37 52	U	-	U	-	-	-
Fla.	20	19	203	185	154	86	-	-	-	-	-	1
E.S. CENTRAL Ky.	26 9	38 5	203 21	196 36	189 37	196 15	-	-	-	-	-	2 2
Tenn.	14	19	80	79	85	85	-	-	-	-	-	-
Ala. Miss.	3	12 2	28 74	33 48	24 43	48 48		-	-	-	-	-
W.S. CENTRAL	29	35	826	2,351	290	466	-	-	-	-	-	3
Ark. La.	- 6	1 9	79 28	21 70	43 50	36 93		-	-	-	-	-
Okla.	21 2	23 2	135	248	56 141	53	-	-	-	-	-	-
Tex. MOUNTAIN		2 51	584 206	2,012		284	-		-	-	- 9	3 1
Mont.	60	1	396 1	627 12	195 3	256 15	-	8 -	-	-	-	-
ldaho Wyo.	2	1 1	14 6	26 4	4	14 5	Ū	-	Ū	-		1
Colo. N. Mex.	11 12	7 11	78 38	116 21	42 44	40 88	-	1	-	1	2	-
Ariz.	30	27	201	372	73	57	-	-	-	-	-	1
Utah Nev.	4 1	2 1	30 28	23 53	12 17	13 24	Ū	3 4	Ū	-	3 4	
PACIFIC	57	56	1,287	1.866	508	606	-	1	-	4	5	30
Wash.	3 17	1 21	128 102	114 130	25 41	25 51	-	-	-	-	-	5 10
Oreg. Calif.	22	28	1,052	1,609	433	516	-	-	-	3	3	15
Alaska Hawaii	2 13	4 2	5	4 9	4 5	9 5	-	1 -	-	- 1	1 1	-
Guam	-	-	-	2	-	2	U	-	U	-	-	1
P.R. V.I.	-	1 U	50	128 U	33	120 U	Ū	-	Ū	-	-	Ū
Amer. Samoa	-	Ŭ	-	Ŭ	-	Ŭ	Ŭ	-	Ŭ	-	-	Ŭ
C.N.M.I.	-	U	-	U · No ror	-	U	U	-	U	-	-	U

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending June 3, 2000, and June 5, 1999 (22nd Week)

N: Not notifiable. U: Unavailable. - : No reported cases. *For imported measles, cases include only those resulting from importation from other countries. *Of 117 cases among children aged <5 years, serotype was reported for 51 and of those, 12 were type b.

			and Ju	ne 5, i	999 (Z	<u>zna w</u>	eek)		i		
	Mening Dise			Mumps			Pertussis			Rubella	
Reporting Area	Cum. 2000	Cum. 1999	2000	Cum. 2000	Cum. 1999	2000	Cum. 2000	Cum. 1999	2000	Cum. 2000	Cum. 1999
UNITED STATES	1,054	1,217	3	168	170	80	1,951	2,550	3	55	89
NEW ENGLAND	60_	63	-	2	3	10	477	259	-	5	7
Maine N.H.	5 4	4 9	-	-	- 1	5	12 59	53	-	- 1	-
Vt. Mass.	2 39	4 38	-	-	2	5	107 274	9 185	-	- 3	-7
R.I. Conn.	3 7	2 6	Ū	1 1	-	Ū	7 18	3 9	Ū	- 1	-
MID. ATLANTIC	101	120	-	9	20	7	155	558	-	2	13
Upstate N.Y. N.Y. City	27 24	32 39	-	6	4 3	7	87	484 10	-	2	9
N.J. Pa.	21 29	21 28	-	- 3	1 12	-	- 68	15 49	-	-	1 3
E.N. CENTRAL	190	206	1	18	23	5	229	201	_	-	-
Ohio Ind.	42 27	76 23	-	7	6	4	160 22	102 10	-	-	-
III.	43	56	1	4	7	-	20	40	-	-	-
Mich. Wis.	60 18	27 24	-	7	7 1	1 -	17 10	18 31	-	-	-
W.N. CENTRAL Minn.	84 7	123 27	-	12	6	11 6	97 53	76 24	-	2	43
lowa	16	23	-	5	1 3	4	15	16	-	-	8
Mo. N. Dak.	48 2	45 3	-	1 -	1 -	-	14 1	17	-	-	-
S. Dak. Nebr.	4 3	5 7	-	2	-	-	1 3	2 1	-	-	- 35
Kans.	4	13	-	4	1	1	10	16	-	2	-
S. ATLANTIC Del.	173	183 3	1	28	30	7	163 4	118	3	32	2
Md. D.C.	16	30 1	-	6	4	1	40	38	-	-	1
Va. W. Va.	29 7	24 4	-	4	8	-	15	13 1	-	-	-
N.C.	28	25 23	1	4	5	5	44	27	3	23	1
S.C. Ga.	12 27	30	U -	8 2	3 1	U 1	16 20	7 15	U -	7	-
Fla.	54	43	-	4	7	-	24 33	17	-	2 4	-
E.S. CENTRAL Ky.	76 16	91 16	-	5	3	2	16	51 12	-	4	2
Tenn. Ala.	35 21	33 25	-	2 2	- 1	2	8 8	25 12	-	- 3	2
Miss.	4	17	-	1	2	-	1	2	-	-	-
W.S. CENTRAL Ark.	82 7	121 22	-	18 1	22	2	66 9	71 5	-	4	4
La. Okla.	25 20	40 19	-	3	3 1	-	3 6	3 8	-	-	-
Tex.	30	40	-	14	18	2	48	55	-	4	4
MOUNTAIN Mont.	59 1	85 2	-	14 1	9	12	362 7	281 2	-	1	15
ldaho Wyo.	6	8 3	Ū	- 1	1	2 U	41	93 2	Ū	-	-
Colo. N. Mex.	18 7	23 10	-	1	3 N	7 2	201 63	78 17	-	1	-
Ariz.	18	28	-	3	-	1	38	57	-	-	13
Utah Nev.	7 2	6 5	Ū	4 3	2 3	Ū	8 4	30 2	Ū	-	1 1
PACIFIC Wash.	229 24	225 34	1	62 3	54 1	24 17	369 121	935 472	-	5	3
Oreg.	31	40	Ň	N	Ň	3	41	19	-	-	-
Calif. Alaska	165 3	142 5	1 -	54 4	47 1	3 1	196 7	423 3	-	5	3
Hawaii	6	4	-	1	5	-	4	18	-	-	-
Guam P.R.	- 4	1 8	U	-	1 -	U -	-	1 7	U	-	-
V.I. Amer. Samoa	-	U U	U U	-	U U	U U	-	U U	U U	-	U U
C.N.M.I. N: Not potifiable	-	available	Ŭ	- No reporte	Ű	Ŭ	-	Ŭ	Ŭ	-	Ŭ

TABLE III. (Cont'd) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending June 3, 2000, and June 5, 1999 (22nd Week)

N: Not notifiable.

U: Unavailable.

- : No reported cases.

		All Cau	ises, Bv	Age (Y		,		All Causes, By Age (Years)							
Reporting Area	All Ages	≥ 6 5	45-64	25-44	1-24	<1	P&l⁺ Total	Reporting Area	All Ages	≥65		25-44	1-24	<1	P&l⁺ Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn Cambridge, Mass Fall River, Mass. Hartford, Conn. Lowell, Mass. New Bedford, Ma New Haven, Conr Providence, R.I. Somerville, Mass Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J.	442 139 50 215 U 222 13 15 U 22 13 15 U 22 23 3 55 2,266 0 0 0 124 26	307 82 44 16 14 26 16 19 38 1,598 48 0 0 33 17 77	6 4 1 U 3 5 3 9 U 8 5 9 425 9 U 8 28 4 2 8 4	29 12 - 1 5 - 1 5 - 1 - 1 2 7 7 156 1 U 1 3	15 10 - - - - - - - - - - - - - - - - - -	6 3 - - U - 1 U - 1 1 48 2 U U 1 -	45 12 8 3 2 U 1 3 4 1 U - 3 - 8 119 7 U 8 2	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla Miami, Fla. Norfolk, Va. Richmond, Va. Richmond, Va. St. Petersburg, f Tampa, Fla. Washington, D.C. Wilmington, D.C. Wilmington, D.C. Wilmington, D.C. Birmingham, Al. Chattanooga, Te Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala.	930 930 279 955 - 102 42 45 45 45 51a. 50 14 - 138 - 138 - 138 - 138 - 138 - 138 - 138 - 162 - 164 - 1	586 U 169 57 64 U 28 24 31 392 61 21 455 795 50 34 104 29	209 U 74 17 23 20 9 8 29 27 - 168 35 20 27 - 168 35 20 15 19 4 4 11	86 U25 8 10 U7 9 3 3 12 9 - 66 16 4 10 7 9 5	25 U 6 5 4 U 1 2 2 4 1 1 3 6 - 1 1	23 U 4 8 1 U 4 - - 4 2 - 5 4 - 5 4	46 U 15 7 5 U 3 3 2 6 2 - 67 17 7 3 8 19 -
Elizabeth, N.J. Erie, Pa.§ Jersey City, N.J. New York City, N. Newark, N.J. Paterson, N.J. Philadelphia, Pa. Pittsburgh, Pa.§ Reading, Pa. Rochester, N.Y. Schenectady, N.Y Scranton, Pa.§ Syracuse, N.Y. Trenton, N.J. Utica, N.Y. Yonkers, N.Y. E.N. CENTRAL	22 45 57 Y. 1,068 42 42 3700 44 21 136 14 28 133 17 17 17 U 1,667	14 32 36 759 21 33 249 33 19 99 92 55 96 12 13 U 1,130	7 12 195 13 4 73 8 1 24 4 1 33 3 3 U 330	5 4 6 81 21 3 1 8 1 2 2 1 U 133	1 20 4 7 - 2 - - - U 39	1 2 12 4 20 - 3 - 1 2 - - - - - - - - - - - - - - - - -	3 40 41 23 6 4 10 2 8 1 U 131	Montgomery, A Nashville, Tenn. W.S. CENTRAL Austin, Tex. Baton Rouge, La Corpus Christi, Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Little Rock, Ark. New Orleans, La San Antonio, Te Shreveport, La. MOUNTAIN	la. 32 118 1,225 64 . 51 Fex. 54 93 325 64 93 325 . U x. 161 102 881	28 76 815 44 39 42 127 65 198 20 114 43 68 592	4 20 234 9 5 7 37 12 13 75 10 U 27 15 24 161	15 105 7 2 20 2 10 39 4 U 14 1 4 80	- 3 35 2 1 - 14 1 2 7 3 U 5 - - 34	- 4 36 2 4 3 7 2 3 6 - U 1 2 6 14	5 8 73 4 3 2 12 - 4 20 3 U 9 8 8 56 56
Akron, Ohio Canton, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Dayton, Ohio Dayton, Ohio Dayton, Ohio Dayton, Mich Fort Wayne, Ind. Gary, Ind. Gary, Ind. Grand Rapids, Mi Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohi W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans Kansas City, Kans	51 323 343 60 1111 101 125 38 52 23 ch. 27 150 34 107 48 47 6 0 48 47 76 0 48 47 76 0 48 47 76 0 48 40 1,400 1,400 4 30 0 2 30 2 41 1,10 1,10 1,10 1,10 1,10 1,10 1,10 1	1,137 377 292 2022 40 75 114 74 74 47 29 39 8 8 19 1066 29 39 8 8 19 1066 29 39 8 8 29 29 20 20 20 20 20 20 20 20 20 20 20 20 20	7 7 2 79 10 0 236 18 25 77 9 4 2 27 8 8 20 0 12 5 7 7 7 9 4 4 2 27 8 8 20 0 15 6 6 6 6 12 6 278 8 10 7 7 64 19 17 7 29 13 13 12 11	3 4 1 42 6 6 12 5 17 2 5 1 - 12 1 2 3 4 1 6 3 85 6 - 12 3 1 5 6 10 6 36 36	32 - 12 - 23 - 6 - 1 1 3 1 1 3 - 2 - 1 1 39 - 1 6 6 6 2 4 1 1 3 1 15	31 - 6425 - 3 2321211 363 - 521542212	7 7 5 34 0 3 9 9 9 13 - 3 1 4 9 9 4 4 4 1 3 2 9 9 1 3 - 3 3 1 4 9 4 4 4 1 3 2 9 1 106 9 2 24 4 2 8 3 - 3 3 51	Albuquerque, N Boise, Idaho Colo. Springs, C Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, U Tucson, Ariz. PACIFIC Berkeley, Calif. Glendale, Calif. Honolulu, Hawa Long Beach, Cal Los Angeles, Cal Pasadena, Calif. Portland, Oreg. Sacramento, Cai San Diego, Calif San Francisco, C San Jose, Calif. Santa Cruz, Cali Seattle, Wash. TotAL	33 30 lo. 45 81 205 366 139 29 tah 103 105 1,328 13 100 9 ii 62 if. 54 if. 303 22 if. 54 if. 303 22 if. 113 . 128 if. 113 . 128 128 if. 114 . 128 . 144 . 14	79 29 52 23 22 24 48 22 66 89 5 89 5 8 49 77 8 49 33 164 144 14 89 57 77 9 75 8 8 9 75 75 8 9 75 8 9 75 75 8 9 75 75 8 9 75 75 29 75 29 75 29 75 29 75 29 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 20 75 75 75 75 75 75 75 75 75 75 75 75 75	$\begin{array}{c} 18\\ 2\\ 8\\ 13\\ 45\\ 2\\ 2\\ 16\\ 6\\ 5\\ 22\\ 2\\ 2\\ 5\\ 22\\ 25\\ 4\\ 20\\ 7\\ 6\\ 2,13$	4 2 6 8 22 6 14 4 11 3 124 4 11 3 124 5 8 8 - 2 7 7 5 8 8 - 7 7 12 U 12 2 6 6 7 7 12 2 8 8 - 8 - 8 8 - 8 8 - 8 8 - 8 8 - 8 -	4 5 5 2 2 9 9 1 3 5 5 4 1 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3	- 23 3 1 4 - 3 1 20 - 1 1 3 6 U U - 1 1 1 3 6 2 31 2 2 31 2 2 31 2 0 - 1 2 2 3 1 2 0 - 1 2 2 3 1 2 3 1 2 2 3 1 2 3 2 3	533812994 11110-51019-101117U82558 754

TABLE IV. Deaths in 122 U.S. cities,* week ending June 3, 2000 (22nd Week)

U: Unavailable. -: No reported cases.

Contributors to the Production of the MMWR (Weekly)

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