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Adverse Events Associated with Ingestion of Gamma-Butyrolactone — Minnesota, New Mexico, and Texas, 1998–1999

Products containing gamma-butyrolactone (GBL)* are marketed for many claimed purposes, including to induce sleep, release growth hormone, enhance sexual activity and athletic performance, relieve depression, and prolong life. GBL is converted by the body into gamma-hydroxybutyrate (GHB), a drug banned outside of clinical trials approved by the Food and Drug Administration (FDA). Recognized manifestations of GHB toxicity include bradycardia, hypothermia, central nervous system depression, and uncontrolled movements (1). This report describes seven cases of GBL toxicity involving the product "Revivarant," which is labeled as containing 1.82 g of GBL per fluid ounce, reported from two hospital emergency departments (EDs) in Minnesota during October–December 1998 and summarizes an additional 34 cases of GBL toxicity reported to poison centers in New Mexico and Texas during October 1998–January 1999.

Minnesota

Patient 1. On November 26, 1998, a 24-year-old man vomited and had seizures shortly after drinking 3–4 oz of Revivarant. His behavior became unusual, and he alternated between extreme agitation and profound calm. Paramedics noted that his skin was warm, flushed, and profusely diaphoretic, and he had bradycardia (pulse as low as 45 beats per minute [bpm]). Systolic blood pressure was 110 mm Hg. Transcutaneous oxygen saturations (SpO₂) were 96% on room air, and blood glucose by fingerstick was 90 mg/dL. During transport to an ED, he had periods of combativeness lasting 30 to 60 seconds followed by coma lasting 1–3 minutes. In the ED, he was unconscious with spontaneous eye opening, a positive withdrawal reflex, and no speech (Glasgow Coma Scale of 7); rectal temperature was 94.8 F (34.9 C). A urine toxicology screen and blood ethanol test were negative. He was intubated and admitted to the intensive-care unit (ICU) with a diagnosis of toxic encephalopathy. During the next 7 hours, his heart rate increased from 42 to 116 bpm and he became more alert. He had no recollection of events except for having ingested Revivarant. He was discharged with normal mental status.

^{*}Also is known as dihydro-2(3H)-furanone; 4-butanolide; 2(3H)-furanone, dihydro; tetrahydro-2-furanone; and butyrolactone gamma.

Gamma-Butyrolactone — Continued

Patient 2. On December 12, 1998, a 46-year-old woman had a seizure and lost consciousness after drinking approximately 2.7 oz of Revivarant in conjunction with ethanol. Paramedics found her unconscious and in severe respiratory depression with a pulse of 54 bpm. Oxygen was administered by mask; she had an SpO₂ of 87%. On arrival in the ED, physical examination identified sinus bradycardia (54 bpm); temperature of 96.1 F (35.6 C); and miotic pupils. A serum ethanol level was 0.11%. She was admitted to the ICU, mechanically ventilated through the night, and awoke in improved condition the next morning; she was discharged with no memory of the events.

Patient 3. On November 8, 1998, a 31-year-old man drank approximately 1 oz of Revivarant, four beers, and a large sip of wine. Shortly thereafter, he gradually lost consciousness and subsequently fell. He regained consciousness but had involuntary muscle movements and episodes of confusion. Paramedics noted that he was ambulatory but confused. On physical examination in the ED, he was agitated, anxious, and unable to recall the preceding events. His shoulders twitched, and he had a small abrasion below his left eye. He had a pulse of 64 bpm and hypothermia (oral temperature of 95.2 F [35.1 C]). Breath ethanol level was 0.08%. He denied previous GBL use or illicit drug use. He recovered completely and was discharged.

Patients 4 and 5. On October 31, 1998, a 24-year-old man (patient 4) and a 26-year-old man (patient 5) each drank 10–13 oz of Revivarant while drinking alcohol at a bar. On leaving the bar, witnesses observed them fall and become unresponsive. On arrival at the ED, they alternated between somnolence and confusion. When awake, neither patient could consistently follow commands. Patient 4 had fecal incontinence. Vital signs for both patients were within normal limits. Breath ethanol levels were 0.09% (patient 4) and 0.15% (patient 5). Neither patient had a history of using medications or illicit drugs. After 2 hours of observation, the patients recovered but were unable to recall most of the evening's events.

Patients 6 and 7. On December 12, 1998, a 19-year-old woman (patient 6) and a 22-year-old woman (patient 7) were brought to an ED by friends because of vomiting and decreased levels of consciousness. These symptoms followed ingestion of Revivarant (2 oz by patient 6 and an unknown amount by patient 7). Patient 6 had drank one beer; patient 7 had had no ethanol. Vital signs were normal except for respiratory depression. On physical examination, patient 6 was lethargic and disoriented. Patient 7 exhibited intermittent periods of extreme agitation, necessitating chemical treatment and physical restraint, punctuated by moments of calm during which her attention focused on minor details. Mental changes for both patients resolved, and they were discharged approximately 4 hours after arrival.

New Mexico

From October 3, 1998, through January 29, 1999, the New Mexico Poison Center identified 14 cases of adverse events resulting in an ED visit among persons who had ingested GBL-containing products. Ten (71%) of the cases were reported in January. Patients' ages ranged from 14 to 36 years; nine were male. Products used included "Firewater" (11 cases), "Blue Nitro Vitality" (two), and "RenewTrient" (one). The approximate amount ingested ranged from 1 to 10 oz (mean: 3 oz). Five (36%) persons also had ingested ethanol and/or other drugs. Most of the patients were discharged from the ED within 13 hours of arrival; three were hospitalized. The most common

Gamma-Butyrolactone — Continued

symptoms and signs were nausea/vomiting (10 [71%]), obtundation (nine [64%]), bradycardia (seven [50%]), prolonged unconsciousness (six [43%]), syncope (six [43%]), seizures (four [29%]), confusion (four [29%]), combativeness (four [29%]), respiratory depression (three [21%]), amnesia (two [14%]), and euphoria (two [14%]). One person had cardiac arrest, one had respiratory arrest, and one had a motorvehicle crash associated with the effects resulting from use of a GBL-containing product. No deaths were reported.

Texas

From October 2, 1998, through January 24, 1999, Texas poison-control centers identified 20 adverse events resulting in ED visits among persons who had ingested GBL-containing products. Twelve (60%) of the cases were reported in January. Patients' ages ranged from 11 to 41 years; 13 were male. Products known to have been used included "RenewTrient" (six cases), "Revivarant" (four), "Revivarant-G" (two), and "Blue Nitro Vitality" (two). Ten persons also ingested ethanol and/or other drugs. Ten patients were admitted to the hospital from the ED. The most common symptoms and signs were obtundation (13 [65%]), prolonged unconsciousness (nine [45%]), respiratory depression (nine [45%]), anxiety/nervousness (seven [35%]), nausea/vomiting (six [30%]), confusion (six [30%]), tremors/twitching (four [20%]), tachycardia (three [15%]), and combativeness (three [15%]). One person had respiratory arrest; no deaths were reported.

Reported by: SW Smith, MD, AR Topliff, MD, M Danigelis, MD, DL Zvosec, PhD, LL Schrag, MD, SA Freiwald, MD, SR Gunn, MD, Dept of Emergency Medicine, Hennepin County Medical Center, Minneapolis; SC Setzer, Hennepin Regional Poison Center, Minneapolis; M Rock, MD, Methodist Hospital, St. Louis Park; MT Osterholm, PhD, State Epidemiologist, Minnesota Dept of Health. BE Benson, PharmD, New Mexico Poison Center, Albuquerque; J Padilla, MS, R Voorhees, MD, CM Sewell, DrPH, State Epidemiologist, New Mexico Dept of Health. L Williams, G Shepherd, North Texas Poison Center, Dallas; G Coody, PharmD, Div of Drugs and Medical Devices, Bur of Food and Drug Safety; DM Simpson, MD, State Epidemiologist, Texas Dept of Health. Center for Food Safety and Applied Nutrition, Food and Drug Administration. Environmental Hazards and Epidemiology Section, Health Studies Br, Div of Environmental Hazards and Health Effects, National Center for Environmental Health; Div of Applied Public Health Training, Epidemiology Program Office; and ElS officers, CDC.

Editorial Note: GBL is metabolized to GHB in the body, but because of better absorption GBL has greater bioavailability than GHB on an equimolar basis (2). Clinical effects of GHB appear to be dose-related and include reports of vomiting, hypotonia, tremors, seizures, aggression, impairment of judgment, coma, respiratory depression, hypothermia, and bradycardia (1). GHB mixed with ethanol acts synergistically to produce central nervous system and respiratory depression (3). Symptoms usually resolve with supportive care within 2–96 hours (4). Death occurring when GHB was the sole intoxicant also has been reported (5). Toxic effects of GBL would be expected to be similar or identical to those of GHB, but previous clinical experience is limited (6,7). There is no antidote for GHB; treatment consists of supportive therapy until symptoms of toxicity subside. A withdrawal syndrome, which can include insomnia, tremor, and anxiety, has been reported following discontinuance of GHB in chronic, high-dose users (8).

GBL is an industrial and household solvent of acrylate polymers, and unintentional poisonings have been reported (6,9). It also is marketed as a dietary supplement at health food stores and on the World-Wide Web under several trade names. Although

Gamma-Butyrolactone — Continued

labeled as dietary supplements, GBL-containing products are illegally marketed, unapproved new drugs that have been involved in at least 55 reports of adverse events, including one death (10). On January 21, 1999, FDA asked manufacturers to recall their GBL-containing products and warned consumers through press releases to avoid taking these products (10). Public education efforts should inform consumers that FDA review procedures for drugs are different than those used for dietary supplements. Consumers should be alert to the potential dangers of these products and understand that terms such as "natural" do not necessarily imply safety. Physicians should counsel patients about these products and be prepared to recognize and treat the toxic reactions that some might produce. Chronic GBL users should be monitored for withdrawal symptoms when discontinuing use of the product. Depending on the severity of the withdrawal symptoms, medical intervention may be required. Physicians are encouraged to report serious adverse events associated with these products to FDA's MedWatch program, telephone (800) 332-1088.

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Decline in Cigarette Consumption Following Implementation of a Comprehensive Tobacco Prevention and Education Program — Oregon, 1996–1998

In November 1996, residents of Oregon approved a ballot measure increasing the cigarette tax by 30¢ (to 68¢ per pack). The measure stipulated that 10% of the additional tax revenue be allocated to the Oregon Health Division (OHD) to develop and implement a tobacco-use prevention program. In 1997, OHD created Oregon's

Cigarette Consumption — Continued

Tobacco Prevention and Education Program (TPEP), a comprehensive, community-based program modeled on the successful tobacco-use prevention programs in California and Massachusetts (1,2). To assess the effects of the tax increase and TPEP in Oregon, OHD evaluated data on the number of packs of cigarettes taxed before (1993–1996) and after (1997–1998) the ballot initiative and implementation of the program. Oregon's results also were compared with national data. This report summarizes the results of the analysis, which indicate that consumption of cigarettes in Oregon declined substantially after implementation of the excise tax and TPEP and exceeded the national rate of decline.

OHD obtained data on the sale of Oregon cigarette tax stamps from the Oregon Department of Revenue for 1993–1998. OHD also obtained data on the proportion of revenue received at the old and new rates after the tax change (February 1997) to calculate the number of packs sold each month. Per capita consumption was calculated by dividing the number of packs sold by the total population of Oregon each year (3).

National comparison estimates were generated using data from the Tobacco Institute on state tax receipts for wholesale cigarette deliveries. Reliable figures were available through December 1997 (4). Data from Oregon and the other three states (Arizona, California, and Massachusetts) with tobacco-use prevention programs funded through state initiatives were excluded from the comparison estimates. National per capita consumption was calculated by dividing the total number of packs sold by the total population in the remaining 46 states and the District of Columbia (5). Calculations for Oregon for 1996–1998 represent the 1 year before and the 2 years after the tax increase.

From 1993 to 1996, taxable per capita consumption of cigarettes increased 2.2% in Oregon and decreased 0.6% in the 46 remaining states and the District of Columbia. In Oregon, from 1996 to 1998, taxable per capita cigarette consumption declined 11.3% (from 92 packs to 82 packs) (Figure 1). Despite a 2.7% increase in the state's population, 25 million fewer cigarette packs were sold in Oregon in 1998 than in 1996. In the United States during 1996–1997, per capita consumption declined 1.0% (from 93 packs to 92 packs).

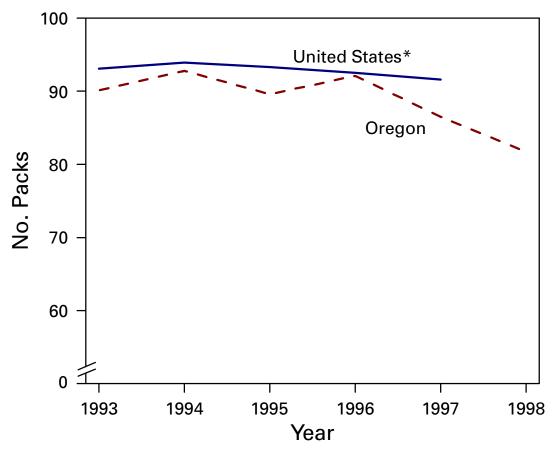
Reported by: B Pizacani, MPH, C Mosbaek, K Hedberg, MD, L Bley, PhD, M Stark, PhD, J Moore, PhD, D Fleming, MD, Oregon Health Div. Epidemiology Br, Office on Smoking and Health, National Center for Chronic Disease Prevention and Health Promotion, CDC.

Editorial Note: Two years after the implementation of a ballot measure to increase the excise tax on tobacco and initiate TPEP, per capita consumption has declined 11.3% in Oregon, or the equivalent of 200 cigarettes (10 packs) per capita. Elements of the program include community-based tobacco-use prevention coalitions in every county; a statewide public awareness and education campaign; comprehensive school-based programs; tribal tobacco-use prevention programs; multicultural outreach and education; a quitters' help line providing smoking cessation support; and projects evaluating new approaches to prevent or reduce tobacco use. TPEP has an annual budget of \$8.5 million, 93% of which is awarded in grants or contracts to external partners (e.g., county health departments, community-based agencies, tribal governments, and private-sector partners implementing the public awareness campaign).

Decreased consumption is probably a result of both the increase in the price of cigarettes and the tobacco-use prevention program. Price elasticity of demand,

Cigarette Consumption — Continued

FIGURE 1. Annual per capita sales of cigarettes — Oregon and United States, 1993–1998



^{*}Excluding Arizona, California, Massachusetts, and Oregon.

defined as the percentage change in demand for cigarettes resulting from a 1% change in price, is an estimated –0.4% (6). A 15.8% increase in the price of cigarettes (the amount of the price increase in Oregon, calculated in 1996 dollars) should result in a 6.3% decrease in cigarette consumption. The findings in this report are consistent with reports from other states with tobacco-use prevention programs and indicate that excise taxes in conjunction with prevention programs reduce cigarette consumption more than excise taxes alone (1,7).

Other factors that could account for the decrease in cigarette consumption in Oregon probably did not contribute to the decline. Smuggling or cross-border sales probably are insignificant because a large proportion of Oregon's population resides in Portland, near Washington, where cigarette prices are higher. Increased sales on Indian reservations in the state probably would not contribute to the decline because cigarettes sold on reservations are taxed, and tribes are reimbursed only for tobacco taxes paid by tribal members. Another possibility is that the observed downward trend for Oregon may reflect national declines. Although reliable national data are not available for 1998, it is unlikely that the decrease in Oregon reflects secular trends.

Cigarette Consumption — Continued

During 1990–1997, the annual rate of decline in consumption for all 50 states averaged only 1.4% (8).

Oregon's decrease in cigarette consumption also appears to be resulting in decreases in smoking prevalence. Preliminary data from the Behavioral Risk Factor Surveillance System for 1996–1998 indicate that prevalence of current smoking among adults in Oregon declined 6.4%, representing 35,000 fewer smokers. The decline in cigarette consumption in Oregon, California, and Massachusetts indicates that an adequately funded, comprehensive tobacco-control program can quickly and substantially reduce tobacco use.

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Neighborhood Safety and the Prevalence of Physical Inactivity — Selected States, 1996

Physical inactivity is an important risk factor for premature morbidity and mortality, especially among high-risk populations. Although health-promotion programs have targeted high-risk groups (i.e., older adults, women, and racial/ethnic minorities) (1), barriers exist that may affect their physical activity level (2). Identifying and reducing specific barriers (e.g., lack of knowledge of the health benefits of physical activity, limited access to facilities, low self-efficacy, and environmental issues [2–6]) are important for efforts designed to increase physical activity. Concerns about neighborhood safety may be a barrier to physical activity (2,3). To characterize the association between neighborhood safety and physical inactivity, CDC analyzed data from the 1996 Behavioral Risk Factor Surveillance System (BRFSS) in Maryland, Montana, Ohio, Pennsylvania, and Virginia. This report summarizes the results of this analysis, which indicate that persons who perceived their neighborhood to be unsafe were more likely to be physically inactive.

The BRFSS is a population-based, random-digit–dialed telephone survey of the civilian, noninstitutionalized U.S. population aged ≥18 years. In 1996, data on physical activity were analyzed for 12,767 persons (5320 men and 7447 women) who

Physical Inactivity — Continued

responded to the Social Context Module included in the 1996 surveys in Maryland, Montana, Ohio, Pennsylvania, and Virginia. Respondents were asked, "How safe from crime do you consider your neighborhood to be?" Possible responses were "extremely safe," "quite safe," "slightly safe," or "not at all safe." Respondents were classified as physically inactive if they reported no physical activity or exercise during the preceding month. Numbers for racial/ethnic groups other than white were combined because, when analyzed separately, data were too small for meaningful analysis. Data were weighted, and standard errors were calculated using SUDAAN (7).

The prevalence of physical inactivity among respondents was approximately 30% (n=3967), which is similar to the levels reported for adults in the United States (1). The prevalence of physical inactivity was highest among adults aged ≥65 years, women, racial/ethnic minorities, persons with a high school education or less, and persons with annual household incomes of <\$20,000 (Table 1). Overall, higher levels of perceived neighborhood safety were associated with lower levels of physical inactivity; the differences were greatest among persons aged ≥65 years (from 38.6% [extremely safe] to 63.1% [not at all safe]) and racial/ethnic minorities (from 29.9% [extremely safe] to 44.6% [not at all safe]). For respondents with more than a high school education, little difference in physical inactivity was noted among persons who perceived their neighborhood as unsafe and persons who perceived their neighborhood as safe (24.5% and 23.0%, respectively).

The prevalence of physical inactivity among men and women differed across neighborhood safety levels among persons aged 18–64 years but not among persons aged ≥65 years (Figure 1). Data stratified by age and sex and controlling for race and education demonstrated an association between neighborhood safety and physical inactivity among older adults (odds ratio=2.3; 95% confidence interval=1.1–4.7).

TABLE 1. Perceived neighborhood safety and the prevalence of physical inactivity among persons aged ≥18 years, by selected characteristics — Maryland, Montana, Ohio, Pennsylvania, and Virginia, Behavioral Risk Factor Surveillance System, 1996

Characteristic Total*		Extre	mely safe	Qu	iite safe	Slig	htly safe	Not at all safe		
		%	(95% CI [†])	%	(95% CI)	%	(95% CI)	%	(95% CI)	
Age (yrs)										
18-64	2898	30.5	(27.6-33.3)	29.1	(27.4 - 30.8)	35.1⁵	(31.7 - 38.4)	39.3	(31.5–47.0)	
≥65	1069	38.6	(33.6–43.7)	40.9	(37.6–44.2)	45.0	(36.9–53.0)	63.1	(48.8–77.5)	
Sex										
Men	1496	30.7	(26.9 - 34.4)	28.3	(26.0 - 34.4)	34.2	(29.6-38.8)	36.7	(25.3-48.1)	
Women	2471	33.8	(30.5–37.0)	33.8	(31.8–35.7)	38.5§	(34.5–42.5)	47.2	(38.8–55.7)	
Race/ Ethnicity [¶]										
White	3188	32.4	(29.8 - 35.1)	30.3	(28.6 - 32.0)	33.1	(29.5-36.7)	40.8	(31.6-49.9)	
Other	779	29.9	(23.0-36.9)	36.8	(32.9-40.7)	45.1§	(39.1–51.0)	44.6§	(34.0-55.3)	
Education level										
≤12 years	2451	41.7	(37.9 - 45.5)	40.3	(38.0-42.7)	44.8	(40.4-49.1)	51.3⁵	(42.9-59.7)	
>12 years	1516	23.0	(19.9–26.1)	22.4	(20.5-24.3)	25.4	(21.4–29.5)	24.5	(15.0-33.9)	
Annual income										
<\$20,000	938	43.8	(37.0-50.6)	42.9	(38.8 - 47.0)	42.5	(36.0-49.0)	44.0	(33.4-54.7)	
≥\$20,000	2269	30.7	(27.8–33.7)	28.6	(26.8–30.5)	34.4	(30.7–38.1)	39.8	(29.2–50.5)	

^{*}n=3967; numbers may not add to total because of missing data.

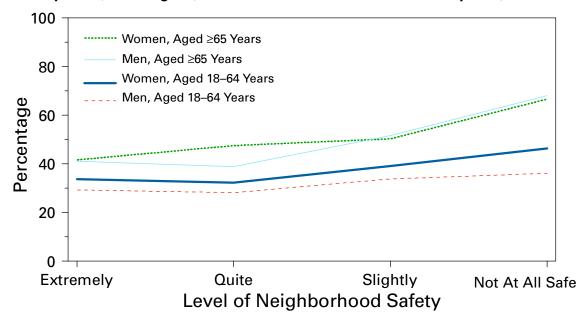
[†]Confidence interval.

[§]p ≤0.05 compared with "extremely safe."

[¶]Numbers for racial/ethnic groups other than white were combined because, when analyzed separately, data were too small for meaningful analysis.

Physical Inactivity — Continued

FIGURE 1. Percentage of respondents who reported physical inactivity, by sex, age group, and perceived neighborhood safety level — Maryland, Montana, Ohio, Pennsylvania, and Virginia, Behavioral Risk Factor Surveillance System, 1996



Reported by the following state BRFSS coordinators: A Weinstein, MA, Maryland; P Feigley, Montana; P Pullen, MS, Ohio; L Mann, Pennsylvania; L Redman, Virginia. Physical Activity and Health Br, Div of Nutrition and Physical Activity, National Center for Chronic Disease Prevention and Health Promotion, CDC.

Editorial Note: This report is the first to document the higher prevalence of physical inactivity among persons who perceive their neighborhoods as unsafe, and this finding remained after controlling for other factors. The findings were similar to those from other studies (1), which found that women were more physically inactive than men, and older adults were more inactive than younger adults. However, among older persons who perceived their neighborhoods as unsafe, the prevalence of physical inactivity in these states was similar among men and women.

Environmental barriers, including neighborhood safety, are not the only factors associated with physical inactivity among adolescents and young adults (3,8). However, many young adults use facilities, and the types of activities available are more varied. Among older adults, environmental barriers studied have been related to access to facilities (e.g., malls, parks, and gymnasiums) for physical activity rather than neighborhood safety issues (4,5,9). Older adults, for whom walking is the major activity, may be more influenced by safety concerns in their neighborhoods. These results suggest an association between perceived neighborhood safety and physical inactivity for adults aged ≥ 65 years.

The findings in this report are subject to at least five limitations. First, BRFSS data are cross-sectional and may not accurately reflect behaviors or conditions over time. Second, data are from only five states and may not represent trends in other states. Third, because the number of respondents in this analysis is relatively small and the data are self-reported, estimates may be unreliable. Fourth, because of the small number of respondents for racial/ethnic minorities, numbers were combined for a

Physical Inactivity — Continued

comparison with whites. Finally, these data may be affected by unmeasured confounding factors (e.g., social and demographic factors).

The survey described in this report suggests that public health action is needed to provide safe alternatives for physical activity in neighborhoods. Such efforts could increase community support and access to safe places for older adults to engage in physical activity. Additional research is needed to increase understanding of how perceived and actual neighborhood safety inhibits or facilitates participation in physical activity.

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Notice to Readers

Recall of Tripedia Vaccine

On January 27, 1999, the Food and Drug Administration initiated a voluntary recall of Tripedia[™] diphtheria and tetanus toxoids and acellular pertussis vaccine (DTaP), lot number 0916490, manufactured by Pasteur Mérieux Connaught USA.* Routine post-release stability testing completed in January 1999 indicated that the potency of the diphtheria toxoid component of this lot was below specification. The potency of the tetanus and pertussis components of this lot was acceptable.

The lot was distributed during February–June 1998. All lots of Tripedia[™] met potency specifications before release. Previously tested lots of Tripedia[™] met diphtheria potency specifications in routine stability testing after release; stability testing of additional lots is in progress.

A primary series (three doses) of fully potent diphtheria toxoid-containing vaccine is required to reliably induce protective antibody levels. Five doses of diphtheria toxoid-containing vaccine are recommended for preschool-aged children in the United States and provide optimal protection against diphtheria.

^{*}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.

Notice to Readers — Continued

The risk for exposure to toxigenic strains of *Corynebacterium diphtheriae* in the United States is low; however, diphtheria remains endemic in many countries. Additional doses of diphtheria toxoid-containing vaccine beyond those recommended in the childhood immunization schedule are associated with an increase in local reactions and should be considered only for children vaccinated with Tripedia™ lot number 0916490 who may be at increased risk for exposure to toxigenic strains of *C. diphtheriae*. CDC, the American Academy of Pediatrics, and the American Academy of Family Physicians have developed recommendations for children who received one or more doses of Tripedia™ lot number 0916490. The complete text of the recommendations is available on CDC's National Immunization Program World-Wide Web site, http://www.cdc.gov/nip/news/recall.htm; in summary, the recommendations are as follows:

- Children remaining in the United States until the scheduled fourth dose of DTaP or traveling to countries where the risk for diphtheria is low do not require any supplemental doses of diphtheria toxoid-containing vaccine.
- Children traveling to a country where the risk for diphtheria is high[†] before their scheduled fourth dose of DTaP may require a supplemental dose of DT or a dose of DTaP on an accelerated schedule; the recommendations vary based on the number of doses of TripediaTM lot number 0916490 received (Table 1).

[†]Travelers may be at substantial risk for exposure to toxigenic strains of *C. diphtheriae*, especially with prolonged travel, extensive contact with children, or exposure to poor hygiene. Countries comprise the following: *Africa*—Algeria, Egypt, and sub-Saharan Africa; *Americas*—Brazil, Dominican Republic, Ecuador, and Haiti; *Asia/Oceania*—Afghanistan, Bangladesh, Cambodia, China, India, Indonesia, Iran, Iraq, Laos, Mongolia, Myanmar, Nepal, Pakistan, Philippines, Syria, Thailand, Turkey, Vietnam, and Yemen; and *Europe*—Albania and all countries of the former Soviet Union.

TABLE 1. Recommendations for children who travel to areas where the risk for diphtheria is high* and who received TripediaTM diphtheria and tetanus toxoids and acellular pertussis vaccine (DTaP) lot number 0916490[†]

Age (mos)	Total doses received of any diphtheria toxoid-containing vaccine	No. doses of Tripedia TM from lot no. 0916490	Recommendation [§]
2–11	1–2	1–2	Complete primary series with DTaP¶
	3	1	Administer fourth dose of DTaP at age 15–18 mos
	3	2–3	Administer supplemental dose of DT, followed by fourth dose of DTaP at age 15–18 mos
≥12	3	1	Administer fourth dose of DTaP at age 15–18 mos
	3	2–3	Administer supplemental dose of DT if <6 months have elapsed since third dose of DTaP, followed by fourth dose of DTaP at age 15–18 mos
			OR
			Administer fourth dose of DTaP as early as age 12 mos if ≥6 months have elapsed since third dose of DTaP
	4	1–3	Administer fifth dose of DTaP at age 4 to 6 yrs
	5	1	Administer routine tetanus and diphtheria toxoids (for adolescent and adult use) boosters

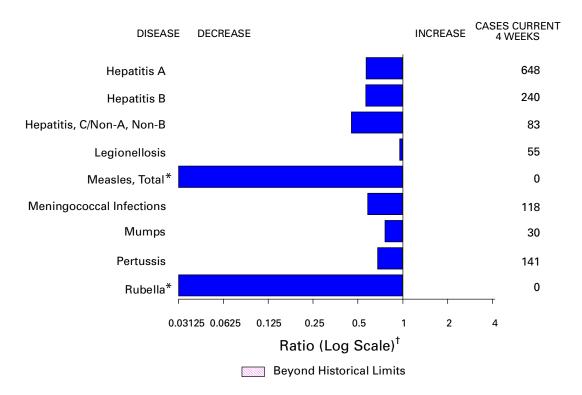
^{*}Travelers may be at substantial risk for exposure to toxigenic strains of *Corynebacterium diphtheriae*, especially with prolonged travel, extensive contact with children, or exposure to poor hygiene conditions. Countries comprise the following: *Africa*—Algeria, Egypt, and sub-Saharan Africa; *Americas*—Brazil, Dominican Republic, Ecuador, and Haiti; *Asia/Oceania*—Afghanistan, Bangladesh, Cambodia, China, India, Indonesia, Iran, Iraq, Laos, Mongolia, Myanmar, Nepal, Pakistan, Philippines, Syria, Thailand, Turkey, Vietnam, and Yemen; and *Europe*—Albania and all countries of the former Soviet Union.

[†]Manufactured by Pasteur Mérieux Connaught USA. 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.

¶If time is sufficient before travel, children who received two doses of Tripedia™ lot number 0916490 should receive their third dose of DTaP (as early as 4 weeks after the previous dose of DTaP) and a supplemental dose of DT (as early as 4 weeks after the third dose of DTaP).

[§]The minimum interval is 4 weeks between a dose of diphtheria and tetanus toxoids (for pediatric use) (DT) and any other dose of diphtheria toxoid-containing vaccine. The minimum interval is 6 months between the third dose of DTaP (including doses of TripediaTM lot number 0916490) and the fourth dose of DTaP.

FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending February 20, 1999, with historical data — United States



^{*}No measles or rubella cases were reported for the current 4-week period, yielding a ratio for week 7 of zero (0).

TABLE I. Summary — provisional cases of selected notifiable diseases, United States, cumulative, week ending February 20, 1999 (7th Week)

	Cum. 1999		Cum. 1999
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*	- 5 - 87 - 1 - - 5 1 5 7	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	- 2 - 20 146 4 - 1 8 1 22

^{-:} no reported cases

[†] 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.

^{*}Not notifiable in all states.

^{*}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 January 24, 1999.

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

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending February 20, 1999, and February 21, 1998 (7th Week)

						erichia 157:H7			Нера	titis
	AII			nydia	NETSS [†]	PHLIS	Gono		C/N/	
Reporting Area	Cum. 1999*	Cum. 1998	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1999	Cum. 1999	Cum. 1998	Cum. 1999	Cum. 1998
UNITED STATES	3,137	4,884	57,395	74,787	136	37	34,046	45,267	240	439
NEW ENGLAND	158	184 2	1,683	2,831	22	11	513	828 6	38	16
Maine N.H.	3 3	10	72 109	136 131	1 1	-	8 7	19	-	-
Vt. Mass.	124	5 70	51 1,143	36 1.156	- 14	- 7	6 407	1 297	38	2 14
R.I.	9	13	299	343	-	-	83	46	-	-
Conn.	19	84	9	1,029	6	4	2	459	-	-
MID. ATLANTIC Upstate N.Y.	489 17	1,005 116	8,745 N	10,787 N	8 6	-	4,464 258	6,181 682	4 4	25 23
N.Y. City	237 162	490	4,910	4,763	2	-	2,376	2,238	-	-
N.J. Pa.	73	237 162	484 3,351	1,579 4,445	N N	-	210 1,620	1,035 2,226	-	2
E.N. CENTRAL	179	317	9,795	11,562	31	4	7,181	9,247	80	72
Ohio Ind.	38 25	56 79	3,003	3,866	19 5	3	1,828 726	2,287 866	-	3 1
III.	77	101	3,558	2,746	2	-	2,211	2,681	_1	10
Mich. Wis.	22 17	57 24	2,829 405	3,081 1,869	5 N	1	2,233 183	2,677 736	79 -	58 -
W.N. CENTRAL	110	64	1,978	4,686	28	9	691	1,866	_	65
Minn.	20 3	15 6	578 234	951 455	12 5	8 1	225 73	331 131	-	- 1
lowa Mo.	72	22	234	1,624	1	-	-	791	-	63
N. Dak. S. Dak.	-	- 4	238	127 224	2	-	23	11 34	-	-
Nebr.	6	9	338	434	2	-	157	161	-	-
Kans.	9	8	590	871	6	-	213	407	-	1
S. ATLANTIC Del.	883 13	1,458 13	15,559 406	13,455 295	14	5	11,804 234	11,181 204	26	12
Md.	81	218	1,106	959	1	-	1,107	1,066	15	2
D.C. Va.	8 54	125 109	N 1,911	N 1,697	5	-	452 1,667	443 964	4	1
W. Va.	10	5	332	691	-	1	76	224	1	-
N.C. S.C.	69 60	105 91	2,981 3,859	2,535 2,203	2 1	2 1	2,662 2,038	2,237 1,633	1	5 -
Ga.	111	113	1,321	2,707	1 4	- 1	857	2,352	- 5	-
Fla. E.S. CENTRAL	477 157	679 266	3,643 4,686	2,368 5,001	7	-	2,711 4,214	2,058 5,156	5 17	4 17
Ky.	15	38	, -	759	-	-	-	507	-	4
Tenn. Ala.	64 31	82 86	1,814 1,783	1,761 1,283	5 2	-	1,527 1,657	1,618 1,746	16 1	11 2
Miss.	47	60	1,089	1,198	-	-	1,030	1,285	-	-
W.S. CENTRAL	532	871	4,143	10,460	3 1	-	2,948	6,594	7	10
Ark. La.	19 27	33 134	630 2,275	375 1,674	1	-	281 2,005	578 1,446	6	1 -
Okla. Tex.	6 480	52 652	1,238	1,032 7,379	- 1	-	662	572 3,998	- 1	9
MOUNTAIN	45	192	2,345	3,693	6	1	543	1,079	14	52
Mont.	-	5	163	107	-	-	3	['] 6	2	4
ldaho Wyo.	4	5 -	186	255 109	-	-	10 -	19 7	3 -	15 13
Colo.	26	39	707	839	2	1	151	376	3	3
N. Mex. Ariz.	4 4	36 61	590 522	576 1,299	2 1	-	118 243	114 455	3 2	7 -
Utah	4 U	22	177	266	1 U	-	18 U	31	1	6
Nev. PACIFIC	584	24 527	U 8,461	242 12,312	17	- 7	1,688	71 3,135	U 54	4 170
Wash.	29	31	341	1,427	1	2	48	258	2	1
Oreg. Calif.	15 525	13 468	508 7,170	763 9,611	6 10	5 -	71 1,491	124 2,652	- 52	1 136
Alaska	5	-	245	243	-	-	45	47	-	-
Hawaii	10	15	197	268	- N1	-	33	54	-	32
Guam P.R.	1 92	88	Ū	37 U	N 1	Ū	42	4 73	-	-
V.I. Amer. Samoa	-	8	N U	N U	N N	Ü	Ü	Ü	U U	U U
C.N.M.I.	-	-	N N	N N	N N	U	U	7	U	U

N: Not notifiable

U: Unavailable

^{-:} no reported cases

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

N: Not notificable U: Unavailable -: no reported cases C.N.M.I.: Commonwealth of Northern Mariana Islands
*Updated monthly from reports to the Division of HIV/AIDS Prevention–Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention, last update January 24, 1999.

†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 February 20, 1999, and February 21, 1998 (7th Week)

Legionellosis Disease Malaria (Primary & Secondary) Tuberco	758 37	Cum. 1999 482 86 14
NEW ENGLAND 6 11 44 56 2 6 12 12 33	37 - - 1	86 14
	- - 1	14
Maine 1 1	- 1	
N.H. 1 2 1 -		4
Vt. 2 1 Mass. 2 1 44 14 2 6 8 11 11	17	14 26
R.I. 1 3 - 2 13 Conn 5 - 39 3 - 8	4 15	7 21
MID. ATLANTIC 13 30 100 233 27 59 33 54 125	152	108
Upstate N.Y. 5 9 51 53 9 11 1 2 - N.Y. City - 5 - 7 3 36 13 5 70	14 102	70 U
N.J. 3 1 41 36 13 6 1 15 55 Pa. 5 15 8 137 2 6 18 32 U	36 U	27 11
E.N. CENTRAL 27 60 14 14 11 15 107 136 25	15	1
Ohio 12 18 8 10 1 1 12 34 U Ind. 5 7 5 3 4 1 32 26 U	U U	-
III 11 7 56 49 U Mich. 10 11 1 1 5 5 7 15 21	Ü	- 1
Wis 13 U U 1 1 - 12 4	15	-
W.N. CENTRAL 1 11 4 4 5 6 2 18 28 Minn 1 20	29 10	46 10
lowa 1 - 1 4 2 1 Mo 6 3 4 - 10 7	17	12
N. Dak 1	-	15
S. Dak 1 Nebr 5 1 4 -	-	1
Kans 2 1 1 3 -	2	8
S. ATLANTIC 22 18 50 55 33 32 274 354 83 Del. 2 1 1 1 1	154 1	204
Md 5 37 51 14 17 54 97 U D.C 2 1 2 5 2 10 9 4	U 13	44 -
Va. 2 2 4 2 21 36 9 W. Va. N N 1 - 1 - 5	5 10	48 10
N.C. 3 3 10 - 1 4 77 93 33 S.C. 3 1 40 45 32	87 38	50 11
Ga 2 - 4 26 24 U	U	19
Fla. 12 4 2 - 8 2 44 50 U E.S. CENTRAL 3 6 6 6 2 5 142 170 41	U 63	22 15
Ky 4 15 U Tenn. 3 1 2 5 2 3 80 87 U	Ü	13
Ala 4 1 - 1 44 36 39	40	2
Miss 1 1 18 32 2 W.S. CENTRAL 1 4 2 73 115 14	23 235	- 1
Ark 10 16 8 La. 1 3 2 27 52 U	5 U	-
Okla 36 7 6	17	1
Tex 1 40 - MOUNTAIN 4 7 1 1 5 8 - 34 12	213 33	- 7
Mont 1	-	, 3 -
Wyo	.1	-
Colo. 1 2 - - 1 3 - 2 U N. Mex. 1 1 1 - 1 3 - 2 3	U 7	1 -
Ariz 1 25 U Utah 2 4 1 - 2 9	U 6	3
Nev. U - U 1 U - U 3 U	19	U
PACIFIC 6 11 34 15 23 25 17 55 49 Wash 2 3 20	40 26	14 -
Oreg 5 - 1 U Calif. 6 11 34 15 20 20 16 51 U	U U	- 14
Alaska 6 Hawaii 1 - 1 - 23	4 10	-
Guam - 1	12	-
P.R 34 31 - V.I. U U U U U U U	3 U	6 U
Amer. Samoa U U U U U U U U U U U U U U U U U U U	Ü 9	ŭ

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

^{*}Cumulative reports of provisional tuberculosis cases for 1998 and 1999 are unavailable ("U") for some areas using the Tuberculosis Information Management System (TIMS).

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending February 20, 1999, and February 21, 1998 (7th Week)

	H. influ	ienzae,	н	epatitis (Vi	-	ое Ое	1	Measles (Rubeola)						
	inva	invasive		A		В	Indig	jenous	lm	orted [†]	Total			
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	115	150	1,603	2,399	548	1,049	-	7	-	2	9	2		
NEW ENGLAND	11	12	22	61	8	17	-	-	-	-	-	1		
Maine N.H.	1 1	1	2 2	8 3	2	2	-	-	-	-	-	-		
Vt.	2	-	-	2	-	-	-	-	-	-	-	-		
Mass. R.I.	7 -	11 -	7 -	16 4	4 2	9	-	-	-	-	-	-		
Conn.	-	-	11	28	-	6	U	-	U	-	-	-		
MID. ATLANTIC Upstate N.Y.	17 11	19 6	70 16	178 41	52 15	160 31	-	-	-	-	-	1		
N.Y. City	-	6	11	74	6	44	-	-	-	-	-	-		
N.J. Pa.	6	7	24 19	31 32	8 23	27 58	-	-	-	-	-	1 -		
E.N. CENTRAL	14	26	470	443	61	265	_	_	-	-	_	-		
Ohio	12	12	93	58	16	11	-	-	-	-	-	-		
Ind. III.	1 1	2 11	29 31	58 116	4	121 37	-	-	-	-	-	-		
Mich. Wis.	-	- 1	316 1	184 27	41	77 19	-	-	-	-	-	-		
W.N. CENTRAL	3	1	28	233	12	61	-	_	-	_	_	-		
Minn.	-	-	2	5	1	2	-	-	-	-	-	-		
lowa Mo.	1 -	-	7 3	77 127	4	10 42	-	-	-	-	-	-		
N. Dak.	-	-	-	-	-	-	-	-	-	-	-	-		
S. Dak. Nebr.	1 -	-	13	1 3	6	1 1	-	-	-	-	-	-		
Kans.	1	1	3	20	1	5	-	-	-	-	-	-		
S. ATLANTIC Del.	37	23	186	156	98	90	-	-	-	-	-	-		
Md.	19	8	49	52	24	21	-	-	-	-	-	-		
D.C. Va.	2	3	9 12	6 20	2 7	1 7	-	-	-	-	-	-		
W. Va.	1	1	-	-	-	-	-	-	-	-	-	-		
N.C. S.C.	2 2	3 -	20 1	13 6	31 10	40	-	-	-	-	-	-		
Ga. Fla.	- 11	7 1	46 49	27 32	4 20	11 10	-	-	-	-	-	-		
E.S. CENTRAL	9	12	62	68	43	60	-	_		-	_	_		
Ky.	-	2	-	2	-	3	Ū	-	U	-	-	-		
Tenn. Ala.	5 4	5 5	42 19	39 14	32 11	44 13	-	-	-	-	-	-		
Miss.	-	-	1	13	-	-	U	-	U	-	-	-		
W.S. CENTRAL	9	7	73	170	19	72	-	-	-	2	2	-		
Ark. La.	3	3	3 6	3 3	6 4	14 3	-	-	-	-	-	-		
Okla. Tex.	4 2	3 1	26 38	55 109	1 8	3 52	-	-	-	2	2	-		
MOUNTAIN	9	31	123	411	58	106	_	1	_	-	1	-		
Mont.	1	-	2	6	1	1	-	-	-	-	-	-		
ldaho Wyo.	1 -	-	4	26 3	4	4 1	Ū	-	Ū	-	-	-		
Colo. N. Mex.	1	3	51 5	43 27	15	12	-	1	-	-	1	-		
Ariz.	2	16	50	246	27 6	38 29	Ū	-	Ū	-	-	-		
Utah Nev.	4 U	2 10	11 U	25 35	5 U	8 13	- U	- U	- U	Ū	- U	-		
PACIFIC	6	19	569	679	197	218	-	6	-	-	6	_		
Wash.	-	-	20	46	2	14	-	-	-	-	-	-		
Oreg. Calif.	5 -	9 8	27 519	49 575	4 189	19 180	-	6	-	-	6	-		
Alaska Hawaii	1	2	2	9	2	1	-	-	-	-	-	-		
Guam	-	_	- -	9	-	4	- U	_	U	-	-	-		
P.R.		1	8	6	9	69	-		-					
V.I. Amer. Samoa	U U	U U	U U	U U	U U	U U	U U	U U	U U	U U	U U	U U		
C.N.M.I.	-	-	-	-	-	7	ŭ	-	Ŭ	-	-	-		

N: Not notifiable

U: Unavailable

^{-:} no reported cases

^{*}Of 16 cases among children aged <5 years, serotype was reported for 4 and of those, 1 was 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 February 20, 1999, and February 21, 1998 (7th Week)

		ococcal ease		Mumps	21, 133		Pertussis			Rubella	
Reporting Area	Cum. 1999	Cum. 1998	1999	Cum. 1999	Cum. 1998	1999	Cum. 1999	Cum. 1998	1999	Cum. 1999	Cum. 1998
UNITED STATES	246	473	7	44	44	29	278	511	-	3	30
NEW ENGLAND	17	29	-	1	-	2	51	108	-	-	10
Maine N.H.	3	3 1	-	- 1	-	-	3	4 8	-	-	-
Vt.	1	1	-	-	-	2	10	19	-	-	-
Mass. R.I.	13 -	10 3	-	-	-	-	38	74 -	-	-	1 -
Conn.	-	11	U	-	-	U	-	3	U	-	9
MID. ATLANTIC Upstate N.Y.	28 7	43 11	2 2	5 2	1 1	2 2	16 15	45 29	-	-	14 12
N.Y. City	7 10	8 14	-	-	-	-	-	3	-	-	2
N.J. Pa.	4	10	-	3	-	-	1	9	-	-	-
E.N. CENTRAL	36	78	1	2	5	10	57	68	-	-	-
Ohio Ind.	20 7	30 8	-	1 -	3	9 1	50 2	28 2	-	-	-
III. Mich.	5 4	20 8	- 1	- 1	2	-	- 5	10	-	-	-
Wis.	-	12	-	-	-	-	5 -	28	-	-	-
W.N. CENTRAL	14	40	-	1	-	-	5	32	-	-	-
Minn. Iowa	4	8	-	1	-	-	3	18 7	-	-	-
Mo. N. Dak.	3	19	-	-	-	-	1	2	-	-	-
S. Dak.	4	4	-	-	-	-	1	-	-	-	-
Nebr. Kans.	1 2	1 8	-	-	-	-	-	2	-	-	-
S. ATLANTIC	48	66	3	9	8	3	41	43	_	3	1
Del. Md.	1 6	- 10	- 1	2	-	-	- 13	- 7	-	-	-
D.C.	-	-	-	-	-	-	-	-	-	-	-
Va. W. Va.	2	7 2	-	-	1	-	6	-	-	-	-
N.C.	6	14	-	1	4	-	16	23	-	3	1
S.C. Ga.	6 8	5 21	-	2	2	-	2	5 -	-	-	-
Fla.	19	7	2	4	1	3	4	8	-	-	-
E.S. CENTRAL Ky.	20	44 8	Ū	-	-	2 U	9	11 -	Ū	-	-
Tenn.	9	15	-	-	-	2	6	3	-	-	-
Ala. Miss.	11 -	19 2	Ū	-	-	Ū	3	8 -	Ū	-	-
W.S. CENTRAL	11	27	-	9	11	1	12	12	-	-	1
Ark. La.	3 6	5 8	-	-	-	-	3	3	-	-	-
Okla.	1	13 1	-	1	- 11	1	2 7	- 9	-	-	-
Tex. MOUNTAIN	1 22	28	1	8	4	- 7	7 79	113	-	-	1 3
Mont.	-	1	-	-	-	-	-	1	-	-	-
Idaho Wyo.	3	1 1	Ū	-	1	6 U	50	54 -	Ū	-	-
Colo. N. Mex.	5 6	11 3	1 N	2 N	- N	1 -	5 7	14 37	-	-	-
Ariz.	5	9	Ü	-	1	Ū	2	3	Ū	-	-
Utah Nev.	3 U	1 1	Ū	1 U	2	Ū	15 U	2 2	Ū	Ū	2 1
PACIFIC	50	118	-	14	15	2	8	79	-	-	1
Wash. Oreg.	6 4	12 28	- N	- N	N	2	4 3	11 8	-	-	-
Calif.	33	75	-	12	9	-	-	60	-	-	1
Alaska Hawaii	3 4	1 2	-	1 1	2 4	-	1 -	-	-	-	-
Guam	-	-	U	-	1	U	-	-	U	-	-
P.R. V.I.	- U	- U	Ū	- U	- U	- U	- U	- U	Ū	- U	Ū
Amer. Samoa	U	U	U	U	Ū	Ū	U	Ü	U	U	Ü
C.N.M.I.	-	-	U	-	2	U	-	-	U	-	-

N: Not notifiable

U: Unavailable

TABLE IV. Deaths in 122 U.S. cities,* week ending February 20, 1999 (7th Week)

	1	All Cau	ises, By	/ Age (Y		u. ,	P&I [†]		,	All Cau	ıses, By	Age (Y	ears)		P&I [†]
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.	66 79 2 36	553 118 34 16 37 60 26 8 33 43 65 2	33 6 6 1 12 5 5 2 16 8	31 10 2 - 3 1 - 2 5 4 - 1	12 4 - - 1 1 - 2 2 2	5 4 - - 1 - - - -	104 31 4 5 12 2 3 11	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.	1,094 U 110 123 127 107 58 76 75 77 226 101	736 U 75 79 87 62 46 47 54 68 145 61	209 U 18 29 34 26 6 19 17 6 32 20 2	73 U 11 12 3 14 2 5 3 1 9	23 U 2 1 2 5 - 1 3 4	18 U 4 2 1 - 4 - 1 1 2 3	85 U 13 14 4 2 9 10 13 16 4
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.	34 68 2,403 54 14 85 34 14 55 33	27 54 1,733 38 12 64 24 9 45 23 917	12 487 11 1 13 6 5 7 8	1 2 124 2 1 5 1 - 2 2 60	1 34 2 - 1 - - - 1 1	25 1 - 2 3 - 1	18 125 3 - 1 3 1 3 -	E.S. CENTRAL Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala. Nashville, Tenn. W.S. CENTRAL	876 217 60 115 67 222 43 27 125	615 149 44 82 48 142 37 23 90	151 43 11 23 10 37 6 3 18	58 15 3 8 1 23 - 1 7	23 3 1 1 2 12 - 4 42	26 5 1 1 5 8 - 6 44	61 18 3 4 5 21 2 5 3
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.	70 22 299 56 41 156 29 47 69 24 39 U	33 13 203 44 34 117 25 38 48 17 29 U	24 5 62 9 3 31 3 7 14 6	99 21 23 55 11 51 10	3 2 7 1 1 3 - 1 2	11 - 6 - - - - - U	30 3 28 8 5 17 5 3 6 7 2 U	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.	84 62	63 41 37 132 37 88 274 55 13 158 48 74	14 11 7 44 10 21 101 8 4 29 10	6 5 2 20 2 5 41 3 3 12 4 5	2 5 2 15 3 4 5 2	1 52 4 7 9 2 1 4 2 7	3 5 5 5 1 24 28 3 - 27 16 7
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, Ill. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind.	2,338 54 42 367 102 138 278 127 268 58 71	1,682 43 34 236 69 95 205 90 195 48	7 6 77 17 28 50 25 49 6	134 2 1 29 5 10 13 8 15 2	47 1 15 3 1 6 2 6	50 2 9 8 4 4 2 3 1	168 6 28 15 5 20 8 6 3	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.	117 209 35 80 34	741 95 44 31 75 154 28 56 27 77	145 16 7 9 19 33 6 15 4 17	66 8 2 3 12 13 1 8 3 7	22 1 - 4 6 - 1 - 5	19 1 - 4 7 3 - - 2 2	105 9 4 - 20 16 7 - 8 14 27
Gary, Ind. Grand Rapids, Micl Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio	221 47 139 65 64 52 93 74	9 51 151 30 104 48 53 44 59 60	10 39 13 25 14 6 6 25 8	3 20 3 7 2 2 2 5 3	2 1 2 1 2 3 1	2 9 - 1 - 1 - 2	4 22 5 19 3 9 2 5 4	PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawaii Long Beach, Calif. Los Angeles, Calif. Pasadena, Calif. Portland, Oreg. Sacramento, Calif. San Diego, Calif.	1,589 16 132 13 61 69 291 26 91 204	1,177 13 102 9 45 50 215 18 74 161 121	260 2 17 3 11 12 40 4 11 31 35	96 - 13 - 3 1 22 - 3 9	28 - 1 1 2 8 - 1 2 2 U	26 1 1 4 6 4 2 1	168 1 15 5 6 17 4 12 43 20
W.N. CENTRAL Des Moines, lowa Duluth, Minn. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	586 U 30 U 118 61 98 75 113 91 U	441 U 26 U 83 45 77 57 81 72 U	2 U 19 9 15 7 19	39 U 2 U 5 7 3 7 11 4 U	10 U 5 - 1 1 3 U	15 U - U 6 - 3 3 1 2 U	46 U 2 U 11 4 12 6 11 U	San Francisco, Calit San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash.	176 156 37 141 65 111 12,084 [¶]	U 115 29 97 48 80	U 27 3 30 11 23	14 U 11 2 8 4 6 729	U 2 3 4 2 - 241	2 - - 2 228	20 U 13 4 5 10 13 986

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.

Contributors to the Production of the *MMWR* (Weekly) Weekly Notifiable Disease Morbidity Data and 122 Cities Mortality Data

Samuel L. Groseclose, D.V.M., M.P.H.

State Support Team Robert Fagan Scott Connolly Gerald Jones David Nitschke Carol A. Worsham

CDC Operations Team Carol M. Knowles Deborah A. Adams Willie J. Anderson Patsy A. Hall Amy K. Henion

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and Prevention Jeffrey P. Koplan, M.D., M.P.H. Deputy Director, Centers for Disease Control and Prevention Claire V. Broome, M.D.

Director, Centers for Disease Control Director, Epidemiology Program Office Stephen B. Thacker, M.D., M.Sc. Editor, MMWR Series John W. Ward, M.D. Managing Editor, MMWR (weekly) Karen L. Foster, M.A.

Writers-Editors, MMWR (weekly) Jill Crane David C. Johnson Teresa F. Rutledge Caran R. Wilbanks Desktop Publishing Morie M. Higgins Peter M. Jenkins

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