



- **1009** Reptile-Associated Salmonellosis
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Reptile-Associated Salmonellosis — Selected States, 1996–1998

During 1996–1998, CDC received reports from approximately 16 state health departments of *Salmonella* infections in persons who had direct or indirect contact with reptiles (i.e., lizards, snakes, or turtles). *Salmonella* infection can result in invasive illness including sepsis and meningitis, particularly in infants. Despite educational efforts, some reptile owners remain unaware that reptiles place them and their children at risk for salmonellosis. This report summarizes clinical and epidemiologic information in four cases and provides information about state regulations to prevent transmission of *Salmonella* spp. from reptiles to humans.

Case Reports

Arizona. During October 1996, a 3-week-old boy was admitted to a hospital emergency department with fever (103.6 F [40 C]), vomiting, and bloody diarrhea of 15 days' duration. Stool and blood cultures yielded *Salmonella* serotype IV 44:z₄,z₂₃-, an extremely rare serotype. The infant was hospitalized for 10 days and treated with intravenous fluids and amoxicillin. To determine the cause of the infant's illness, a stool specimen was obtained from the family's pet iguana, which also yielded *Salmonella* IV 44:z₄,z₂₃-. In an attempt to prevent reinfection, local health officials informed the parents of the importance of the infant avoiding direct and indirect contact with the reptile, and the iguana was moved to a relative's home. One month later, the infant spent 2 days in the relative's home where the iguana was housed; 48 hours after this visit, the infant was again treated at an emergency department for fever and diarrhea. A stool culture again yielded *Salmonella* IV 44:z₄,z₂₃-.

Kansas. During April 1997, a 6-year-old boy had bloody diarrhea of 10 days' duration, abdominal cramps, vomiting, and fever (104.9 F [41 C]). Stool culture yielded Salmonella serotype Typhimurium. The child was treated with ceftriaxone and amoxicillin/clavulanate. Nine days after the boy started therapy, his 3-year-old brother also developed diarrhea, and a stool sample yielded S. Typhimurium. No other family members became ill. The two boys shared a room with two corn snakes that they handled regularly. Stool cultures from the corn snakes yielded S. Typhimurium. The parents reported to health department staff that they were unaware that snakes are a source of salmonellosis.

Massachusetts. During May 1997, an 8-year-old boy with a congenital immune deficiency developed severe vomiting, abdominal cramps, bloody diarrhea, and head-

aches. Stool samples yielded *Salmonella* serotype St. Paul. The boy was ill for 14 days and received extensive supportive care at home. Three days before the boy became ill, the family had purchased two iguanas from a local pet store. The family was not informed by pet store personnel that reptiles are a source of salmonellosis; the child handled the reptiles, including putting them on his head and face. Before diagnostic testing could be performed, the reptiles were returned to the pet store. The parents informed the pet store owner of the child's illness, and the pet store owner reportedly was unaware that reptiles carry *Salmonella* spp.

Wisconsin. In December 1998, a previously healthy 5-month-old boy suddenly died at home. No significant macroscopic or histologic findings were revealed during autopsy; however, culture of a heart blood sample yielded *Salmonella* serotype Marina. The cause of death was attributed to *S.* Marina septicemia. The family had a pet iguana that had not come into direct contact with the infant. Culture of a stool sample taken from the iguana yielded *S.* Marina. Based on an interview, the parents were unaware that the infant was at risk for salmonellosis from indirect or direct contact with the iguana.

State Regulations for Preventing Reptile-Associated Salmonellosis

During March 1999, CDC contacted all 50 state health departments to determine whether state regulations existed for sale of reptiles and distribution of information about salmonellosis. Of the 48 states that responded, three (California, Connecticut, and Michigan) had regulations requiring pet stores to provide information about salmonellosis to persons purchasing a turtle; two states (Kansas and Maryland) require salmonellosis information to be provided to persons purchasing any reptile. Three states (Arizona, Minnesota, and Wyoming) prohibit reptiles in day care centers and long-term–care facilities.

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Editorial Note: In the United States, pet turtles were an important source of salmonellosis until commercial distribution of pet turtles <4 inches long was banned in 1975 (1). This ban led to a 77% reduction in the frequency of turtle-associated *Salmonella* serotypes isolated from humans during 1970–1976 (1). The popularity of other reptiles as pets is growing and has raised concerns about their impact on public health. This and other reports (2) demonstrate that reptile-related salmonellosis continues to pose a substantial threat to human health. Approximately 93,000 (7%) cases per year of *Salmonella* spp. infections are attributable to pet reptile or amphibian contact (3). An estimated 3% of households in the United States have a reptile (CDC, unpublished data, 1999). Many reptiles are colonized with *Salmonella* spp. and intermittently shed the organism in their feces (4). Persons become infected by ingesting *Salmonella* after handling a reptile or objects contaminated by a reptile and then failing to wash their hands properly. Either direct or indirect contact with infected reptiles and their environment can cause human illness (5,6).

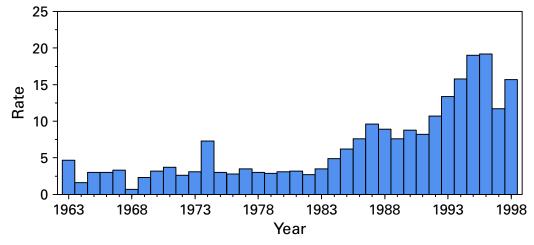
Rare Salmonella serotypes, such as Java, Marina, Stanley, Poona, and Chameleon associated with reptiles, increasingly have been isolated from humans (7) (Figure 1). For example, S. Marina isolates from humans increased from two in 1989 to 47 in 1998, and S. Poona increased from 199 in 1989 to 341 in 1998 (8). Isolation of rare serotypes of Salmonella spp. can alert public health staff about trends in the transmission of infection from reptiles to humans.

Most persons who contract reptile-associated salmonellosis are infants and young children. In 1994, 413 (81%) of 513 *S*. Marina cases occurred in children aged <1 year, whereas 4301 (14%) of 30,723 reported salmonellosis cases occurred in children aged <1 year (6). During 1989–1998, 516 (24%) of 2150 *Salmonella* isolates with reptile-associated serotypes were from children aged <4 years, whereas 50,755 (19%) of 267,131 other serotypes were from this age group (CDC, unpublished data, 1999). Because infants and immunocompromised persons are more susceptible to illness, many reptile-associated *Salmonella* infections involve serious complications, including septicemia and meningitis (9).

The risks for transmission of *Salmonella* spp. from reptiles to humans can be reduced by thoroughly washing hands with soap and water after handling reptiles or objects that have been in contact with reptiles and by preventing reptile contact with food-preparation areas. Children aged <5 years and immunocompromised persons should avoid direct and indirect contact with reptiles. Reptiles also should not be kept in homes of persons with children aged <1 year and in child care facilities (see box). All pet store personnel and reptile owners should be aware that reptiles can carry and transmit *Salmonella* spp. Pet stores are in a unique position to educate consumers because reptile owners obtain most of their information about their pet from pet store personnel. CDC and the Pet Industry Joint Advisory Council (PIJAC) have developed educational posters and brochures for use by veterinarians and pet stores on safe pet reptile handling.*

^{*}Posters are available on request from PIJAC, telephone (800) 553-7387.





^{*}Per 10,000,000 population.

[†]Reptile-associated serotypes are isolates from nonhumans reported to CDC and the U.S. Department of Agriculture that are isolated from reptiles ≥50% of the time.

Recommendations for Preventing Transmission of *Salmonella* from Reptiles to Humans

- Pet store owners, veterinarians, and pediatricians should provide information to owners and potential purchasers of reptiles about the risk for acquiring salmonellosis from reptiles.
- Persons should always wash their hands thoroughly with soap and water after handling reptiles or reptile cages.
- Persons at increased risk for infection or serious complications of salmonellosis (e.g., children aged <5 years and immunocompromised persons) should avoid contact with reptiles.
- Pet reptiles should be kept out of households where children aged <1 year and immunocompromised persons live. Families expecting a new child should remove the pet reptile from the home before the infant arrives.
- Pet reptiles should not be kept in child care centers.
- Pet reptiles should not be allowed to roam freely throughout the home or living area.
- Pet reptiles should be kept out of kitchens and other food-preparation areas to prevent contamination. Kitchen sinks should not be used to bathe reptiles or to wash their dishes, cages, or aquariums. If bathtubs are used for these purposes, they should be cleaned thoroughly and disinfected with bleach.

The effectiveness of educating the public about reptile-associated salmonellosis needs to be evaluated. To enhance efforts to educate the public in a systematic, consistent, and timely manner, the National Association of State Public Health Veterinarians and the Council of State and Territorial Epidemiologists jointly recommend "that the appropriate state and local agencies enact legislation prohibiting the sale or gift of reptiles unless there is written point-of-sale education provided to consumers on the risks for and prevention of reptile-associated salmonellosis" (10). CDC will provide assistance to states interested in developing point-of-sale educational material; however, if these educational efforts should prove unsuccessful, states may wish to adopt restrictions for the sale of reptiles similar to those for turtles.

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Assessment of Laboratory Tests for Plasma Homocysteine — Selected Laboratories, July–September 1998

Cardiovascular disease, including coronary heart disease and stroke, is the leading cause of death in the United States. Elevated plasma homocysteine (Hcy), generally defined as fasting plasma Hcy levels >15 μ mol/L, is an independent risk factor for vascular diseases (1,2). It is unknown whether Hcy is a cause of or a marker for atherosclerosis. A recent statement by the Nutrition Committee of the American Heart Association concluded that until results of clinical trials are available, population-wide Hcy screening is not recommended (3). However, Hcy tests are used in the clinical setting and information on interlaboratory variation, on method variation, is limited. To assess the status of interlaboratory and intralaboratory variation for Hcy analysis, CDC conducted a study of selected laboratories during July–September 1998. This report summarizes findings from the study, which indicates a need to improve analytic precision and to decrease analytic differences among laboratories (4).

Fourteen laboratories participated in the study, including three manufacturers, two government, eight academic, and one clinical research laboratory. Each of three laboratories used two different methods. Selection of laboratories was based on the type of method used for Hcy testing: high-performance liquid chromatography (HPLC), gas chromatography-mass spectrometry (GC-MS), and immunoassay. Laboratories that used HPLC were further subdivided based on the type of detection each laboratory used (electrochemical or fluorescence) and by each of the types of reducing and labeling reagent each used to convert protein-bound and oxidized Hcy into free Hcy and to attach a fluorescent tag to the free Hcy for detection purposes. Laboratories using immunoassay were subdivided into two groups: those using fluorescence polarization immunoassay (FPIA) and those using enzyme immunoassay (EIA). Laboratories participated in a 2-day analysis of 46 blinded plasma samples, four blinded plasma samples with added Hcy, and three plasma quality-control (QC) pools. Interlaboratory and intralaboratory (i.e., between tests run in a laboratory) variation was expressed as a relative standard deviation*. In the absence of target values for the samples analyzed, the GC-MS method was considered arbitrarily as a reference method. Because it used stable-isotopically labeled Hcy as an internal standard, this method is considered to be the most accurate and precise assay available.

For all tests, the mean interlaboratory variation was 9.2% for plasma samples, 8.8% for plasma samples with added Hcy, and 7.6% for the QC pools (Table 1). The mean interlaboratory variation in each method group ranged from 3% to 13%. The group of laboratories performing the FPIA assay had the lowest interlaboratory variation (4.9% for plasma, 3.2% for plasma with added Hcy, and 3.2% for the QC pools). The mean intralaboratory variation was 5.6% for plasma samples, 4.9% for plasma samples with added Hcy, and 4.2% for the QC pools (Table 1). For most laboratories, the intralaboratory variation was <10% and the analytical recovery of added Hcy was 85%–115%.

^{*}Relative standard deviation=standard deviation/mean x 100.

Plasma Homocysteine — Continued

TABLE 1. Mean interlaboratory and intralaboratory variations of homocysteine (Hcy) plasma samples from 14 laboratories, July–September 1998

Plasma	Plasma and Hcy	Quality-control pools
RSD*	RSD	RSD
9.2%	8.8%	7.6% 4.2%
	RSD* 9.2%	RSD* RSD

^{*}Relative standard deviation.

Two of the HPLC methods (HPLC with electrochemical detection and HPLC with fluorescence detection using sodium borohydride as a reducing agent and monobromobimane as a labeling agent) and the EIA method produced results that were, on average, 7.5%, 8.1%, and 7.4% higher than GC-MS results. One HPLC method (HPLC with fluorescent detection using trialkylphosphine as a reducing agent and ABD-F as a labeling agent) produced results that were on average 16.1% lower than GC-MS results. The FPIA method and the two remaining HPLC methods (HPLC with fluorescence detection using either TCEP or TBP as a reducing agent and SBD-F as a labeling agent) showed no deviation compared with results of the GC-MS method.

Analytical quality specifications analysis was performed to test whether the precision[†] and bias[§] of each method were satisfactory (5,6). On the basis of intralaboratory variations, none of the laboratories showed optimum performance for all three types of samples, two laboratories showed desirable performance, and six laboratories exceeded the requirements for minimum performance for at least two types of samples. Three methods performed best regarding analytical precision: GC-MS, FPIA, and HPLC with fluorometric detection using a water-soluble phosphine as reducing agent (TCEP) and SBD-F as fluorescent tag.

With regard to apparent analytical bias, nine laboratories met the requirements for optimum performance with respect to GC-MS, and two laboratories did not meet the requirements for minimum performance. The following three methods performed best regarding apparent analytical bias (versus GC-MS): FPIA, HPLC with fluorometric detection using water-soluble phosphine as reducing agent (TCEP) and SBD-F as fluorescent tag, and HPLC with fluorometric detection using the classical tri-butyl phosphine as reducing agent (TBP) and SBD-F as fluorescent tag.

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Editorial Note: The findings in this report indicate that the fully automated FPIA assay performed best with respect to lowest interlaboratory variation, analytical precision, and apparent analytical bias relative to a GC-MS method. Previous findings have shown a good agreement between the FPIA assay and an HPLC assay with internal standardization for approximately 800 serum and plasma samples (7).

Although both the mean interlaboratory variation and the mean intralaboratory variation were <10% in this study overall, this variation might be clinically unacceptable because of a graded increase in risk for vascular diseases with increasing plasma

[†]Analytical precision is <0.25 x within-patient variability for optimum performance, <0.5 x within-patient variability for desirable performance, and <0.75 x within-patient variability for minimum performance.

[§]The bias of a method is <0.125 x combination of within-patient and between-patient variability for optimum performance, <0.25 x combination of within-patient and between-patient variability for desirable performance, and <0.375 x combination of within-patient and between-patient variability for minimum performance.

Plasma Homocysteine — Continued

Hcy, starting at plasma Hcy concentrations well within the normal range of the population. The findings in this report also indicate that laboratories performing the same method sometimes vary more among themselves than laboratories performing different methods. The analysis also suggests that improvements are needed in the analytical precision to assure that laboratories in an area can use the same reference intervals. To aid this improvement, researchers need to evaluate individual laboratory performance through a program that includes standard reference materials and comparisons with other laboratories. Such an improvement is needed because Hcy has developed from an esoteric test to a clinical test.

The major limitations of this study were the small number of laboratories using each method and the arbitrary selection of the GC-MS method as a reference method, which may itself be biased. As a result, conclusions regarding interlaboratory variation within a method group and with respect to method-specific bias should be interpreted with caution. A high-quality reference method for Hcy is needed to better evaluate a laboratory's or a method's quality.

CDC is promoting efforts to develop a high-order reference method for plasma Hcy by tandem mass spectrometry, the state-of-the-art methodology for accuracy and precision. This method will be a standard for the development and characterization of reference materials. Future studies will assess the variability between laboratories before and after the introduction of standard reference materials.

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Surveillance for Acute Pesticide-Related Illness During the Medfly Eradication Program — Florida, 1998

The Mediterranean fruit fly (Medfly) (*Ceratitis capitata*, Wiedemann) is an exotic insect that can damage approximately 250 fruit and vegetable plant species and is a serious threat to domestic agriculture. During the spring and summer of 1998, pesticides were used by federal and state agriculture authorities to eradicate Medfly infestations that had been detected in portions of five Florida counties (Table 1). This report summarizes surveillance data, describes probable and possible cases of illness asso-

TABLE 1. Number of gallons of malathion applied, number of square miles covered, estimated number of persons exposed, and number of persons adversely affected in the Medfly Eradication Program, by county — Florida, 1998

County	Dates of eradication effort	Total malathion volume, excluding bait (gallons)*	Total treatment area (square miles)	Estimated exposed population	Probable and possible cases of illness	
Dade [†]	April 4–24	1.4	0.08	1,500	0	
Lake/Marion§	April 30-July 13	2,125.0	35.20	2,500	11	
Manatee	May 14-June 26	1,665.0	50.00	120,000	103	
Highlands	July 9-Sept 6	2,494.0	42.50	8,000	9	
Total		6,285.4	127.78	132,000	123	

^{*}In addition, small amounts of diazinon, another organophosphate insecticide, were applied from the ground as a soil drench in Dade (0.04 gallon applied over 0.0001 square mile), Lake (0.3 gallon applied over 0.0007 square mile), and Manatee counties (0.1 gallon applied over 0.0003 square mile).

ciated with the eradication effort, and provides recommendations for future Medfly-eradication programs.

The Medfly Eradication Program began on April 4, 1998, with ground applications of malathion/bait* and diazinon, followed by aerial malathion/bait application that began on April 30. All insecticide applications were completed on September 6. The respective county health departments estimated that 132,000 persons resided in the areas treated with these pesticides.

Surveillance for Illness

Reports of potential adverse health effects attributed to the Medfly Eradication Program pesticide applications were solicited by state health and agriculture authorities and collected through telephone hotlines maintained by the Florida Poison Information Network and county health departments. The public was advised of the pesticide use and the hotline number through public meetings hosted by federal and state agriculture department officials, news articles, and radio and television reports. During April 30–September 30, 1998, 230 reports of illness were received from Florida residents and physicians and were investigated by the Florida Department of Health. Reports were classified according to a standard case classification system. Of the 230 reports, 34 (15%) cases were classified as probable pesticide-related illness based

[†]No aerial application of malathion/bait; ground application only.

[§] Aerial applications of malathion/bait in Lake County also included small portions of Marion County.

^{*}Malathion (Fyfanon® ULV, Cheminova Inc., Wayne, New Jersey) combined with a corn protein bait, Nu-Lure® (Miller Chemical and Fertilizer Co., Hanover, Pennsylvania) was applied at a rate of 2.4 fluid ounces malathion and 9.6 fluid ounces bait per acre per week. The reportedly nontoxic bait comprises hydrolyzed corn gluten meal and inert ingredients including corn syrup. Backpack sprayers or truck-mounted pressure sprayers were used for ground applications; UH-1 "Huey" helicopters and DC-3 aircraft conducted the aerial applications. (Use of trade names and commercial sources is for identification only and does not imply endorsement by CDC or the U.S. Department of Health and Human Services.)

[†]CDC's National Institute for Occupational Safety and Health classifies a case of acute pesticiderelated illness and injury as being definite, probable, possible, or suspicious as determined by the level of certainty of exposure, whether health effects were observed by a health-care provider, and whether sufficient toxicologic information supports a causal relation between the exposure and the reported health effects. When toxicologic evidence for an exposure-health effect relation is not present, the case is classified as unlikely.

on abnormal medical signs compatible with malathion/bait or diazinon toxicity observed by a licensed health-care professional, and 89 (39%) were classified as possible based on symptoms compatible with malathion/bait or diazinon toxicity reported to health-care providers or a state health authority. Of the remaining 107 (47%), 24 were excluded because of insufficient information, 32 were asymptomatic or had symptoms unrelated to exposure, and 51 were classified as unlikely. No reports were classified as definite cases of pesticide-related illness because this category requires confirmation by laboratory testing of clinical or environmental samples that were not available.

The 123 probable or possible cases represent a crude rate of nine cases per 10,000 residents in the exposed areas. Of the 123, 89 (72%) occurred in females; the median age was 46.5 years (range: 6 months–82 years). Eight reports (7%) involved children aged ≤5 years, and 20 (16%) involved persons aged ≥65 years. Four reports (3%) described persons whose illnesses were considered work-related (i.e., Medfly Eradication Program pesticide applicator, lawn-care worker, health department hotline worker, and hotel worker).

Among the 123 cases, signs and symptoms for 87 (71%) were respiratory (e.g., dyspnea, wheezing, coughing, and upper respiratory tract pain/irritation); 77 (63%) involved the gastrointestinal system (e.g., nausea, vomiting, diarrhea, melena, and abdominal cramping); 74 (60%) involved the neurologic system (e.g., headache, vertigo, ataxia, peripheral paresthesia, disorientation, and confusion); 28 (23%) involved the skin (e.g., erythema [with or without maculopapular rash], pruritis, and burning sensations); and 23 (19%) involved the eyes (e.g., lacrimation, conjunctivitis, blepharitis, and blurred vision)§.

Case Reports

- **Case 1.** A 49-year-old man experienced dyspnea, upper respiratory irritation, and headache after being exposed to aerial malathion/bait applications while working on the roof of his house. His physician diagnosed severe bronchitis and reported that the illness probably resulted from malathion/bait exposure.
- **Case 2.** A 31-year-old man reported a blistering rash over his arms, legs, and neck following an aerial application of malathion/bait. He was exposed to malathion/bait while conducting his lawn maintenance business. He reported that the rash developed where grass trimmings coated with pesticide stuck to his skin. His physician diagnosed allergic contact dermatitis secondary to malathion/bait exposure.
- **Case 3.** A 35-year-old man reported a pruritic rash on exposed skin surfaces. He had covered his pool in accordance with recommendations and was exposed to malathion/bait while removing the cover, which he had folded and carried under his right arm. He was not wearing a shirt, and the rash developed at those points where the pool cover had contacted his arm and torso. His physician diagnosed allergic dermatitis.
- **Case 4.** A 32-year-old woman with a history of asthma complained of multiple symptoms in reaction to ground applications of malathion/bait and diazinon in her neighborhood. Symptoms included nausea, diarrhea, abdominal cramping, cough, upper respiratory irritation, dyspnea, wheezing, headache, and fatigue. Her physician diagnosed acute aggravation of asthma secondary to pesticide exposure from the Medfly Eradication Program.

[§]Total is 289 cases because some persons experienced signs and symptoms in more than one system.

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Editorial Note: The Environmental Protection Agency (EPA) classifies malathion as an acute toxicity category III compound \P , and it is considered safer than many other organophosphates because it is rapidly detoxified by the body. Nevertheless, adverse health effects have been reported by persons exposed to malathion (1). Self-reported health effects previously associated with aerial spraying of malathion/bait include respiratory symptoms (particularly among persons with pre-existing respiratory conditions), gastrointestinal symptoms, neurologic symptoms, contact dermatitis, and conjunctivitis (2–4). These effects may represent irritant or allergic responses to either component of the malathion/bait formulation (5,6). Cholinesterase inhibition (3) or anxiety about aerial malathion/bait application (2,7) also may be responsible for some symptoms.

The findings in this report suggest that for most persons, aerial application of malathion/bait does not pose an acute risk to health; however, at least 123 probable or possible pesticide-related cases of illness were associated with pesticide exposure. Each case-patient had signs and/or symptoms consistent with pesticide exposure, and illness probably resulted from sensitivity to the irritant/allergic effects of malathion/bait. Although ground application of diazinon, another acute toxicity category III organophosphate, was employed in some locations, this agent was considered less likely to be responsible for the observed health effects because it was used in only three counties, was applied focally (without aerial application), and was used in minimal quantities.

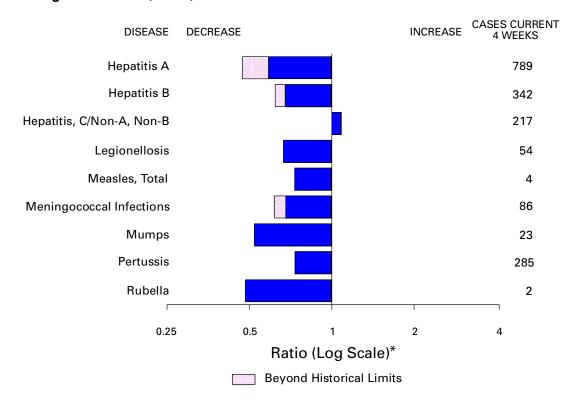
The findings in this report are subject to at least three limitations. First, because this was a passive surveillance effort, persons may have become ill who did not seek medical attention or were not reported to the surveillance system. Second, rates of the health outcomes in the exposed population could not be compared with those for the general population because baseline incidence data for many of the effects attributed to the malathion/bait application are not available. Third, the role of cholinesterase inhibition was not determined because blood cholinesterase levels were not obtained.

Certain malathion formulations are registered by EPA for aerial spraying over urban areas in mosquito-control programs. The use of malathion in these programs provides an important public health benefit by controlling mosquitoes that transmit human diseases such as encephalitis, dengue fever, and malaria. Spraying malathion/bait over urban populations for Medfly eradication has generated controversy in part because these applications are directed not at preventing human illness but at eradicating an agricultural pest. Federal law does not permit spraying malathion/bait over urban areas without an emergency EPA exemption**. To reduce the risk for illness among persons sensitive to the effects of malathion/bait applications, federal and state agricultural authorities are encouraged to pursue and enhance alternative methods for Medfly control. These methods include preventing Medfly importation into the United States, quickly detecting Medfly infestations (e.g. through

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

^{** 40} CFR Part 166.

FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending November 6, 1999, with historical data — United States



^{*}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.

TABLE I. Summary — provisional cases of selected notifiable diseases, United States, cumulative, week ending November 6, 1999 (44th Week)

		Cum. 1999		Cum. 1999
Cyclosporiasi Diphtheria Encephalitis: Ehrlichiosis Hansen Disea Hantavirus pu	California* eastern equine* St. Louis* western equine* human granulocytic (HGE)* human monocytic (HME)*	42 3 5 49 2 50 6 6 128 35 87 18 88	HIV infection, pediatric*§ Plague Poliomyelitis, paralytic Psittacosis* Rabies, human Rocky Mountain spotted fever (RMSF) Streptococcal disease, invasive Group A Streptococcal toxic-shock syndrome* Syphilis, congenital* Tetanus Toxic-shock syndrome Trichinosis Typhoid fever Yellow fever	121 5 - 15 - 465 1,790 31 175 27 98 8 261

^{-:} no reported cases

^{*}Not notifiable in all states.

† Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases (NCID).

§ 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 October 24, 1999.

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

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending November 6, 1999, and November 7, 1998 (44th Week)

									erichia 157:H7*	
	Al	DS	Chla	mydia	Cryptosp	oridiosis	NE.	TSS		LIS
Reporting Area	Cum. 1999 [†]	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998
UNITED STATES	37,420	38,690	505,533	500,295	1,987	3,356	2,877	2,575	1,936	1,977
NEW ENGLAND	1,904	1,517	16,989	17,174	129	142	286	292	293	248
Maine N.H.	68 38	26 25	738 809	894 836	25 17	29 14	35 28	33 42	29	42
Vt.	15	18	414 7,808	357	34 49	26	32	19 125	18 171	17 141
Mass. R.I.	1,231 90	766 110	1,974	7,099 1,944	49	66 7	164 27	135 11	6	141
Conn.	462	572	5,246	6,044	-	-	U	52	69	47
MID. ATLANTIC Upstate N.Y.	9,663 1.146	10,367 1,250	52,184 N	52,034 N	370 144	506 304	249 190	267 190	76 -	84
N.Y. City	5,100	5,843	21,963	22,404	114	180	9	12	15	12
N.J. Pa.	1,741 1,676	1,894 1,380	8,990 21,231	10,051 19,579	36 76	22 N	50 N	65 N	32 29	51 21
E.N. CENTRAL	2,519	2,736	76,100	84,807	451	676	606	405	425	322
Ohio	403	567	19,151	22,939	56 37	68	199 91	108	168	62 47
Ind. III.	285 1,201	446 1,037	9,299 29,105	9,405 22,783	60	52 80	205	91 104	55 81	47 74
Mich. Wis.	504	530	18,545 U	17,813	45 253	37 439	111 N	102	73 48	62 77
W.N. CENTRAL	126 846	156 750	28.087	11,867 29.702	253 191	439 304	549	N 435	46 370	77 377
Minn.	161	146	5,758	5,999	71	125	219	183	158	197
lowa Mo.	72 408	60 363	3,864 9,298	3,824 10,640	53 27	63 25	108 51	89 44	70 57	58 60
N. Dak.	6	5	707	876	18	30	16	10	14	15
S. Dak. Nebr.	13 61	15 60	1,337 2.897	1,325 2,384	7 14	20 35	44 90	30 47	59 -	34
Kans.	125	101	4,226	4,654	1	6	21	32	12	13
S. ATLANTIC	10,275	10,032	108,441	96,421	331	302 3	295 6	218	148 3	161 2
Del. Md.	147 1,242	122 1,394	2,280 9,922	2,213 6,307	- 17	18	36	39	2	14
D.C. Va.	496 689	750 771	N 12,367	N 11,306	8 21	21 20	- 68	1 N	U 52	U 51
W. Va.	61	70	1,204	2,056	3	1	10	9	8	9
N.C. S.C.	688 847	703 638	18,914 10,284	18,723 14,264	22	N	64 19	52 11	49 14	47 11
Ga.	1,466	1,060	29,077	20,362	121	104	29	69	-	-
Fla.	4,639	4,524	24,393	21,190	139	135	63	37	20	27
E.S. CENTRAL Ky.	1,666 236	1,596 248	38,629 6,396	34,842 5,450	26 6	24 10	113 43	109 33	56 -	62
Tenn.	643 423	590 417	11,791 10,761	11,645	6 11	8 N	43 22	49 21	36 16	40 18
Ala. Miss.	364	341	9,681	8,651 9,096	3	6	5	6	4	4
W.S. CENTRAL	3,822	4,742	70,980	76,195	80	894	123	88	109	96
Ark. La.	158 742	177 814	5,075 10,879	3,257 12,687	1 22	6 15	13 9	11 5	8 13	10 7
Okla.	113	254	6,790	8,217	10	N	28	15	24	8
Tex. MOUNTAIN	2,809	3,497	48,236	52,034	47 87	873 110	73	57 335	64 167	71 235
Mont.	1,469 11	1,359 26	26,692 1,287	27,565 1,152	10	119 10	288 24	15	167 -	235 5
ldaho Wyo.	21 10	27 3	1,428 653	1,695 599	7 1	17 2	56 14	38 53	20 14	24 55
Colo.	271	254	5,099	6,581	12	17	106	72	86	61
N. Mex. Ariz.	78 745	188 550	3,151 10,697	3,114 9,820	38 12	46 18	11 30	18 43	5 19	19 26
Utah	129	114	1,808	1,823	N	N	32	72	21	21
Nev.	204	197	2,569	2,781	7	9	15	24	2	24
PACIFIC Wash.	5,256 305	5,591 369	87,431 10,116	81,555 9,523	322 N	389 N	368 144	426 94	292 119	392 124
Oreg.	185	146	5,072	4,796	88	65	73	102	68	95
Calif. Alaska	4,673 13	4,915 17	68,196 1,611	63,468 1,599	234	321 -	141 1	223 7	94 1	159 -
Hawaii	80	144	2,436	2,169	-	3	9	-	10	14
Guam P.R.	5 1,094	1 1,498	302 U	355 U	-	- N	N 5	N 5	U U	U U
V.I.	36	31	U	U	Ų	U	U	U	U	U
Amer. Samoa C.N.M.I.	-	-	U	U U	U U	U U	U U	U U	U U	U U
N. Not potificals	Hilling				CNN		nuscalth of N			

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

^{*}Individual cases may be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the

Public Health Laboratory Information System (PHLIS).

†Updated monthly from reports to the Division of HIV/AIDS Prevention–Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention, last update October 24, 1999.

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending November 6, 1999, and November 7, 1998 (44th Week)

-	Gond	orrhea	Hepa C/N/		Legion	ellosis	Lyr Dise	
Reporting Area	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998
UNITED STATES	282,710	298,684	2,868	2,816	778	1,107	11,143	14,034
NEW ENGLAND	5,214	5,132	59	55	70	76	3,245	4,327
Maine N.H.	42 92	57 79	2	-	3 6	1 7	41 16	73 42
Vt. Mass.	41 2,167	33 1,920	6 48	5 47	13 28	7 31	20 1,033	11 666
R.I. Conn.	,496 2,376	327 2,716	3	3	9 11	19 11	450 1,685	571 2,964
MID. ATLANTIC	33,170	32,371	116	188	165	273	6,303	7,743
Upstate N.Y. N.Y. City	5,751 11,762	6,135 10,115	81 -	97 -	53 9	86 34	3,390 30	3,567 218
N.J. Pa.	5,408 10,249	6,775 9,346	- 35	U 91	18 85	15 138	905 1,978	1,701 2,257
E.N. CENTRAL	55,411	58,589	1,377	595	218	365	1,376	709
Ohio Ind.	11,535 5,136	15,051 5,557	3 1	7 5	65 36	114 62	68 19	42 34
III.	25,722	18,930	39	38	22	48	12	14
Mich. Wis.	13,018 U	13,659 5,392	743 591	412 133	59 36	76 65	1 5	12 607
W.N. CENTRAL Minn.	11,211 2,211	14,561 2,302	260 9	36 9	42 9	60 6	205 137	196 148
Iowa	1,027	1,306	-	8	11 14	9 16	19	24
Mo. N. Dak.	4,686 71	7,499 72	240 -	12 -	1	-	26 1	11 -
S. Dak. Nebr.	160 1,230	195 1,007	- 5	4	3 4	3 18	10	3
Kans.	1,826	2,180	6	3	- 110	8	12	10 700
S. ATLANTIC Del.	81,311 1,415	80,563 1,291	181 1	92	113 11	124 12	1,010 51	790 63
Md. D.C.	8,575 3,166	8,187 3,725	38 1	13 -	26 3	31 6	716 4	563 4
Va. W. Va.	8,206 363	7,685 749	10 17	11 6	28 N	18 N	109 16	59 12
N.C. S.C.	16,786 5,840	16,192	33 22	19 7	14 8	11	66 5	50 6
Ga.	19,795	9,146 17,234	1	9	1	10 8	-	5
Fla. E.S. CENTRAL	17,165 31,009	16,354 33,698	58 213	27 251	22 36	28 59	43 71	28 98
Ky.	2,901	3,183	17	19	18	26	9	25
Tenn. Ala.	9,569 9,691	10,169 11,131	79 1	149 4	14 4	21 <u>5</u>	30 19	41 19
Miss. W.S. CENTRAL	8,848 39,702	9,215 46,691	116 290	79 484	23	7 30	13 43	13 19
Ark.	2,669	3,318	18	21	2	1	4	6
La. Okla.	8,653 3,354	10,860 4,488	102 14	95 14	3	4 12	4	4 2
Tex. MOUNTAIN	25,026 7,922	28,025 7,692	156 126	354 341	18 41	13 65	35 16	7 17
Mont.	45	37	5 7	7 86	-	2 2	5	- -
ldaho Wyo.	71 27	145 29	37	86	2	1	3	1
Colo. N. Mex.	2,032 632	1,744 762	20 8	27 84	11 1	15 2	- 1	4
Ariz. Utah	3,858 181	3,553 192	35 6	11 21	6 15	16 21	- 5	1
Nev.	1,076	1,230	8	19	6	6	2	6
PACIFIC Wash.	17,760 1,787	19,387 1,686	246 16	774 21	70 13	55 12	145 10	135 7
Oreg. Calif.	734 14,630	689 16,306	17 213	18 681	N 56	N 41	11 124	20 107
Alaska Hawaii	260 349	265 441	-	- 54	1	1	N	1 N
Guam	349	61	1	5 4 1	-	2	- IN	1
P.R. V.I.	282 U	327 U	Ü	Ū	- U	- U	N U	N U
Amer. Samoa C.N.M.I.	Ü	Ü	Ŭ	Ŭ	Ŭ	Ŭ	Ü	Ü
C.I V.IVI.I.	U	U	U	U	U	U	U	U

N: Not notifiable

U: Unavailable

-: no reported cases

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending November 6, 1999, and November 7, 1998 (44th Week)

				-		Salmonellosis*						
	Ma	laria	Rabies,	Animal	NE	TSS	PH	LIS				
Reporting Area	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998				
UNITED STATES	1,106	1,265	5,181	6,412	31,605	36,258	24,551	29,849				
NEW ENGLAND Maine	58	55	766 150	1,281 208	1,427 122	2,162 150	1,813 90	2,046				
N.H.	3 2	5 5	48	74	113	170	121	58 204				
Vt. Mass.	4 21	1 17	86 185	59 453	83 989	123 1,177	73 993	97 1,211				
R.I.	4	9	84	85	120	127	147	34				
Conn. MID. ATLANTIC	24 257	18 376	213 983	402 1,401	U 3,721	415 5,793	389 3,244	442 5,200				
Upstate N.Y.	64	83	701	974	1,146	1,405	900	1,238				
N.Y. City N.J.	115 43	212 51	U 155	U 195	1,163 508	1,705 1,283	853 535	1,323 1,218				
Pa.	35	30	127	232	904	1,400	956	1,421				
E.N. CENTRAL Ohio	127 18	131 14	141 33	119 54	4,643 1,138	5,536 1,337	2,986 913	4,234 1,013				
Ind.	18	10	12	11	454	589	355	464				
III. Mich.	46 37	53 43	10 83	N 35	1,425 858	1,698 1,022	399 841	1,354 944				
Wis.	8	11	3	19	768	890	478	459				
W.N. CENTRAL Minn.	63 33	85 51	630 101	631 105	1,983 576	2,022 497	1,952 608	2,074 582				
lowa	13	7	147	136	237	338	195	265				
Mo. N. Dak.	13 -	14 2	14 127	37 122	646 41	545 56	792 48	752 67				
S. Dak. Nebr.	-	- 1	151 3	147 7	85 179	101 166	106	113 42				
Kans.	4	10	87	77	219	319	203	253				
S. ATLANTIC	302	260	1,852	2,100	7,496	7,319	4,599	5,397				
Del. Md.	1 84	3 76	37 349	43 404	123 775	70 811	144 827	108 794				
D.C. Va.	17 63	16 52	- 497	500	65 1,129	64 964	U 872	U 784				
W. Va.	2	2	94	69	138	131	137	146				
N.C. S.C.	26 17	25 6	371 132	517 134	1,158 590	1,058 566	1,173 418	1,242 480				
Ga. Fla.	21 71	33 47	204 168	261 172	1,258 2,260	1,442 2,213	651 377	1,342 501				
E.S. CENTRAL	21	31	229	247	1,645	2,002	924	1,404				
Ky. Tenn.	7 6	7 15	34 82	29 126	359 317	324 519	473	124 619				
Ala.	7	6	112	90	536	608	374	520				
Miss.	1	3	1	2	433	551	77	141				
W.S. CENTRAL Ark.	16 3	34 1	89 14	28 28	3,335 571	4,163 529	2,838 120	2,838 329				
La. Okla.	10 2	14 3	- 75	- N	334 380	619 434	472 291	718 204				
Tex.	1	16	-	-	2,050	2,581	1,955	1,587				
MOUNTAIN Mont.	41 4	60 1	177 54	233 51	2,646 53	2,235 72	2,200 1	1,797 43				
ldaho	3	8	-	N	99	108	81	85				
Wyo. Colo.	1 16	- 18	42 1	55 42	63 633	57 478	49 644	51 453				
N. Mex.	2 8	12 8	9 58	6 47	341 836	265 723	217 692	232				
Ariz. Utah	4	1	8	26	455	314	463	613 122				
Nev.	3	12	5	6	166	218	53	198				
PACIFIC Wash.	221 25	233 17	314 -	372 -	4,709 577	5,026 438	3,995 670	4,859 577				
Oreg. Calif.	19 166	15 194	1 306	7 342	382 3,399	269 4,023	452 2,610	290 3,697				
Alaska	1	2	7	23	51	53	15	32				
Hawaii	10	5 2	-	-	300 24	243 34	248 U	263 U				
Guam P.R.	- . .	-	63	47	255	694	U	U				
V.I. Amer. Samoa	U U	U U	U U	U U	U U	U U	U U	U U				
C.N.M.I.	ŭ	ŭ	ŭ	ŭ	Ŭ	ŭ	ŭ	ŭ				

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

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending November 6, 1999, and November 7, 1998 (44th Week)

		Shige			Sypt			•
	NE	TSS		ILIS	(Primary &		Tubero	culosis
Reporting Area	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999†	Cum. 1998†
UNITED STATES	13,368	18,442	6,167	10,449	5,709	6,105	11,715	14,211
NEW ENGLAND	642	378	638	331	50	65	353	378
Maine N.H.	5 16	12 15	14	19	1	1 2	16 10	11
Vt.	6	6	4	1	3	4	1	4
Mass. R.I.	592 23	249 34	563 9	238 13	31 2	37 1	201 35	214 49
Conn.	Ü	62	48	60	13	20	90	100
MID. ATLANTIC	806	2,115	398	1,565	218	272	2,118	2,474
Upstate N.Y. N.Y. City	246 242	533 641	45 82	188 550	24 79	35 64	266 1,120	325 1,217
N.J.	195	611	121	583	48 67	85	442 290	520
Pa. E.N. CENTRAL	123 2,424	330 2,506	150 1,121	244 1,342	1,326	88 890	1,106	412 1,407
Ohio	370	435	116	117	75	124	200	204
Ind. III.	280 919	143 1,377	90 592	36 1,122	592 451	170 363	82 495	131 665
Mich.	388	236	255	4	208	176	246	316
Wis.	467	315	68	63	U	57	83	91
W.N. CENTRAL Minn.	993 218	916 276	617 209	537 306	103 9	115 9	409 169	404 125
lowa	51	63	44	43	9	2	39	40
Mo. N. Dak.	607 2	131 8	320 2	95 3	67	86	143 6	151 8
S. Dak.	13	31	6	21	-	1	17	16
Nebr. Kans.	65 37	345 62	36	19 50	8 10	4 13	16 19	23 41
S. ATLANTIC	2,090	3,670	397	1,139	1,743	2,241	2,448	2,611
Del.	12	34	8	28	. 8	20	12	32
Md. D.C.	141 46	186 25	49 U	63 U	305 59	589 71	230 35	265 94
Va. W. Va.	117 8	177 11	48 5	80 7	136 2	132 2	247 35	250
N.C.	185	264	79	155	400	636	348	36 365
S.C. Ga.	111 204	158 957	56 37	79 225	230 343	294 248	210 521	246 438
Fla.	1,266	1,858	115	502	260	249	810	885
E.S. CENTRAL	927	1,080	450	832	967	1,060	745	1,003
Ky. Tenn.	220 508	117 490	393	45 572	88 532	92 498	156 272	138 355
Ala.	106	423	47	208	190	247	261	323
Miss.	93	50	10	7	157	223	56	187
W.S. CENTRAL Ark.	2,312 73	3,713 191	1,806 23	1,169 58	818 66	915 100	1,253 145	2,112 122
La.	118 447	294	99 149	255 130	200	363 79	U 112	246
Okla. Tex.	1,674	446 2,782	1,535	726	163 389	373	112 996	146 1,598
MOUNTAIN	977	1,120	560	655	205	217	372	481
Mont. Idaho	7 25	8 18	9	3 13	1 1	2	10 14	18 10
Wyo.	3	18 3	1	1	-	1	3	4
Colo. N. Mex.	169 113	186 265	123 62	143 153	2 11	10 22	U 52	56 58
Ariz.	524	546	341	295	182	163	183	187
Utah Nev.	58 78	39 55	18 6	28 19	2 6	4 15	35 75	45 103
PACIFIC	2,197	2,944	180	2,879	279	330	2,911	3,341
Wash.	101 78	191 172	79 74	163	63 9	27 4	151	227 119
Oreg. Calif.	1,989	2,527	-	138 2,527	204	295	86 2,481	2,797
Alaska	3 26	8 46	2 25	5 46	1 2	1 3	48 145	46 152
Hawaii Guam	2 0 8	46 34	25 U	46 U	1	3 1	145	78
P.R.	62	53	U	U	140	158	41	140
V.I. Amer. Samoa	U U	U U	U U	U U	U U	U U	U U	U U
C.N.M.I.	Ü	ŭ	ŭ	ŭ	Ŭ	ŭ	ŭ	Ŭ

N: Not notifiable U: Unavailable -: no reported cases
*Individual cases may 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).

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending November 6, 1999, and November 7, 1998 (44th Week)

	H. influ	ienzae,		epatitis (Vi		Measles (Rubeola)							
		sive		A	E		Indi	genous		orted*		tal	
Reporting Area	Cum. 1999 [†]	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998	1999	Cum. 1999	1999	Cum. 1999	Cum. 1999	Cum. 1998	
UNITED STATES	981	919	14,458	19,074	5,414	8,169	-	54	-	23	77	84	
NEW ENGLAND	82	63	242	246	85	186	-	6	-	5	11	3	
Maine N.H.	7 17	3 10	11 15	17 12	1 13	4 18	-	-	-	1	1	-	
Vt. Mass.	5 31	8 36	18 78	15 109	3 34	8 67	-	- 5	-	3	- 8	1 2	
R.I.	5	5	21	15	34	63	-	-	-	-	-	-	
Conn. MID. ATLANTIC	17 154	1	99	78 1 497	-	26	-	1	-	1	2	- 14	
Upstate N.Y.	154 74	147 50	836 235	1,487 309	528 161	1,056 203	-	-	-	2 2	2 2	14 2	
N.Y. City N.J.	34 45	39 50	244 112	518 309	168 41	368 183	-	-	-	-	-	- 8	
Pa.	1	8	245	351	158	302	-	-	-	-	-	4	
E.N. CENTRAL Ohio	150 51	157 45	2,421 566	3,071 270	563 81	1,233 68	- U	1	- U	2	3	15 1	
Ind.	22	39	100	132	36	97	-	1	-	1	2	3	
III. Mich.	63 13	55 11	574 1,123	681 1,813	1 431	209 394	-	-	-	- 1	- 1	10	
Wis.	1	7	58	175	14	465	U	-	U	-	-	1	
W.N. CENTRAL	84 40	83	750 75	1,228	313 49	351 43	-	2	-	-	2 1	-	
Minn. Iowa	9	64 2	123	115 388	34	51	-	1 -	-	-	-	-	
Mo. N. Dak.	26 1	10	451 2	573 3	188	210 4	-	1 -	-	-	1	-	
S. Dak.	1	-	9	31	1	2	U	-	U	-	-	-	
Nebr. Kans.	3 4	1 6	50 40	25 93	14 27	18 23	Ū	-	Ū	-	-	-	
S. ATLANTIC	214	162	1,740	1,668	1,038	853	-	10	-	5	15	8	
Del. Md.	- 55	- 50	2 316	3 359	1 145	3 120	U	-	U	-	-	1 1	
D.C.	4	-	54	55	21	11	U	-	U	-	-	-	
Va. W. Va.	17 6	16 6	149 32	182 7	77 22	90 8	Ū	10 -	Ū	3	13 -	2	
N.C. S.C.	31 5	23 3	140 42	110 34	204 63	195 39	-	-	-	-	-	-	
Ga.	55	39	423	549	155	127	-	-	-	-	-	2	
Fla.	41	25	582	369	350	260	-	-	-	2	2	2	
E.S. CENTRAL Ky.	52 6	53 7	339 61	352 29	347 37	435 42	-	2 2	-	-	2 2	2	
Tenn. Ala.	28 15	31 13	142 49	199 66	165 76	241 67	-	-	-	-	-	1 1	
Miss.	3	2	87	58	69	85	-	-	-	-	-	-	
W.S. CENTRAL Ark.	45 2	48	3,481 50	3,419 78	773 58	1,813 95	-	8 3	-	4	12 3	-	
La.	7	20	73	92	77	145	Ū	- -	Ū	-	-	-	
Okla. Tex.	32 4	25 3	402 2,956	520 2,729	110 528	87 1,486	-	- 5	-	4	9	-	
MOUNTAIN	97	106	1,119	2,822	497	715	_	3	_	-	3	2	
Mont. Idaho	3 1	- 1	17 36	89 224	17 26	5 38	U	-	U	-	-	-	
Wyo.	1	1	7	35	12	9	-	-	-	-	-	-	
Colo. N. Mex.	11 18	21 6	197 43	281 135	82 155	93 275	Ū	-	Ū	-	-	-	
Ariz.	52	54	657	1,687	130	159	-	1	-	-	1	2	
Utah Nev.	8 3	4 19	46 116	172 199	31 44	65 71	U U	2	U U	-	2	-	
PACIFIC	103	100	3,530	4,781	1,270	1,527	-	22	-	5	27	40	
Wash. Oreg.	5 38	8 38	286 218	884 385	60 81	95 169	Ū	9	Ū	-	9	1 -	
Calif. Alaska	46 6	44 3	3,004 9	3,444 16	1,102 14	1,237 13	-	13	-	4	17 -	8 31	
Hawaii	8	7	13	52	13	13	-	-	-	1	1	-	
Guam	-	-	2	1	2	2	U	1	U	-	1	-	
P.R. V.I.	1 U	2 U	112 U	63 U	102 U	220 U	Ū	Ū	Ū	Ū	Ū	Ū	
Amer. Samoa C.N.M.I.	U U	U	U U	U U	U	U U	U U	U U	U U	U	U U	U U	

N: Not notifiable

U: Unavailable

^{-:} no reported cases

^{*}For imported measles, cases include only those resulting from importation from other countries.

†Of 187 cases among children aged <5 years, serotype was reported for 98 and of those, 26 were type b.

TABLE III. (Cont'd.) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending November 6, 1999, and November 7, 1998 (44th Week)

	_	ococcal			7, 100	7 (4461)	vveek)		B			
	Dise Cum.	ease Cum.		Mumps Cum.	Cum.		Pertussis Cum.	Cum.		Rubella Cum.	Cum.	
Reporting Area	1999	1998	1999	1999	1998	1999	1999	1998	1999	1999	1998	
UNITED STATES	2,020	2,263	6	296	577	57	4,718	5,584	1	227	345	
NEW ENGLAND Maine	100 5	103 6	-	8 -	8	8 -	558 -	888 5	-	7 -	38	
N.H. Vt.	12 5	11 5	-	1 1	-	3	78 60	103 69	-	-	-	
Mass. R.I.	58 5	49 7	-	4 2	5 1	-	360 33	662 9	-	7	8 1	
Conn.	15	25	-	-	2	5	27	40	-	-	29	
MID. ATLANTIC Upstate N.Y.	185 58	242 65	-	30 10	184 6	32 32	780 645	560 291	-	22 18	146 114	
N.Y. City	46	30	-	3	155	-	10	37	-	-	18	
N.J. Pa.	44 37	55 92	-	17	6 17	-	12 113	24 208	-	1 3	13 1	
E.N. CENTRAL Ohio	347 122	348 125	- U	34 14	74 27	4 U	415 184	725 247	- U	2	-	
Ind.	58	63	-	4	7	1	63	149	-	1	-	
III. Mich.	93 42	87 42	-	9 7	10 27	1 2	66 54	106 59	-	1 -	-	
Wis. W.N. CENTRAL	32 223	31 195	U -	- 13	3 31	U 1	48 335	164 511	U 1	- 124	- 39	
Minn.	49	30	-	1	13	-	187	291	-	5	-	
lowa Mo.	40 90	38 70	-	7 2	10 3	1 -	49 51	66 35	1	29 3	2	
N. Dak. S. Dak.	3 11	5 7	Ū	-	2	Ū	4 5	3 8	- U	-	-	
Nebr. Kans.	12 18	15 30	Ū	3	3	Ū	4 35	16 92	Ū	87	- 37	
S. ATLANTIC	356	374	1	47	45	2	359	279	-	36	18	
Del. Md.	8 51	2 27	U 1	6	-	U 1	5 98	5 55	U -	- 1	- 1	
D.C. Va.	1 47	1 36	U -	2 10	- 8	U -	- 29	1 30	U -	-	- 1	
W. Va. N.C.	6 40	16 53	U	- 8	11	U	3 86	1 91	U	- 35	: 13	
S.C.	42	52	-	4	6	-	15	26	-	-	-	
Ga. Fla.	58 103	86 101	-	4 13	1 19	1 -	38 85	24 46	-	-	3	
E.S. CENTRAL Ky.	125 28	177 33	1	13	15	-	72 21	112 49	-	1	2	
Tenn.	43	63	-	-	1	-	27	33	-	-	2	
Ala. Miss.	32 22	47 34	1 -	10 3	8 6	-	21 3	24 6	-	1 -	-	
W.S. CENTRAL Ark.	166 31	269 27	-	33	55 12	1	157 18	333 77	-	15 6	87	
La.	34	52	U	3	7	U	3	9	U	-	-	
Okla. Tex.	27 74	38 152	-	1 29	36	1	12 124	31 216	-	9	87	
MOUNTAIN Mont.	125 2	127 4	1 U	26	36	8 U	619 2	963 9	- U	16	5	
Idaho	10	10	1	2	4	4	139 2	215	-	-	-	
Wyo. Colo.	4 32	5 23		5	1 6	4	183	8 243	-	1		
N. Mex. Ariz.	14 42	25 39	N -	N 8	N 6	U -	133 100	89 191	U -	13	1 1	
Utah Nev.	14 7	13 8	U U	6 5	5 14	U U	55 5	167 41	U U	1 1	2 1	
PACIFIC	393	428	3	92	129	1	1,423	1,213	-	4	10	
Wash. Oreg.	61 69	58 74	- N	2 N	10 N	1 U	594 47	284 85	- U	-	5 -	
Calif. Alaska	251 5	288 3	3	76 2	93 2	-	745 4	813 14	-	4	3	
Hawaii	7	5	-	12	24	-	33	17	-	-	2	
Guam P.R.	2 5	2 9	U -	1 -	5 3	U -	1 16	1 6	U -	-	14	
V.I. Amer. Samoa	Ü	Ü	U U	U U	Ü	U U	Ü	Ü	U U	U U	U U	
C.N.M.I.	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	ŭ	

N: Not notifiable

U: Unavailable

-: no reported cases

TABLE IV. Deaths in 122 U.S. cities,* week ending November 6, 1999 (44th Week)

	P	All Cau	ıses, By	/ Age (Y	ears)		P&I [†]		/	All Cau	ıses, By	/ Age (Y	ears)		P&l [†]
Reporting Area	All Ages	>65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	>65	45-64	25-44	1-24	<1	Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn. Cambridge, Mass. Fall River, Mass. Hartford, Conn. Lowell, Mass. Lynn, Mass. New Bedford, Mass. New Haven, Conn. Providence, R.I. Somerville, Mass.	46 62 2 40	186 U 8 23 U 13 7 16 25 41 22 32 19	U 1 3 U 3 4 3 11 13	22 UU 2 - U - 2 3 8 5 - 2	4 U U - 1 U - - 1 1 - 1	5 UU - - - - 1 3 - 1	29 UU36U31135'34	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla. Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, Fla. Tampa, Fla. Washington, D.C. Wilmington, Del.	777 U 150 83 116 U 42 63 50 98 145 U	494 U 80 51 73 U 28 33 39 71 98 U 21	170 U 36 22 29 U 11 19 7 12 25 U	68 U 22 5 7 U 9 1 9	31 U 10 4 4 U 3 - 2 4 U	13 U 2 1 3 U - 2 1 2 2 U -	51 U 12 7 4 U 1 5 3 5 14 U
Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Elizabeth, N.J. Jersey City, N.J. New York City, N.Y. 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.	21 U 995 41 U 80 20 13 49 54 U U 27 27 389 48 31 109 U 28 75 15 21 U	725 29 0 60 14 11 35 38 0 0 18 267 29 85 0 57 12 17	U 170 8 U 12 4 10 U U 6 6 79 11 2 15 U 2 11 2 4	0 61 2 U 5 1 2 6 6 U U 1 25 5 4 U 1 2 1 - U	18 1 U 1 1 1 1 1 1 1 3 U 1 1 1 1 1 1 1 1 1	20 1 U 1 4 - U U 2 7 3 - 2 U U	4U 565U6''4'UU'142511U171'U	E.S. CENTRAL Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala. Nashville, Tenn. W.S. CENTRAL Austin, Tex. Baton Rouge, La. Corpus Christi, Tex. Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La. San Antonio, Tex. Shreveport, La. Tulsa, Okla.	97 70 196 62 U 121 1,338 70 11	530 114 500 67 48 134 41 U 76 894 48 10 46 111 54 84 203 52 U 161 355 90	177 43 17 21 15 37 13 U 31 280 15 1 10 47 15 29 79 19 U 38 5 5 29 79 29 29 29 20 20 20 20 20 20 20 20 20 20 20 20 20	60 10 5 3 20 7 10 88 5 2 12 1 9 29 5 U 17 5 3	14 5 2 2 1 2 - U 2 31 1 2 5 4 2 10 2 0 3 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	11 1 1 2 2 3 1 U 1 45 1 - 9 2 3 15 3 U 3 3 6	73 17 5 10 10 18 1 1 12 113 5 1 10 13 7 10 34 3 13 5 12
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, Ill. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Gary, Ind. Grand Rapids, Mich Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, Ill. Rockford, Ill. South Bend, Ind. Toledo, Ohio Youngstown, Ohio W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	1,910 39 36 362 65 124 191 121 200 47 53 23 24 143 26 112 45 65 59 92 64 446 U 30 35 87 38	1,322 31 29 206 427 139 90 117 37 42 122 122 37 98 34 52 40 70 52 323 U 21 56 30 105 U U U U U U U U U U U U U U U U U U U	363 2 4 815 30 32 255 8 9 4 5 25 7 14 6 8 11 15 8 72 U 5 7 11 28 U U 6	144 52 47 34 15 81 12 13 12 13 12 23 33 52 29 0 - 50 10 47 70 0 0 1 2	421 - 131 122 - 6 - 131 4 121 22 9U - 15 - 2UU - 1	377 - 1 111 4 1 1 3 1 1 4 4 1 1 2 2 4 4 4 1 1 1 5 5 4 U U U - 2	13123622110153713102 · 553883 31U43519UU63	MOUNTAIN Albuquerque, N.M. Boise, Idaho Colo. Springs, Colo Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, Utah Tucson, Ariz. PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawaii Long Beach, Calif. Los Angeles, Calif. Pasadena, Calif. Portland, Oreg. Sacramento, Calif. San Diego, Calif. San Francisco, Calif. San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash.	807 107 47 . 65 . U 156 . U 156 . U 134 1,113 . U 111 . U 70 . 49 . U . 21 . U . U . 27 . 139 . 27 . 139 . 63 . 100	544 74 74 34 41 104 U 101 129 66 95 783 U 82 U 46 36 36 U 19 U 107 17 7 75 49 69 5,801	172 18 10 16 0 36 0 40 9 15 28 197 17 10 8 0 2 0 0 16 0 17 0 17 0 17 0 17 0 17 0 17 0	58 10 3 5 11 11 16 3 3 7 9 0 0 7 0 9 1 1 1 2 1 2 1 1 2 1 6 5 6 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 6 7 6 7 6 6 7 7 6 7 6 7 6 7 6 7 7 6 7 6 7 7 7 6 7 7 6 7 6 7 7 6 7 6 7 7 6 7 7 6 7 7 6 7 7 6 7 7 6 7 7 6 7 7 6 7 7 6 7 7 6 7 7 6 7 6 7 7 6 7 6 7 7 6 7 6 7 7 6 7 7 6 7 6 7 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 6 7 7 6 7 7 6 7 7 6 7	193 - 1 U 2 U 7 2 2 2 2 2 3 U 5 U 1 2 U - U U 4 U 4 2 1 1 1 3 3 191	13 2 - 2 U 2 U 2 U 4 - 1 1 2 18 U U 4 2 U U U 3 3 U 5 - 2 2 175	62 66 37 11 11 12 10 10 78 11 10 76 10 10 10 10 10 10 10 10 10 10 10 10 10

U: Unavailable -: no reported cases

*Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of 100,000 or more. 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.

increased sentinel trapping densities), releasing sterile male Medflies to interrupt the reproductive cycle, and identifying and using safer eradication agents.

During aerial malathion applications for mosquito control and Medfly eradication, the public should be advised to stay indoors and, when appropriate, persons with exposure-related health concerns should seek medical attention. The public also should be provided with an opportunity to ask questions and receive timely responses about the malathion applications (i.e., through telephone hotlines and community meetings). When malathion/bait applications are used for Medfly eradication, additional precautions are recommended, including immediately washing any skin surfaces that come into contact with malathion/bait-contaminated surfaces; providing advance public notification of spray schedules; performing aerial malathion/bait applications when residents are usually indoors (e.g., at night); directing the homeless to shelters; advising highly sensitive persons to leave the area during spraying; and convening a health advisory committee, an action that has been shown to be useful for mitigating risk (7). Medfly Eradication Program workers should be trained in the safe handling of pesticides, and consideration should be given to measuring plasma and red blood cell cholinesterase in these workers before beginning exposure and periodically thereafter (8). Workers should wear the personal protection equipment (PPE) listed on the pesticide label. Supplementary PPE also may be indicated.

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Erratum: Vol. 48, No. 43

In the article, "Tobacco Use—United States, 1900–1999," on page 989, in the last line of the fourth full paragraph, reference (15) should be (17), and on page 990, in the first line of the second paragraph under Future Challenges, reference (17) should be (18).

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