



MORBIDITY AND MORTALITY WEEKLY REPORT

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Varicella-Related Deaths Among Children — United States, 1997

During the first quarter of 1998, the Texas Department of Health and the lowa Department of Public Health notified CDC of three fatal cases of varicella (chickenpox) that occurred in children during 1997. All three children were unvaccinated. Two children contracted chickenpox from unvaccinated siblings, and the mode of exposure was unknown for the third. This report summarizes these cases and indicates that varicella-related deaths continue to occur among children in the United States despite the availability of vaccine and recommendations for its use in all susceptible children (1,2).

Case 1

On February 28, 1997, a previously healthy, unvaccinated 21-month-old boy developed a typical varicella rash. He had no reported exposure to varicella. On March 1, he was taken to a local emergency department (ED) with a high fever and was started on oral acetaminophen and diphenhydramine. On March 3, his primary-care physician prescribed oral acyclovir. On March 4, his mother noted a new petechial-like rash. The next morning, his primary-care physician noted lethargy, a purpuric rash, and poor perfusion. He was transferred to a local ED. Fluid resuscitation and intravenous ceftriaxone were initiated, but the child continued to deteriorate rapidly, requiring intubation, mechanical ventilation, and inotropic support with dopamine. Blood cultures were negative for bacterial pathogens. Laboratory tests indicated disseminated intravascular coagulation and severe dehydration. Approximately 1½ hours after arrival at the ED, he was transported to a tertiary-care center. Within 10 minutes of arrival, he suffered cardiac arrest and died. The death was attributed to varicella with hemorrhagic complications.

Case 2

On December 21, 1997, a 5-year-old unvaccinated boy with a history of asthma was taken to a local ED with a fever of 104.5 F (40.3 C) and a typical varicella rash in multiple stages of healing. The child was treated with antipyretic and antipruritic medications and discharged.

That evening, the boy developed mild dyspnea and was treated at home for a presumed asthma attack with metered-dose inhalers and one dose of oral prednisone. He Varicella-Related Deaths — Continued

returned to the ED on December 22 with shortness of breath and a 4-hour history of abdominal and leg pain. On presentation to the ED, one of the patient's siblings had active varicella and another had recently recovered from varicella. Physical examination revealed numerous chickenpox lesions, one of which appeared infected. He was tachypneic, and his extremities were mottled consistent with peripheral septic emboli. Chest and abdominal radiographs revealed a right pleural effusion, pneumonia, and mild ileus. Thoracostomy produced pleural fluid containing gram-positive cocci, confirmed 8 hours later to be group A *Streptococcus* (GAS). A peripheral blood sample revealed gram-positive cocci. He was admitted to the hospital and treated with intravenous ceftriaxone, nafcillin, and acyclovir.

After admission, his breathing became labored and his extremities increasingly mottled. He rapidly developed hypotension, obtundation, and bradycardia. Despite efforts at cardiopulmonary resuscitation, the child died 5 hours after arriving at the ED. A post-mortem examination attributed the death to GAS septicemia, pneumonia, and pleural effusion, complicating varicella infection.

Case 3

On December 14, 1996, a previously healthy, unvaccinated 23-month-old boy developed fever and a typical varicella rash. Approximately 1-2 weeks earlier, his unvaccinated 4-year-old sibling had contracted varicella. He was taken to his physician on December 17 because of persistent fever and cellulitis of the left foot, and he was hospitalized on December 19 for failure to improve on an unspecified outpatient antibiotic regimen. Because his condition deteriorated despite intravenous methicillin and ceftriaxone, he was transferred to a regional hospital on December 21. Sepsis, possible viral meningoencephalitis, and mild pleural effusion were diagnosed. A cerebrospinal fluid examination revealed lymphocytic pleocytosis, and blood and urine cultures grew penicillin-resistant Staphylococcus aureus. Antibiotics were changed to nafcillin and gentamycin, and intravenous acyclovir was added on December 23. On December 24, the child developed an aortic insufficiency murmur, and an echocardiogram revealed a 9x9 mm vegetation on the aortic valve, consistent with bacterial endocarditis. Serial echocardiograms displayed growth of the vegetation and development of a pericardial effusion. He was transferred to a cardiac surgery center on December 26. While awaiting surgery, he developed refractive heart failure secondary to staphylococcal endocarditis. He became incoherent, probably secondary to a major embolic neurologic event, and died on January 8, 1997.

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Editorial Note: The three cases described in this report indicate that healthy children continue to die from complications of varicella, a disease that is preventable through vaccination. Although commonly viewed as a benign disease of childhood, serious complications and death can occur following varicella. Varicella is the leading cause of vaccine-preventable deaths in children in the United States.

During 1990–1994, varicella was the underlying cause of death in an average of 43 children aged <15 years each year (CDC, unpublished data, 1998). During

Varicella-Related Deaths — Continued

1988–1995, up to 10,000 children were hospitalized each year for varicella or its complications (CDC, unpublished data, 1998). Ninety percent of the children who died did not have high-risk conditions for severe varicella. The most common severe complications from varicella among fatal cases in children are secondary bacterial infections and pneumonia. Other complications include encephalitis, hemorrhagic complications, hepatitis, arthritis, and Reye syndrome. Reports of severe invasive infections from GAS-complicating varicella have heightened awareness that varicella is a well-defined risk factor for GAS disease (3,4).

Varicella vaccine was licensed in the United States in March 1995, is widely available, and is recommended for routine vaccination of children aged 12–18 months and for vaccination of susceptible older children, adolescents, and adults (1,2). The Vaccines For Children (VFC) program provides varicella vaccine for VFC-eligible children aged >12 months who were born on or after January 1, 1983, and for VFC-eligible children aged <19 years who are family members of an immunocompromised person.

National coverage levels among children aged 19–35 months for varicella vaccine have increased from 14% during July–September 1996 to 25% during March–June 1997 (5). Barriers to vaccine use include the perception that varicella is a benign disease, concerns that immunity will not persist, the potential that varicella disease burden will shift to older age groups among whom the disease is more severe, and concerns about vaccine efficacy and safety (4). A recent study documented 100% vaccine efficacy for prevention of moderate or severe varicella and 86% for prevention of all varicella (6). In addition, vaccinated children who developed varicella caused by wild virus or "breakthrough disease" had very mild disease of short duration with <50 lesions (7). Persistence of immunity for more than 20 years post vaccination has been demonstrated (8). As disease incidence and exposure to wild virus declines, continuing surveillance will determine the need for and timing of additional doses of vaccine.

To monitor the impact of varicella vaccination programs throughout the United States, varicella surveillance is needed, and surveillance for varicella deaths in all states is a key first step in this process. States also are encouraged to develop additional sustainable surveillance systems, including monitoring hospitalizations and establishing statewide aggregate reporting for cases by schools, day care centers, and/or health-care provider offices, and to consider instituting vaccine requirements for day care and school entry (1).

Efforts to increase routine and catch-up varicella vaccination among children should include educating health-care providers that deaths and severe morbidity from varicella are preventable (1,2). Policies that delay vaccination of susceptible children until adolescence accept the considerable disease burden that occurs among children aged 2–11 years. The most effective vaccination strategy focuses on vaccinating children routinely at age 12–18 months and vaccinating all susceptible older children and adolescents. Children have the highest disease incidence and are the group that serve as the primary source of transmission of varicella to groups at higher risk for severe disease, including adults (9) and persons who are not eligible for vaccination. Most deaths and severe morbidity from varicella in children and in adults can be prevented by implementing recommended policies for childhood vaccination.

Varicella-Related Deaths — Continued

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Pregnancy-Related Death Associated with Heparin and Aspirin Treatment for Infertility, 1996

In 1996, a 38-year-old nulliparous woman died from complications of a cerebral hemorrhage. She was approximately 9 weeks' pregnant with triplets at the time of her death. The patient had undergone in vitro fertilization (IVF) and was being treated with anticoagulants (heparin and aspirin) and intravenous immunoglobulin at the time of her death. This report summarizes the investigation of this case by state and county health departments with assistance from CDC.

The patient had undergone 3 years of infertility therapy, including the use of clomiphene citrate with intrauterine insemination, before beginning IVF in 1995. She had no history of recurrent pregnancy loss at initiation of IVF. Her infertility workup included a normal hysterosalpingogram; her husband had a normal semen analysis. An autoantibody screen revealed positive antithyroid antibodies (antimicrosomal [76.0 μ g/mL] and antithryroglobulin [19.9 μ g/mL]; normal: <0.5 μ g/mL for both assays). Antiphospholipid antibodies were negative. In 1985, she had a transphenoidal resection of a pituitary adenoma, with normal prolactin levels thereafter.

She underwent three IVF cycles (ovulation induction, IVF, and embryo transfer). The first ended with a spontaneous abortion at 8 weeks in 1995; the second IVF cycle did not result in a pregnancy; and the third cycle resulted in a pregnancy with triplets in 1996. The patient was treated with estrogen and progesterone during each pregnancy. In addition, with each IVF cycle she received 5000 units heparin subcutaneously twice a day, 81 mg aspirin daily, and intravenous gamma globulin each month. Platelets and prothrombin time (PT) and partial thromboplastin time (PTT) were normal throughout her treatment.

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During her ninth week of pregnancy, the patient experienced an acute headache, anxiety, and nausea while visiting a clinic. She was transferred to a general hospital and lost consciousness en route. On admission to the hospital, she underwent immediate radiologic and neurosurgic evaluation. Her platelets and PT and PTT were normal. Neurosurgery identified a hemorrhagic arteriovenous malformation, which was surgically clipped. A postoperative computerized axial tomography (CAT) scan revealed no rebleeding, but her condition worsened. Massive cerebral swelling could not be controlled, and her condition became critical. On her third day of hospitalization, she was pronounced brain-dead, and life support was discontinued the following day.

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Editorial Note: CDC, in collaboration with state health departments, maintains a pregnancy-related mortality surveillance system. In 1990, CDC received reports of 417 pregnancy-related deaths in the United States. A pregnancy-related death is one that occurs during or within 1 year of pregnancy and was caused by the pregnancy or its complications. No national surveillance system exists for morbidity associated with infertility therapy.

Treatment of IVF patients with immunotherapy (anticoagulation or immunoglobulin) is aimed at preventing early pregnancy loss. Heparin and aspirin therapy substantially reduces the risk for recurrent spontaneous abortion (more than two pregnancy losses) for women with elevated antiphospholipid antibodies (APA) (1) by modifying the effect of APA on platelet activity, which can cause placental thrombosis and lead to fetal loss (2). Heparin and aspirin are widely used in the United States to treat women with recurrent spontaneous abortion and APA. However, the woman described in this report had no antiphospholipid antibodies and no history of recurrent spontaneous abortion at the initiation of her infertility therapy.

Two recent studies have investigated the role of treating IVF patients with heparin and aspirin to prevent early pregnancy loss. One study documented higher pregnancy rates among women with APA following IVF cycles treated with heparin and aspirin (3). A prospective nonrandomized study did not demonstrate substantially higher pregnancy rates among women with APA undergoing IVF when treated with heparin and aspirin (4). A randomized prospective study investigating the efficacy of heparin and aspirin in women undergoing IVF is under way (4).

Anticoagulation therapy can increase the risk for fatal hemorrhagic stroke (5,6). The inhibition of platelet activity with aspirin doses lower than 81 mg daily are well documented (7). Although heparin decreases the risk for death from pulmonary embolism in surgical patients, it has been associated with increased postoperative bleeding (8). A meta-analysis of randomized clinical trials of low-dose heparin (5000 units/twice daily) to prevent thromboembolism demonstrated an increase in wound hematoma formation associated with heparin treatment (9). In surgical patients receiving heparin, the concomitant use of aspirin has been associated with increased risk for serious bleeding (10).

Although data about the risks and benefits of anticoagulation and immunoglobulin therapy in IVF patients are limited, use of this therapy is becoming more common in

Pregnancy-Related Death — Continued

the United States. Neither aspirin or heparin, alone or in combination, are approved by the Food and Drug Administration (FDA) for this use. In July 1997, a survey of medical practices that provide assisted reproductive technology services indicated that combination therapies of heparin and aspirin for infertility treatment were used at least once by 74% of respondents (Society for Assisted Reproductive Technology, unpublished data, 1997). Of those providing immunotherapy treatment, 94% reported that they considered women who had had recurrent spontaneous abortions as potential candidates for anticoagulation treatment. In addition, 49% considered women who previously had an unsuccessful IVF attempt as potential candidates for immunologic treatment, and 19% considered new IVF patients as potential candidates for therapy.

This case is the first reported pregnancy-related death associated with the use of heparin and aspirin for infertility. The patient died from a cerebral hemorrhage associated with a congenital arteriovenous malformation. Although a causal relation between anticoagulation and hemorrhage from an arteriovenous malformation cannot be established, pregnant women have the risks for bleeding associated with anticoagulation therapy found in the general population (cerebrovascular accidents, gastric ulcers, and trauma) in addition to unique hemorrhagic risks such as ectopic pregnancy. Both heparin and aspirin therapy have been associated with increased risks for and severity of bleeding. The patient in this report did not have recurrent spontaneous abortions or a history of antiphospholipid antibodies, widely accepted as indications for heparin and aspirin therapy. Because the potential for bleeding exists with heparin and aspirin, the risks for and benefits of anticoagulation therapy to improve success rates in IVF patients require vigorous scientific investigation before being accepted as routine practice.

The regular monitoring of all pregnancy-related deaths is essential to the reproductive health of women. To further assess the potential health threat of anticoagulation therapy in the treatment of infertility, CDC requests that deaths or severe morbidity associated with the use of heparin and aspirin for the prevention of pregnancy loss be reported to CDC, telephone (770) 488-5372, or to FDA's MedWatch, telephone (800) 332-1088. Until the results of further studies are available, women undergoing IVF and their health-care providers should carefully review all information about the risks and benefits of heparin and aspirin therapy.

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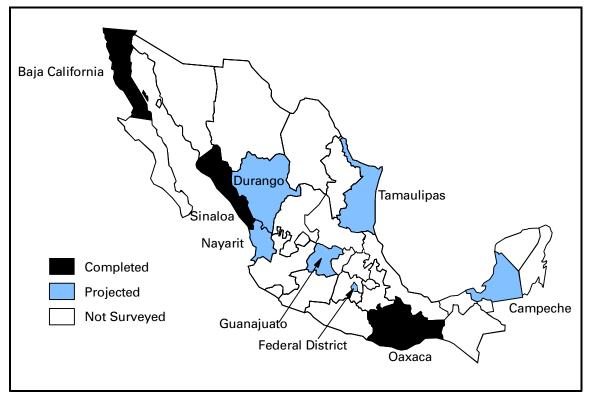
Population-Based Survey for Drug Resistance of Tuberculosis — Mexico, 1997

The World Health Organization (WHO) estimates that 90 million cases of tuberculosis (TB), resulting in 30 million deaths, will occur during the 1990s (1). To address this problem, WHO has recommended a comprehensive strategy of directly observed treatment, short-course (DOTS)* (2). Although DOTS results in cure rates of ≥80% (3), the worldwide emergence of strains of *Mycobacterium tuberculosis* (MTB) resistant to antimycobacterial agents threatens this strategy for TB control (2). In 1994, WHO and the International Union Against Tuberculosis and Lung Disease (IUATLD) proposed the establishment of a global surveillance program to monitor drug resistance (2). In 1997, the Secretary of Health of Mexico, in collaboration with CDC, developed and implemented a national survey of drug resistance for TB as part of the global project on TB drug resistance. This report describes study results for three states in Mexico (Baja California, Oaxaca, and Sinaloa) and presents the first population-based TB drug-resistance data available for that country.

For this study, the 31 states and Federal District of Mexico were categorized by reported TB incidence in 1994 into three strata (high, medium, and low incidence). Nine of these 32 areas were randomly chosen in proportion to the number of cases reported in each strata. Baja California (high), Sinaloa (high), and Oaxaca (medium) were selected as the first of the nine to participate in the survey (Figure 1). Cases were enrolled from two of the country's five major public-sector health-care agencies, the Secretaria de Salud Administracion (Secretary of Health) (SSA) and the Instituto Mexicano de Seguro Social (Mexican Institute of Social Security) (IMSS); these two agencies together provide health-care service to approximately 80% of the population and diagnose and manage 90% of reported TB cases. During January-April 1997, physicians, epidemiologists, and laboratory workers from these agencies in all three states received extensive training from SSA in conducting the survey. During April 1-October 31, physicians completed patient enrollment forms for all patients submitting at least one sputum sample for evaluation for pulmonary TB. All acid-fast bacilli (AFB) smear-positive samples were sent to the state laboratories for inoculation onto Lowenstein-Jensen media and were forwarded to the Instituto Nacional de Diagnostico y Referencia Epidemiologicos (National Diagnostic and Epidemiologic Reference Institute) (INDRE) in Mexico City for species identification and testing for drug susceptibility to isoniazid, rifampin, pyrazinamide, streptomycin, and ethambutol using the

^{*}DOTS consists of 1) committing to a sustainable national TB program; 2) detecting cases among symptomatic patients self-reporting to health services, using smear microscopy; 3) administering standardized short-course chemotherapy with direct observation of treatment; 4) establishing a regular drug supply of essential anti-TB drugs; and 5) establishing and maintaining a standardized recording and reporting system that allows assessment of treatment results.

FIGURE 1. States surveyed or projected to be surveyed for multidrug-resistant *Mycobacterium tuberculosis* — Mexico, 1997



radiometric method (4). The reference institute and CDC exchanged and tested 20 MTB isolates on two separate occasions for quality-control monitoring; there was a discrepancy in one drug for one isolate, for an accuracy rate of 97.5%.

In this analysis, resistance to one or more drugs was defined as resistance to isoniazid, rifampin, or pyrazinamide—the three drugs that constitute first-line treatment in Mexico. Resistance to one or more drugs was defined as primary for patients who had never taken anti-TB drugs and as acquired for patients reporting previous treatment with anti-TB drugs. Multidrug-resistant (MDR) TB was defined as resistance to at least isoniazid and rifampin (2). Primary resistance was considered to reflect infection with a resistant organism, and acquired resistance was considered to reflect the development of resistance during the course of previous therapy.

During the study period, 816 patients were officially reported with AFB smear-positive pulmonary TB: 351 from Baja California, 110 from Oaxaca, and 355 from Sinaloa (Table 1). Of these, 602 (74%) were enrolled in the study; MTB isolates were available for drug-susceptibility testing from 440 (73%) patients. Of the remaining specimens, 22% had no growth, 4% were contaminated, and 1% had nontuberculous mycobacteria. Of patients with MTB isolates, 24% had a history of prior TB treatment. The median age of patients was 36 years (range: 10–99 years); 69% were male. No difference was observed between patients with culture-positive and culture-negative isolates by age or prior history of TB.

Primary resistance to one or more of the three current first-line drugs used in Mexico was 12%; acquired resistance was 50% (Table 2). Levels for both primary and acquired drug resistance did not differ significantly by state or by patient age or sex. Levels of combined resistance (primary and acquired), which represent an approximation of the overall level of drug resistance to isoniazid, rifampin, pyrazinamide, ethambutol, or streptomycin the community, were 26% (113 of 440) for one or more of the five drugs, 18% (79 of 440) for isoniazid resistance, and 6% (28 of 440) for MDR TB.

TABLE 1. Number of acid-fast bacilli smear-positive pulmonary tuberculosis (TB) cases, number and percentage of enrolled patients with smear positive pulmonary TB, and number and percentage of isolates recovered from those patients — selected states, Mexico, April–October, 1997

	No. cases	Patients en	rolled in study	Isolates recovered†		
State	reported*	No.	(%)	No.	(%)	
Baja California	351	298	(85)	216	(72)	
Oaxaca	110	159	(>100)§	104	(65)	
Sinaloa	355	145	(41)	120	(83)	
Total	816	602	(74)	440	(73)	

^{*}Source: Secretary of Health of Mexico.

TABLE 2. Number and percentage of sputum-smear positive pulmonary tuberculosis (TB) patients with drug-resistant *Mycobacterium tuberculosis* isolates, by drug to which the isolate was resistant and by TB treatment history[†] — Baja California, Oaxaca, and Sinaloa states, Mexico, April–October 1997

	ΔII c	ases	Pr	evious t	reatmei			
	(n=440)		No [†] (n=308)		Yes§ (n=99)		Prevalence	
Drug	No.	(%)	No.	(%)	No.	(%)	rate ratio¶	(95% CI**)
Isoniazid	79	(18)	35	(11)	41	(41)	3.6	(2.4- 5.4)
Rifampin	37	(8)	7	(2)	27	(27)	12.0	(5.4-26.7)
Pyrazinamide	23	(5)	4	(1)	18	(18)	14.0	(4.9-40.4)
Ethambutol	24	(6)	9	(3)	14	(14)	4.8	(2.1-10.8)
Streptomycin	66	(15)	34	(11)	28	(28)	2.6	(1.6- 4.0)
Any first-line drug ^{††}	90	(21)	38	(12)	49	(50)	4.0	(2.8- 5.7)
Multidrug-resistant ^{§§}	28	(6)	5	(2)	20	(20)	12.4	(4.8–32.3)
Five drugs¶	10	(2)	2	(1)	8	(8)	12.4	(2.7–57.6)

^{*}Patients with culture-positive TB for whom complete data about treatment history was available; n=407.

[†] Mycobacterium tuberculosis isolates available for drug susceptibility testing.

[§]Because of underreporting and reporting delays, some patients enrolled in the study may not have been officially reported.

[†]Isolates from persons who reported no history of TB treatment are considered to have primary drug resistance.

[§]Isolates from persons who reported a history of previous treatment with anti-TB drugs are considered to have acquired drug resistance.

[¶]Rate in the previously treated group divided by the rate in the previously untreated group.

^{**} Confidence interval for the prevalence rate ratio.

^{††}Resistance to Isoniazid, rifampin, or pyrazinamide.

^{§§}Resistance to at least isoniazid and rifampin.

[¶]Resistance to Isoniazid, rifampin, pyrazinamide, ethambutol, and streptomycin.

Patients with acquired resistance were significantly more likely than patients with primary resistance to have resistance to one or more of the three first-line drugs (prevalence rate ratio [PRR]=4.0; 95% confidence interval [CI]=2.8–5.7), to have isoniazid resistance (PRR=3.6; 95% CI=2.5–5.4), and to have MDR TB (PRR=12.4; 95% CI=4.8–32.3).

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Editorial Note: This is the first population-based study of TB drug resistance from Mexico. Compared with results from 35 countries participating in the WHO/IUATLD global project on TB drug-resistance surveillance during 1994–1997, Mexico would have had the ninth highest level for primary resistance to at least one of the four first-line drugs (isoniazid, rifampin, ethambutol, and/or streptomycin) at 18% (pyrazinamide resistance was not evaluated). The United States ranked 14th with a level of 12% (2).

In 1996, 8% of TB cases in the United States occurred in persons born in Mexico (5). The 1993–1996 U.S. surveillance data about persons with TB who were born in Mexico and the findings from the survey of persons born in Mexico described in this report indicate similar rates among patients for primary isoniazid resistance (9% and 11%, respectively) and primary MDR TB (2% and 2%, respectively) (6).

The findings in this report are subject to at least three limitations. First, although surveillance for TB improved in the three surveyed states during the study period, the ability to assess data representativeness is limited by underreporting and notification delays. For example, the study in Oaxaca enrolled more patients than the number of persons officially reported as having smear-positive pulmonary TB. Second, 26% of the persons reported to the SSA with AFB smear-positive TB were not enrolled in the study, and 27% of the samples submitted could not be cultured. However, patients with positive cultures did not differ significantly from those with negative cultures by age or prior treatment history. Third, findings presented here are from only three of 31 states in Mexico and the Federal District; although the states are geographically dispersed, they may not be representative of the nation.

The findings of this survey have lead to improved TB control in Baja California, Oaxaca, and Sinaloa. All three state laboratories now have implemented the capacity to culture for MTB. Although smears rather than cultures are recommended by WHO as the basis of initial TB diagnosis in countries with limited resources, the newly developed culture capacity in the three states will be useful in surveillance efforts and in the management of cases not responding to routinely recommended treatment regimens.

In part as a result of this survey, the Secretary of Health of Mexico, in an effort to limit increases in drug resistance, is planning to initiate a four-drug treatment regimen by adding ethambutol to the current three-drug regimen. Four-drug regimens are recommended by CDC and the American Thoracic Society for communities with primary isoniazid resistance of ≥4% (7). A second action to limit drug resistance that is being implemented by the Secretary of Health is to expand the DOTS program to the entire country. In addition to preventing the development of drug resistance, national strategies that are feasible in Mexico are needed to treat patients with MDR TB. As changes are made in the TB program, trends in MTB drug resistance will need to be monitored by implementing ongoing surveillance or performing periodic surveys. Further collaborative international efforts will be needed to improve TB control in the United States and Mexico.

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Notice to Readers

Satellite Broadcast on Antimicrobial Use and Resistance

Antimicrobial Use and Resistance: Solutions to the Problem, a live, interactive satellite broadcast, will be held Thursday, August 20, 1998, from 9 a.m. to 11:30 a.m. eastern daylight time (EDT) with a repeat broadcast from 1 p.m. to 3:30 p.m. EDT. Cosponsors are CDC, the National Foundation for Infectious Diseases, in collaboration with the Association for Professionals in Infection Control and Epidemiology and the Society for Healthcare Epidemiology of America.

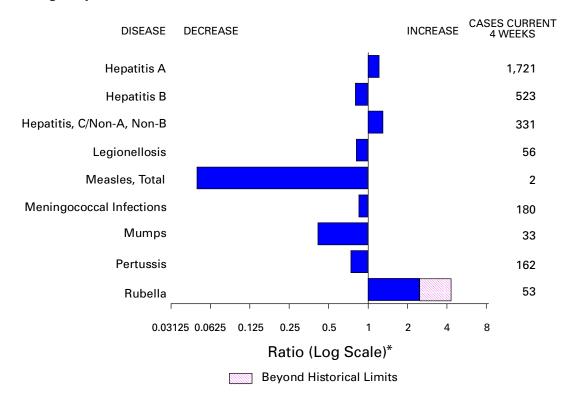
This broadcast will provide an overview of the increasing problem and emergence of antimicrobial resistant pathogens and will describe methods for the surveillance of antimicrobial resistance and assessment of antimicrobial use. Participants also will learn various strategies to improve antimicrobial use and prevent and control the spread of antimicrobial resistant pathogens. Continuing education credits will be awarded for various professions based on 2.5 hours of instruction.

Notice to Readers — Continued

This course is designed for physicians, nurses, infection-control professionals, pharmacists, laboratorians, hospital administrators, and others involved in the prevention and control of antimicrobial resistant pathogens.

Registration information is available through CDC's fax information system, telephone (888) 232-3299; request document number 130018.

FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending May 9, 1998, 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 May 9, 1998 (18th Week)

	Cum. 1998		Cum. 1998
Anthrax Brucellosis Cholera Congenital rubella syndrome Cryptosporidiosis* Diphtheria Encephalitis: California* eastern equine* St. Louis* western equine* Hansen Disease Hantavirus pulmonary syndrome* Hemolytic uremic syndrome, post-diarrheal* HIV infection, pediatric*	- 7 - 1 607 - - - - 43 2 8 8	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	13 - 24 829 25 50 5 52 2 102

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

Updated monthly to the Division of HIV/AIDS Prevention–Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP), last update April 26, 1998.

One suspected case of polio with onset in 1998 has also been reported to date.

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

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending May 9, 1998, and May 3, 1997 (18th Week)

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Penperfusing Area 1998 1997 1998 1998 1998 1997 1998 1998 1997 1998 1997 1998 1997 1998 1997 1998 1998 1998 1997 1998 1		All	DS	Chlai	nydia			Gono	rrhea		
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Nev. 87 144 253 1,046 4 3 92 290 12 4 PACIFIC 2,223 3,232 25,584 27,000 59 34 5,826 6,343 274 176 Wash. 165 240 3,918 3,184 15 11 660 708 8 9 Oreg. 64 128 1,955 1,635 18 17 269 248 2 2 2 Calif. 1,947 2,822 18,385 21,111 26 3 4,661 5,067 222 111 Alaska 11 18 690 491 - - 107 159 1 - - Hawaii 36 24 636 579 N 3 129 161 41 54 Guam - 2 8 151 N - 2 20 - - - PR. 20 -	Ariz.	200	157	3,315	3,464	N	5	1,213	1,089	1	11
PACIFIC 2,223 3,232 25,584 27,000 59 34 5,826 6,343 274 176 Wash. 165 240 3,918 3,184 15 11 660 708 8 9 Oreg. 64 128 1,955 1,635 18 17 269 248 2 2 Calif. 1,947 2,822 18,385 21,111 26 3 4,661 5,067 222 111 Alaska 11 18 690 491 - - 107 159 1 - Hawaii 36 24 636 579 N 3 129 161 41 54 Guam - 2 8 151 N - 2 20 - - PR. 666 517 U U - U 144 229 - 32 VI. 15		45 87									
Wash. 165 240 3,918 3,184 15 11 660 708 8 9 Oreg. 64 128 1,955 1,635 18 17 269 248 2 2 Calif. 1,947 2,822 18,385 21,111 26 3 4,661 5,067 222 111 Alaska 11 18 690 491 - - 107 159 1 - Hawaii 36 24 636 579 N 3 129 161 41 54 Guam - 2 8 151 N - 2 20 - - PR. 666 517 U U - U 144 229 - 32 VI. 15 28 N N N U - - - - - - Amer. Samoa											
Caliř. 1,947 2,822 18,385 21,111 26 3 4,661 5,067 222 111 Alaska 11 18 690 491 - - 107 159 1 - Hawaii 36 24 636 579 N 3 129 161 41 54 Guam - 2 8 151 N - 2 20 - - PR. 666 517 U U - U 144 229 - 32 VI. 15 28 N N N U - - - - - Amer. Samoa - - - - - N U - - - - - -	Wash.	165	240	3,918	3,184	15	11	660	708	8	9
Alaska 11 18 690 491 - - 107 159 1 - Hawaii 36 24 636 579 N 3 129 161 41 54 Guam - 2 8 151 N - 2 20 - - PR. 666 517 U U - U 144 229 - 32 VI. 15 28 N N N U - - - - Amer. Samoa - - - - N U - - - -											
Guam - 2 8 151 N - 2 20 - - PR. 666 517 U U - U 144 229 - 32 VI. 15 28 N N N U - - - - - Amer. Samoa - - - - N U - - - -	Alaska	· 11	18	690	491	-	-	107	159	1	-
P.R. 666 517 U U - U 144 229 - 32 V.I. 15 28 N N N N U Amer. Samoa N U		36								41	54
V.I. 15 28 N N N U Amer. Samoa N U		-								-	32
	V.I.	15				N	U	-			-
Calvillation IN IN IN U / 1.5 - /	Amer. Samoa C.N.M.I.	-	-	- N	- N	N N	U U	- 7	13	-	2

N: Not notifiable

U: Unavailable

-: no reported cases

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

^{*}Updated monthly to the Division of HIV/AIDS Prevention—Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention, last update April 26, 1998.

†National Electronic Telecommunications System for Surveillance.

§Public Health Laboratory Information System.

TABLE II. (Cont'd.) Provisional cases of selected notifiable diseases, United States, weeks ending May 9, 1998, and May 3, 1997 (18th Week)

	Legionellosis		Lyı Dise		Ma	laria		hilis Secondary)	Tubero	culosis	Rabies, Animal
Reporting Area	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1997	Cum. 1998*	Cum. 1997	Cum. 1998
UNITED STATES	353	295	1,232	1,024	350	442	2,353	3,037	1,955	5,333	2,350
NEW ENGLAND	20	24	252	185	17	19	27	59	90	128	466
Maine N.H.	1 2	1 3	7	3 4	1 3	1 2	1 1	-	U 2	11 1	78 33
Vt. Mass.	1 6	3 10	2 69	2 41	- 11	1 13	1 19	32	1 71	66	26 141
R.I. Conn.	4 6	3 4	25 149	32 103	2	2	- 5	- 27	16 U	7 43	30 158
MID. ATLANTIC	72	50	768	671	93	117	84	154	178	977	524
Upstate N.Y. N.Y. City	24 10	12 2	470 -	80 52	27 41	19 67	9 19	17 29	U U	125 509	366 U
N.J. Pa.	3 35	5 31	55 243	166 373	16 9	21 10	18 3 8	71 37	178 U	201 142	67 91
E.N. CENTRAL	116	116	23	13	24	45	366	268	148	550	16
Ohio Ind.	52 16	56 15	22 1	6 4	2 1	4 4	60 54	88 62	5 U	106 47	15 -
III. Mich.	12 23	5 29	-	1 2	6 14	20 14	160 72	24 35	143 U	281 77	-
Wis.	13	11	U	U	1	3	20	59	Ü	39	1
W.N. CENTRAL Minn.	27 3	22 1	10 3	9 7	20 8	9 4	56 -	65 13	65 U	157 43	213 40
lowa Mo.	2 10	3 2	6	- 1	2 7	2 2	45	3 33	U 52	20 59	43 12
N. Dak. S. Dak.	-	2 1	-	-	1	-	-	-	U 7	2 2	42 33
Nebr. Kans.	9 3	9 4	- 1	1	2	1	4 7	- 16	, 3 U	4 27	1 42
S. ATLANTIC	46	35	122	101	86	84	994	1,201	332	882	772
Del. Md.	6 9	5 10	94	20 67	1 31	2 27	9 228	8 350	90	9 92	17 188
D.C. Va.	3 4	2	4	5	5 9	6 21	30 71	44 99	39 53	28 111	221
W. Va.	N	N	4	-	-	-	-	3	20	18	32
N.C. S.C.	4 4	5 2	1	2 1	7	5 5	287 120	245 128	130 U	117 87	136 53
Ga. Fla.	- 15	7	2 12	1 5	13 17	12 6	171 78	221 103	U U	152 268	45 80
E.S. CENTRAL	11	10	15 2	23	9	12	376	670	- U	406	95 14
Ky. Tenn.	8 3	4	7	2 8	1 5	3	43 192	60 272	Ū	60 136	14 60
Ala. Miss.	-	2 4	6	2 11	3 -	3 3	80 61	166 172	U U	137 73	21 -
W.S. CENTRAL	8	5	3 2	2	9	7 1	253	432	38	770	68
Ark. La.	-	1	-	1	3	4	46 102	55 138	38	64 48	1
Okla. Tex.	3 5	1 3	1	1	1 5	2	16 89	41 198	U U	58 600	67 -
MOUNTAIN Mont.	20 1	18 1	1	2	18	27 2	80	64	93 2	158 2	52 16
ldaho	-	1	-	-	1	-	-	-	3	4	32
Wyo. Colo.	1 4	1	-	-	6	1 12	4	2	1 U	2 35	32 -
N. Mex. Ariz.	2 3	1 4	-	1	6 4	4 3	10 61	54	7 59	6 66	4
Utah Nev.	8 1	4 2	- 1	- 1	1 -	1 4	3 2	2 6	21 U	6 37	-
PACIFIC	33	15	38	18	74	122	117	124	1,011	1,305	144
Wash. Oreg.	3	3	1	7	6 7	4 7	6 2	6	U U	103 44	- -
Calif. Alaska	30	11 -	34	11 -	60 -	108 2	109 -	113 1	942 15	1,051 31	131 13
Hawaii	-	1	-	-	1	1	-	1	54	76	-
Guam P.R.	-	-	-	-	-	3	81	3 73	46	13 -	23
V.I. Amer. Samoa	-	-	-	-	-	-	-	-	-	-	-
C.N.M.I.	-	-	-	-	-	-	1	5	8	-	-

N: Not notifiable U: Unavailable -: no reported cases

^{*}Additional information about areas displaying "U" for cumulative 1998 Tuberculosis cases can be found in Notice to Readers, MMWR Vol. 47, No. 2, p. 39.

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending May 9, 1998, and May 3, 1997 (18th Week)

	H. influ	ienzae,	Н	I	•	Meas	les (Rube	ola)				
		sive	A			В	Indi	genous		ported [†]		tal
Reporting Area	Cum. 1998*	Cum. 1997	Cum. 1998	Cum. 1997	Cum. 1998	Cum. 1997	1998	Cum. 1998	1998	Cum. 1998	Cum. 1998	Cum. 1997
UNITED STATES	392	422	7,034	9,276	2,526	3,014	-	3	-	10	13	46
NEW ENGLAND	23	23	97	233	29	64	-	-	-	1	1	1
Maine N.H.	2 1	3 3	10 6	26 12	- 7	3 5	-	-	-	-	-	-
Vt.	2	-	7	5	-	1	-	-	-	-	-	-
Mass. R.I.	16 2	15 1	22 8	121 17	11 11	34 6	-	-	-	1 -	1 -	1 -
Conn.	-	1	44	52	-	15	-	-	-	-	-	-
MID. ATLANTIC Upstate N.Y.	59 24	53 3	418 123	852 93	359 106	470 79	-	-	-	1	1	11 4
N.Y. City	10	18	118	404	96	194	-	-	-	-	-	5
N.J. Pa.	23 2	20 12	84 93	132 223	60 97	90 107	-	-	-	1	1	1 1
E.N. CENTRAL	53	66	841	1,176	249	583	-	-	_	2	2	6
Ohio	27	34 5	122	158	26	34 39	Ū	-	Ū	-	- 1	-
Ind. III.	9 16	18	66 123	111 295	20 38	115	-	-	-	1 -	-	5
Mich. Wis.	- 1	9	475 55	532 80	157 8	179 216	-	-	-	1	1	1
W.N. CENTRAL	29	23	635	661	114	193	_	_	_	-	-	10
Minn.	17	14	28	47	11	9	-	-	-	-	-	1
lowa Mo.	1 7	2 3	307 241	86 381	17 68	12 150	-	-	-	-	-	- 1
N. Dak.	-	2	2 3	7 6	2 1	1	-	-	-	-	-	-
S. Dak. Nebr.	-	1	13	22	5	7	-	-	-	-	-	8 -
Kans.	4	1	41	112	10	14	-	-	-	-	-	-
S. ATLANTIC Del.	89	78 -	622 1	472 10	360	387 3	-	1 -	-	5 1	6 1	2
Md.	24	31	132	112	48	65	-	-	-	1	1	1
D.C. Va.	11	6	24 95	13 64	6 33	18 41	-	-	-	2	2	1 -
W. Va. N.C.	3 11	3 12	36	5 6 8	2 77	6 86	-	-	-	-	-	-
S.C.	1	3	12	42	-	37	-	-	-	-	-	-
Ga. Fla.	18 21	16 7	116 206	43 115	59 135	38 93	U	1	U	1	1 1	-
E.S. CENTRAL	21	25	132	254	156	219	-	-	_	_	-	1
Ky. Tenn.	3 13	4 15	7 92	28 157	16 114	13 135	-	-	-	-	-	-
Ala.	5	6	33	37	26	28	-	-	-	-	-	1
Miss.	-	-	-	32	-	43	U	-	U	-	-	-
W.S. CENTRAL Ark.	23	19 1	1,200 17	1,411 91	383 21	190 19	Ū	-	Ū	-	-	4
La.	11	3	13	75	9	41	-	-	-	-	-	-
Okla. Tex.	11 1	13 2	193 977	590 655	25 328	10 120	-	-	-	-	-	4
MOUNTAIN	56	42	1,146	1,454	295	304	-	-	-	-	-	1
Mont. Idaho	-	-	19 85	43 63	3 14	4 8	-	-	-	-	-	-
Wyo.	-	1	24	16	7	8	-	-	-	-	-	-
Colo. N. Mex.	11 4	5 2	91 65	170 101	36 119	60 106	-	-	-	-	-	-
Ariz. Utah	31 4	12 3	725 74	662 279	71 23	64 36	-	-	-	-	-	1
Nev.	6	19	63	120	22	18	Ū	-	Ū	-	-	-
PACIFIC	39	93	1,943	2,763	581	604	-	2	-	1	3	10
Wash. Oreg.	2 24	1 17	373 140	195 134	46 46	19 44	-	-	-	-	-	-
Calif.	10	72	1,402	2,363	480 4	526	-	2	-	1	3	7
Alaska Hawaii	1 2	1 2	9 19	15 56	4 5	10 5	-	-	-	-	-	3
Guam	-	-	-	-	_ =	1	U	-	U	-	-	-
P.R. V.I.	2	-	12 -	122	226	462 -	Ū	-	- U	-	-	-
Amer. Samoa	-	-	-	-	-	-	Ū	-	U	-	-	-
C.N.M.I.	-	4	-	1	7	20	U	-	U	-	-	1

N: Not notifiable

U: Unavailable

^{-:} no reported cases

 $^{^{*}\!\!}$ Of 93 cases among children aged <5 years, serotype was reported for 47 and of those, 23 were type b.

[†]For imported measles, cases include only those resulting from importation from other countries.

TABLE III. (Cont'd.) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending May 9, 1998, and May 3, 1997 (18th Week)

	Mening Dise	ococcal ease		Mumps Pertussis						Rubella	
Reporting Area	Cum. 1998	Cum. 1997	1998	Cum. 1998	Cum. 1997	1998	Cum. 1998	Cum. 1997	1998	Cum. 1998	Cum. 1997
UNITED STATES	1,111	1,509	10	161	225	40	1,256	1,881	22	191	23
NEW ENGLAND	59	92	-	-	7	5	220	451	3	30	-
Maine N.H.	4 1	8 9	-	-	-	-	5 19	6 54	-	-	-
Vt.	1	2	-	-	-	-	23	150	-	-	-
Mass. R.I.	27 3	52 5	-	-	2 4	5	167	224 12	-	4	-
Conn.	23	16	-	-	1	-	6	5	3	26	-
MID. ATLANTIC	114	147	-	6	29	4	152	171	8	88	9
Upstate N.Y. N.Y. City	29 13	33 25	-	3	4 1	4	95 -	59 43	8	88	1 8
N.J.	33	30	-	-	4	-	-	9	-	-	-
Pa. E.N. CENTRAL	39 147	59	-	3	20 30	-	57 139	60 104	-	-	-
Ohio	147 58	219 81	-	24 11	30 9	-	138 53	194 57	-	-	3
Ind. III.	25 32	22 74	U	2 1	4 9	U	40 10	19 26	U	-	-
Mich.	15	20	-	10	7	-	18	28	-	-	-
Wis.	17	22	-	-	1	-	17	64	-	-	3
W.N. CENTRAL Minn.	94 16	112 17	-	16 9	7 3	1	97 58	104 63	-	2	-
Iowa	14	22	-	5	3	1	17	7	-	-	-
Mo. N. Dak.	38	56	-	1 1	-	-	9	17 2	-	1	-
S. Dak.	6	3	-	-	-	-	4	1	-	-	-
Nebr. Kans.	4 16	4 10	-	-	1 -	-	3 6	2 12	-	1	-
S. ATLANTIC	192	247	7	29	33	3	97	164	_	5	1
Del.	1	4	-	-	-	-	-	-	-	-	-
Md. D.C.	18 -	26 5	-	-	4	1 -	19 1	68 2	-	-	-
Va. W. Va.	19 5	23 9	-	4	4	-	6 1	18 3	-	-	1
N.C.	25	41	-	6	6	-	40	35	-	3	-
S.C. Ga.	30 40	34 47	1 U	4 1	7 4	Ū	10 1	8 2	Ū	1	-
Fla.	54	58	6	14	8	2	19	28	-	1	-
E.S. CENTRAL	79	105	-	-	13	-	34	37	-	-	-
Ky. Tenn.	13 35	27 33	-	-	2 3	-	15 9	10 12	-	-	-
Ala.	31	29		-	4	-	10	9	-	-	-
Miss.	-	16	U	-	4	U	-	6	U	-	-
W.S. CENTRAL Ark.	123 14	145 22	Ū	22	27 -	5 U	68 8	35 2	11 U	51 -	1 -
La.	24	28	-	1	7	-	-	7	-	-	-
Okla. Tex.	21 64	14 81	-	21	20	5	6 54	5 21	11	51	1
MOUNTAIN	68	88	-	14	11	7	279	453	-	5	-
Mont. Idaho	2 3	5 5	-	- 1	2	2	1	2 298	-	-	-
Wyo.	3	-	-	1	1	-	131 7	3	-	-	-
Colo. N. Mex.	16 12	27 15	N	2 N	2 N	- 5	43 55	114 21	-	- 1	-
Ariz.	22	16	-	4	-	-	22	9	-	1	-
Utah Nev.	7 3	11 9	Ū	1 5	3 3	Ū	13 7	2 4	Ū	2 1	-
PACIFIC	235	354	3	50	68	15	, 171	272	-	10	9
Wash.	26	43	-	4	5	15	101	129	-	8	-
Oreg. Calif.	45 159	70 238	N 3	N 32	N 49	-	8 58	10 127	-	1	4
Alaska	1	1	-	2	4	-	-	2	-	-	-
Hawaii	4	2	- U	12	10	-	4	4	- U	1	5
Guam P.R.	2	1 6	-	2	1 4	U -	2	-	-	-	-
V.I. Amer. Samoa	-	-	U U	-	-	U U	-	-	U U	-	-
C.N.M.I.	-	-	Ü	-	1	Ü	-	-	Ü	-	-

C.N.M.I.
N: Not notifiable

U: Unavailable

-: no reported cases

TABLE IV. Deaths in 122 U.S. cities,* week ending May 9, 1998 (18th Week)

-						, 5,	.50	o (10tii vveek)						-	
Dan author Assa		All Cau	ises, By	/ Age (Y	ears)		P&I [†]	Dan autin a Ausa		All Cau	ses, 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.	30 U 9 51	347 83 19 18 20 39 16 8 22 20 U 6 34	30 7 3 2 6 2 3 2 5 U	34 62 - 61 - 3U33	8 2 1 - - 1 - - 1 U	8 4 - 1 - - 1 U	43 14 2 2 3 2 2 U 1 6	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.	970 U 185 87 129 107 50 59 35 148 100	625 U 113 58 78 70 34 35 29 108 63 3	211 U 44 20 32 20 11 14 20 3 23 18 6	80 U 21 6 14 13 1 6 3	25 U 4 2 2 2 3 1 3 5 3	27 U 3 1 3 2 1 4 1 - 4 8	47 U 17 9 - 4 - 4 7 2
Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Erie, Pa. Jersey City, N.J. New York City, N.Y. Newark, N.J. Paterson, N.J. Philadelphia, Pa. Fleading, Pa. Rochester, N.Y. Schenectady, N.Y. Scranton, Pa. Syracuse, N.Y. Trenton, N.J. Utica, N.Y. Yonkers, N.Y.	32 58 2,123 43 52 U 152 51 44 1,109 28 299 50 22 134 26 32 83 28 14 U	23 39 1,492 32 17 40 U 12 28 762 31 15 212 31 18 100 19 24 68 19	10 410 11 4 5 U 4 4 5 225 17 7 62 22 4 11 11 7	2 8 160 2 2 2 8 8 8 13 5 16 4 2 9 1 4 3 1	1 29 1 U 1 6 2 1 U 1 U U 1 U U	1 - 32 - 4 U - 3 18 1 - 3 1 - 1 - U	36 120 12 U ' 2 ' 51 4 ' 18 54 8 2 1 7 4 ' 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.	83 48 U 71 43 128 1,231 82 45	403 113 300 566 366 U 588 311 79 824 466 31 389 40 71 152 666 79 127 U 85	113 30 14 21 8 U 9 8 23 249 24 8 7 51 9 10 44 16 29 37 U	43 17 25 1 1 15 88 7 2 2 11 8 7 17 6 15 11 U 2	19 5 3 1 3 1 5 3 2 3 5 1 5 7 5 3 5 5 0 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	13 4 	36 14 2 1 3 U 2 2 12 69 2 - 3 4 3 11 7 6 - 2 0 U 13
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. 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, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans.	1,863 49 40 425 112 140 154 33 192 43 67 77 70 190 44 123 U 50 0 45 U 77 646 60 22 16	1,257 30 34 261 74 93 100 244 118 31 500 4 60 127 33 88 8 U 36 31 U 63 435 39 177 12	359 7 3 98 226 29 6 41 8 11 4 40 7 24 U 11 10 12 104 12 2 1	140 2 1 40 8 11 15 20 4 1 17 3 4 U 3 7 U 1 45 5 1 1	47 2 1 15 3 4 1 -7 -3 3 1 3 4 -2 0 -1 0 -2 1 1 0 -1 1 0 -1 1 0 -1 1 0 -1 1 0 -1 1 0 -1 1 0 -1 1 0 -1 1 0 -1 1 0 -1 1 1 1	59 8 1 10 5 6 9 1 1 6 2 2 2 2 2 1 5 5 0 0 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	124 6 44 16 3 11 1 6 2 9 4 11 U 5 2 U 4 39 3 2 2	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 Jose, Calif. San Jose, Calif. Santa Cruz, Calif. Santa Cruz, Calif. Santa Cruz, Calif.	104 175 23 156 32 93 121 1,200 0 0 74 60 0 74 67 0 31 U 143	605 59 25 36 72 115 18 101 24 61 94 843 843 49 U 55 U 99 97 73 135 26 106	156 14 7 7 18 42 19 5 17 16 227 2 10 U 12 U 5 U 23 31 24 44 44 43 36	79 5 4 9 10 14 3 16 3 8 7 7 9 - 5 U 7 4 U - U 7 9 12 20 1 6	22 1 - 5 2 2 1 1 1 4 - 5 3 3 28 - 1 1 U - 2 2 U 1 U 5 4 4 8 - 2	16 1 2 2 2 3 3 - 6 2 2 U 1 U 6 2 2 1 6 4	52 3 2 2 9 111 4 8 3 5 5 5 112 - - - - - - - - - - - - - - - - - -
Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	105 44 113 66 86 92 39	70 28 80 48 45 69 27	7 20 10 22 12	6 7 8 3 7 3 4	6 1 2 2 5 2 1	1 1 3 7 6 1	3 7 8 8 3 2	Spokane, Wash. Tacoma, Wash. TOTAL	58 97	45 76 6,831	10 14	2 6 748	1 - 234	1 240	5 4 642

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.

Quarterly Immunization Table

To track progress toward achieving the goals of the Childhood Immunization Initiative (CII), CDC publishes quarterly a tabular summary of the number of cases of nationally notifiable diseases preventable by routine childhood vaccination reported during the previous quarter and year-to-date (provisional data). In addition, the table compares provisional data with data for the previous year and highlights the number of reported cases among children aged <5 years, who are the primary focus of CII. Data in the table are reported through the National Electronic Telecommunications System for Surveillance (NETSS).

Number of reported cases of diseases preventable by routine childhood vaccination — United States, January–March 1998 and January–March 1997*

	No. cases, January–March	Total (January	cases ⁄-March	No. cases among children aged <5 years [†] January–March			
Disease	1998	1997	1998	1997	1998		
Congenital rubella							
syndrome	0	2	0	2	0		
Diphtheria	0	1	0	0	0		
Haemophilus influenzae§	255	283	255	56	55		
Hepatitis B¶	1497	1941	1497	52	19		
Measles	7	17	7	9	5		
Mumps	98	126	98	25	16		
Pertussis	807	1101	807	438	332		
Poliomyelitis, paralytic**	0	0	0	0	0		
Rubella	98	9	98	3	4		
Tetanus	2	8	2	0	1		

^{*}Data for 1997 and 1998 are provisional.

[†]For 1997 and 1998, age data were available for ≥97% cases.

[§]Invasive disease; *H. influenzae* serotype is not routinely reported to the National Notifiable Diseases Surveillance System. Of 55 cases among children aged <5 years, serotype was reported for 22 cases, and of those, 12 were type b, the only serotype of *H. influenzae* preventable by vaccination.

Because most hepatitis B virus infections among infants and children aged <5 years are asymptomatic (although likely to become chronic), acute disease surveillance does not reflect the incidence of this problem in this age group or the effectiveness of hepatitis B vaccination in infants.

^{**}Two cases with onset in 1997 have been confirmed; two suspected cases are under investigation, of which one is in a child aged <5 years. One suspected case in a child aged <5 years with onset in 1998 is also under investigation.

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☆U.S. Government Printing Office: 1998-633-228/67078 Region IV