

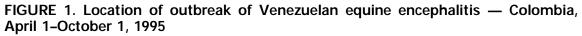
October 6, 1995 / Vol. 44 / No.39

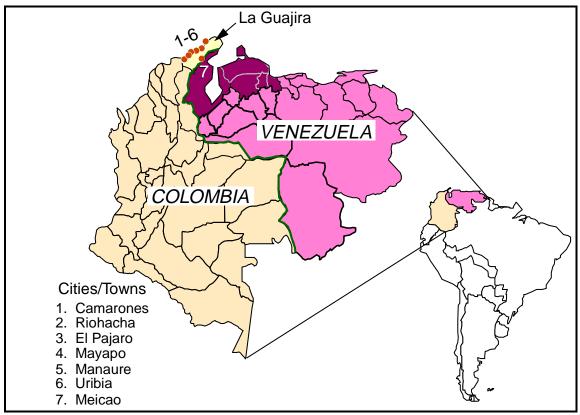
- 721 Venezuelan Equine Encephalitis Colombia, 1995
- 724 Outbreak of Gastrointestinal Illness Associated with Consumption of Seaweed — Hawaii, 1994
- 727 Acute Lower Respiratory Tract Illness in Illicit Drug Users — South Carolina, 1995
- 734 Patterns of Homicide Cali, Colombia, 1993–1994
- 737 National Surveillance
- for Infectious Diseases, 1995 739 Monthly Immunization Table

Venezuelan Equine Encephalitis — Colombia, 1995

MORBIDITY AND MORTALITY WEEKLY REPORT

An outbreak of Venezuelan equine encephalitis (VEE) that began in northwestern Venezuela in April 1995 has spread westward to the Guajira peninsula and to Colombia (Figure 1), resulting in an estimated minimum of 13,000 cases in humans and an undetermined number of equine deaths. Governments of both countries have initiated efforts to control the spread of this outbreak by quarantining and vaccinating equines and applying insecticides. This report summarizes the ongoing investigation of the outbreak in Colombia.





U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES / Public Health Service

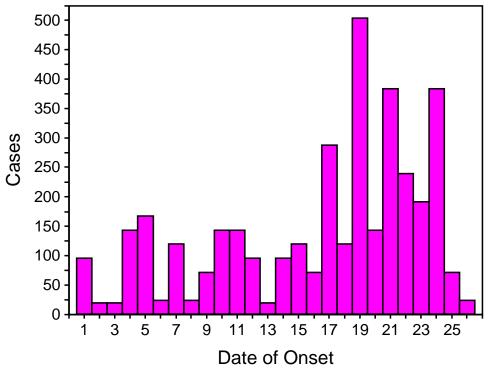
Venezuelan Equine Encephalitis — Continued

During the first week of September, rural health clinics in the towns of Mayapo, Manaure, and El Pajaro in the municipality of Manaure in La Guajira state reported an increased number of patients seeking care for acute febrile illnesses characterized by intense headache, muscle pain, prostration, and vomiting. Illness in some patients was complicated by convulsions and other neurologic symptoms.

As of September 28, a total of 8320 persons with acute febrile illness compatible with VEE had been treated at public hospitals and clinics in La Guajira, and large outbreaks had been reported from the towns of Manaure, Riohacha, El Pajaro, Mayapo, Uribia, and Meicao. Based on a random survey of 250 residents of Manaure, a recent history of acute illness compatible with VEE was present in 57% of respondents (Figure 2); 4% reported associated convulsions, and one person died (case-fatality rate=0.7%). All age groups were equally affected. In Manaure, the epidemic peaked on September 19, and malathion spraying was initiated on September 21.

In Riohacha, the state capitol, hospital visits for acute febrile illness increased steadily in September, reaching 143 visits on September 24 with no evidence of a decline. A similar pattern occurred in Uribia. Based on interviews and physical examinations of 23 inpatients at local hospitals on September 27–28, prominent manifestations included fever (100%); convulsions (98%); headache (56%); photophobia (56%); myalgias (56%); and chills, vomiting, and diarrhea (48% each). Ten associated deaths were reported statewide. Unidentified viral isolates have been recovered from four of 18 human blood samples submitted to the Colombian National Institute of Health.

FIGURE 2. Number of Venezuelan equine encephalitis cases estimated from a random household survey (n=250), by date of onset — Manaure,* Colombia, September 1–26, 1995



^{*} Population 6283.

Venezuelan Equine Encephalitis — Continued

Because of a prolonged rainy season (the heaviest in 20 years in La Guajira), mosquito abundance has increased dramatically. *Aedes aegypti* house indices increased in August to 70% in Manaure and to 22% in Riohacha. Entomologic surveys in Manaure detected large numbers of *Psorophora confinnis* and *Ae. taeniorhynchus* breeding in estuarine waters in the town's vicinity. The equine population in La Guajira consists of approximately 70,000 unvaccinated horses, donkeys, and mules owned by native Wayuu people, who constitute approximately 35% of the inhabitants of Riohacha. Control measures instituted by the government of Colombia include vaccination of equines in La Guajira, restriction of equine movement from and within the state, large-scale application of insecticides, public education and community mobilization campaigns to eradicate mosquito breeding sites, issuance of guidelines on case-management and referral, and surveillance of humans and equines.

Reported by: E Daza, V Frias, A Alcola, I Lopez, I Bruzon, La Guajira Health Dept; JT Montero, G Alvarez, MA Garcia, R Rodriguez, Colombia Ministry of Health; J Boschell, F de la Hoz, F Rivas, V Olano, LA Diaz, FM Caceras, G Aristizabal, V Cardenas, Colombian National Institute of Health, Colombia. J Cuellar, Pan American Health Organization, Colombia. E Gonzalez, Pan American Health Organization, Venezuela. A Ruiz, F Pinheiro, R Gusmao, Pan American Health Organization, Washington, DC. S Weaver, R Tesh, Univ of Texas Medical Br, Galveston, Texas. R Ricco-Hesse, Yale Arbovirus Research Unit, Yale Univ, New Haven, Connecticut. Div of Vector-Borne Infectious Diseases, National Center for Infectious Diseases, CDC.

Editorial Note: Major epizootics and concurrent epidemics of VEE have occurred periodically in northern areas of South America, resulting in hundreds of thousands of human and equine cases. The current epidemic is the largest in the region since 1962– 1971, when outbreaks affected Colombia, Ecuador, Peru, Venezuela, all the countries of Central America (except Panama), Mexico, and Texas (1,2). No major outbreaks had been recognized since the outbreak that occurred on La Guajira peninsula in 1973, suggesting that epizootic viral strains (subtypes IAB and IC) had become extinct. However, based on recent molecular phylogenetic studies, these strains may have evolved from enzootic ID strains maintained in silent cycles of rodent-mosquito transmission. This hypothesis predicts that strains with epidemic and epizootic potential will continue to emerge periodically from enzootic reservoirs (1,3,4). Partial nucleotide sequencing and antigenic analysis of three human isolates from the current epidemic indicates they are related to the IC epizootic strain of VEE virus isolated during a large outbreak in Venezuela and Colombia during 1962-1964 (S. Weaver, University of Texas, and R. Ricco-Hesse, Yale University, personal communications, 1995) and to a strain isolated from a mosquito pool in Venezuela in 1983 (1).

Although conditions leading to the emergence of VEE epidemics have not been clearly defined, previous outbreaks also were associated with heavy rains and flooding in arid rural areas, especially during the dry season. During epidemics and epizootics, VEE virus is transmitted rapidly among equines and from equines to humans by a variety of mosquito species. Horses are the principal amplifying hosts in epidemic transmission because they develop and sustain high levels of viremia and attract large numbers of biting mosquitoes. Cases among humans generally occur 2 weeks after epizootic infections in horses, and epidemic transmission ceases after susceptible horses have been either infected or vaccinated. Because the total number of horses and other equines in South America has declined since the last major outbreaks in 1971, the current outbreak suggests a possible role of human and other animal infections in sustaining the epidemic cycle. VEE virus levels in human blood

Venezuelan Equine Encephalitis — Continued

are sufficiently high to infect mosquitoes, and virus has been isolated from the pharynx of ill persons, indicating the virus could be transmitted between humans by mosquitoes or by close direct contact (5).

The clinical features of cases in the current outbreaks are consistent with those reported in previous VEE epidemics in which neurologic symptoms developed in 4% of cases, primarily among children and the elderly (6). VEE infections during pregnancy may result in fetal infection and malformations. VEE outbreaks have the potential for substantial social impact: for example, during the 1967 outbreak in Colombia, an estimated 220,000 human cases immobilized villages and local clinics (7).

The Pan American Health Organization coordinates a surveillance system for equine encephalitis in the Americas and is assisting countries of Latin America and the Caribbean in strengthening their diagnostic capacity. Equine vaccination with the live attenuated TC-83 vaccine provides rapid immunity and, when combined with restriction of equine movements, may limit the spread of epizootics and prevent their emergence (1).

No commercially licensed human VEE vaccine is available. Persons who cannot defer travel to these areas should use insect repellents, stay in air-conditioned or well-screened accommodations when possible, and wear long-sleeved shirts and long pants. The incubation period for VEE is 2–5 days. Health-care providers who suspect VEE in ill returned travelers can refer clinical specimens through state health departments to CDC for diagnosis.

References

- 1. Weaver SC, Belew LA, Rico-Hesse R. Phylogenetic analysis of alphaviruses in the Venezuelan equine encephalitis complex and identification of the source of epizootic viruses. Virology 1992; 191:282–90.
- 2. Walton TE, Grayson MA. Venezuelan equine encephalomyelitis. In: Monath TP, ed. The arboviruses: epidemiology and ecology. Boca Raton, Florida: CRC Press, 1989;203–32.
- 3. Ricco-Hesse R, Weaver SC, deSiger J, Medina G, Salas RA. Emergence of a new epidemic/ epizootic Venezuelan equine encephalitis virus in South America. Proc Natl Acad Sci U S A 1995;92:5278–81.
- 4. Kinney RM, Tsuchiya KR, Sneider JM, Trent DW. Genetic evidence that epizootic Venezuelan equine encephalitis (VEE) viruses may have evolved from enzootic VEE subtype I-D virus. Virology 1992;191:579–80.
- 5. Bowen GS, Calisher CH. Virological and serological studies of Venezuelan equine encephalomyelitis in humans. J Clin Microbiol 1976;4:22–7.
- 6. Briceno-Rossi AL. Rural epidemic encephalitis in Venezuela caused by a group A arbovirus (VEE). Prog Med Virol 1967;9:176–203.
- 7. Sanmartin C, Mackenzie RB, Trapido H, et al. Equine encephalitis in Colombia, 1967. Bol Oficina Sanit Panam 1973;74:108–37.

Outbreak of Gastrointestinal Illness Associated with Consumption of Seaweed — Hawaii, 1994

Seaweed is frequently served as a side dish at meals in the Pacific Islands and is a common component in the diet of many persons living in the Pacific Rim. Seaweed is often harvested at beaches, gathered in nearshore waters, or purchased at local markets. It is served either raw or cooked and is commonly prepared with salt and/or other

Seaweed-Related Illness — Continued

spices and herbs (e.g., chili pepper, ginger, and garlic). Previous reports have documented a toxic illness associated with seaweed harvested in some locations in the Pacific (1,2). This report summarizes the investigation of an outbreak of acute gastro-intestinal illness associated with consumption of seaweed during a picnic in Hawaii in September 1994.

On September 9, 1994, a 60-year-old woman contacted the Maui County Health Department (MCHD) to report a burning sensation in her mouth and throat that occurred 15 minutes after she tasted a seaweed ("ogo") preparation. The woman had received the seaweed from a friend on September 1 and had prepared the seaweed on September 2 by washing it, then boiling it in water for 1–2 minutes, and finally mixing it with vinegar, sugar, soy sauce, and sliced cucumbers. The seaweed mixture then was refrigerated. Approximately 4 hours after preparation, she tasted the mixture and, 15 minutes later, had onset of a sore throat and mouth, and headache. She was examined by a physician on September 3; a throat culture obtained during the visit was negative.

MCHD determined that the seaweed from which the woman's portion had come was to be served at a picnic on September 5 on the island of Hawaii. When contacted by the Hawaii County Health Department, some picnic attendees reported having had onset of nausea and diarrhea soon after eating at the picnic.

To assess the role of different foods as risk factors for illness, the Hawaii Department of Health (HDOH) surveyed all 13 picnic attendees. A questionnaire was administered by telephone to the attendees regarding menu items consumed, onset of illness, and symptoms. Menu items included poi, two types of fish (mamo and awe), beef stew, ogo (seaweed), salmon salad, steak, rice, tuna salad, chips, crackers, beer, soda, and water. A case was defined as onset of a burning sensation in the mouth or throat or two or more of the following symptoms: vomiting, diarrhea, nausea, or lethargy within 2 hours after eating food items from the picnic.

The seaweed had been prepared on September 4 by washing, removing debris, and cooking in boiling water. It was then mixed with codfish, vinegar, onion, soy sauce, and chili peppers. The woman who had prepared the seaweed and her husband had tasted it soon after preparation, and both noted a burning sensation in their throats that lasted for 4 hours. They attributed the burning sensation to an excessive amount of chili peppers. Before preparation, the seaweed had been stored in previously unused plastic bags and refrigerated.

Of the 13 persons who attended the picnic, eight were aged \geq 18 years. Illness in seven (54%) attendees met the case definition, and onset occurred 15–90 minutes after eating the meal. All seven ill persons were adults, and five were males. Symptoms included diarrhea (71%), nausea (71%), vomiting (60%), and a burning sensation (57%), and the mean duration of gastrointestinal symptoms was 22 hours. None of the ill persons were examined by a physician.

The only foods statistically associated with risk for illness were seaweed (illness in seven of seven who consumed versus none of six who did not [relative risk (RR)= undefined]) and salmon salad (RR=undefined). Two of the seven persons who had consumed seaweed had eaten less than one serving (i.e., one serving spoon) and experienced only a burning sensation in the mouth and throat; in comparison, the five persons who ate one or more servings experienced gastrointestinal illness.

Seaweed-Related Illness — Continued

A leftover sample of the seaweed served at the picnic and another sample harvested from the same site were examined at the University of Hawaii for species identification and toxicity studies. The seaweed was identified as *Gracilaria coronopifolia*. It was rinsed thoroughly with fresh water and extracted with acetone. The toxicity of the crude extract was assessed by intraperitoneal injection into mice; the mice were observed for symptoms of toxicity. Manifestations of toxicity in the mice included diarrhea at lower dosages and death within 15 minutes at the most concentrated dosages. The compound that induced the most extreme signs of toxicity in the mice was isolated using high-performance liquid chromatography; however, the isolate was not identified.

Bacterial organisms were identified microscopically both on the surface and in the seaweed. These bacteria were cultured or isolated using 2% NaCl Heart Infusion Agar (DIFCO,* Detroit, Michigan). Colonies of two different species of *Pseudomonas* and *Vibrio* grew on the culture media. Each culture was then extracted, and the extracted fraction was subjected to the mouse-toxicity test. Mice injected with the extracted fraction from these bacteria exhibited transient weakness but no other signs of toxic-ity.

The seaweed had been harvested from a site in a bay on a northeast-facing shore in Maui County in water at a depth of 3–5 feet. Two fresh-water streams flow into the bay, and discharge from storm drains pours into one of the streams. In addition, the tides carry tree branches and other debris into the bay. The State Wastewater Management Division of the Hawaii Department of Health collects water samples from this area on a monthly basis but had not documented increased levels of fecal coliforms or any other potential pollutants during the period before the seaweed was harvested. There was no evidence the seaweed had been exposed to pesticides.

HDOH has notified physicians throughout the state about the potential for seaweed-induced toxicity and has requested that physicians report any cases to HDOH. Seaweed samples will be collected from the same site from which the original samples were obtained for toxin surveillance.

Reported by: M Hanne, H Matsubayashi, R Vogt, MD, State Epidemiologist, C Wakida, Hawaii State Dept of Health; S Hau, State Dept of Land and Natural Resources; H Nagai, Y Hokama, Univ of Hawaii. L Solorzano, Food and Drug Administration. Div of Field Epidemiology, Epidemiology Program Office, CDC.

Editorial Note: The investigation of this outbreak in Hawaii indicated that consumption of seaweed was associated with acute illness in picnic attendees. In addition to the epidemiologic findings, this conclusion was supported by the isolation of toxin that caused both similar illness and more severe illness in mice. The toxin most likely was elaborated by the seaweed itself or by a coexistent microorganism and probably was heat resistant because both samples of seaweed were boiled before consumption.

Although this outbreak was the first reported episode of seaweed-related illness in Hawaii, this problem has been reported previously in other areas in the Pacific Rim. For example, in 1991, a total of 13 persons became ill, and three of them died after eating seaweed harvested in Guam (3); the seaweed species was identified as *Gracilaria tsudai*, and manifestations included gastrointestinal illness, fever, wheezing, muscle fasciculations, and hypotension. In 1992, three persons had onset of illness after

^{*}Use of trade names and commercial sources is for identification only and does not imply endorsement by the Public Health Service or the U.S. Department of Health and Human Services.

Seaweed-Related Illness — Continued

eating seaweed harvested on a beach in California (4); the seaweed species implicated in that episode was *Grasilariopsis lemanaeformis*. In 1993, two persons became ill, and one of them died after eating *Gracilaria verrucosa* seaweed in Japan (1). In the episodes in both Guam and California, the implicated seaweed previously had not been known to develop toxicity.

Although the mechanism of development of toxicity in seaweed has not been clearly determined, findings of previous studies suggest that some species of seaweed may become toxic at the end of their reproductive cycle, and thereby exhibit a seasonal variation in toxin production (2). Other suggested mechanisms are that stress from over-harvesting may cause seaweed to begin elaborating toxin as a method of protection (5) and that environmental changes and increased pollution promote colonization of toxin-producing bacteria.

State and local health departments should inform persons who may eat seaweed that seaweed consumption can be associated with illness and that varieties previously consumed with safety may undergo changes that increase their potential for causing illness. Episodes of seaweed-related illness should be reported promptly to state and local health departments.

References

- 1. Noguchi T, Matsui T, Miyazawa K. Poisoning by the red alga 'Ogonori' (*Gracilaria verrucosa*) on the Nojima Coast, Yokohama, Kanagawa Prefecture, Japan. Toxicon 1994;32:1533–8.
- 2. Yasumoto T. Guam seaweed poisoning: biochemistry of the *Grasilaria* toxin. Micronesica 1993;26:53–7.
- 3. Haddock R. Guam seaweed poisoning: food histories. Micron 1993;26:35-8.
- 4. Snyder B. Seaweed suspect in poisoning of Livermore family. Oakland Tribune, Nov 2, 1992: 1(col 1).
- Meyer KD, Paul V. Intraplant variation in secondary metabolite concentration in three species of Caulerpa (Chlorophyta: Caulerpales) and its effects on herbivorous fishes. Marine Ecology Progress Series 1992;82:249–57.

Acute Lower Respiratory Tract Illness in Illicit Drug Users — South Carolina, 1995

On July 31, 1995, the South Carolina Department of Health and Environmental Control was notified of a cluster of five patients with acute, severe lower respiratory illnesses among previously healthy residents of a small rural community in Berkeley County (1990 population: 128,776). All five patients were users of illicit drugs. This report summarizes the preliminary findings of an investigation initiated to describe the clinical features and epidemiology of this syndrome and to determine an etiology.

Based on information about the five cases obtained from interviews with the patients and reviews of records, a case was defined as an unexplained acute, severe respiratory illness in a previously healthy person aged <65 years characterized by shortness of breath and/or pleuritic pain with onset of symptoms during July 15–31. One additional case was identified by contacting local physicians, intensive-care units, and pulmonary and infectious disease specialists. No cases of similar acute respiratory illness were noted in household contacts of patients.

CASES CURRENT INCREASE DISEASE DECREASE 4 WEEKS Hepatitis A 1,853 Hepatitis B 566 Hepatitis, C/Non-A, Non-B 231 72 Legionellosis Malaria 107 Measles, Total* 4 Meningococcal Infections 111 Mumps 30 Pertussis 448 Rabies, Animal 500 Rubella 4 0.03125 0.0625 0.125 0.25 0.5 1 2 4 Ratio (Log Scale)[†]

FIGURE I. Notifiable disease reports, comparison of 4-week totals ending September 30, 1995, with historical data — United States

* The large apparent decrease in the number of reported cases of measles (total) reflects dramatic fluctuations in the historical baseline.

[†]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 — cases of specified notifiable diseases, United States, cumulative, week ending September 30, 1995 (39th Week)

	Cum. 1995		Cum. 1995
Anthrax Brucellosis Cholera Congenital rubella syndrome Diphtheria <i>Haemophilus influenzae</i> * Hansen Disease Plague Poliomyelitis, Paralytic	69 13 4 875 101 6	Psittacosis Rabies, human Rocky Mountain Spotted Fever Syphilis, congenital, age < 1 year [†] Tetanus Toxic shock syndrome Trichinosis Typhoid fever	50 1 444 280 22 140 24 236

*Of 856 cases of known age, 204 (24%) were reported among children less than 5 years of age. ¹Updated quarterly from reports to the Division of STD Prevention, National Center for Prevention Services. This total through second quarter 1995.

-: no reported cases

Beyond Historical Limits

			-								
Reporting Area	AIDS*	Gonor	rhea	А		В		C/N/	A,NB	Legion	ellosis
	Cum. 1995	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994
UNITED STATES	54,704	257,118	302,033	20,137	18,077	7,289	8,571	3,173	3,124	924	1,176
NEW ENGLAND	2,653 81	4,313 63	6,340 66	211 23	223 20	163 7	259 11	89	116	23 5	58 4
Maine N.H.	77	89	82	8	16	18	20	12	9	1	-
Vt. Mass.	30 1,137	46 2,140	23 2,368	5 88	8 83	1 63	6 150	1 71	12 75	- 14	- 38
R.I. Conn.	192 1,136	400 1,575	355 3,446	25 62	19 77	8 66	6 66	5	20	3 N	16 N
MID. ATLANTIC	14,696	25,277	34,101	1,186	1,278	909	1,113	311	362	146	181
Upstate N.Y. N.Y. City	1,736 7,624	3,846 8,598	8,044 12,893	314 566	438 483	296 279	293 249	167 1	174 1	39 3	44 5
N.J. Pa.	3,575 1,761	3,256 9,577	3,922 9,242	140 166	228 129	190 144	287 284	108 35	157 30	21 83	34 98
E.N. CENTRAL	4,122	56,160	60,894	2,188	1,793	709	882	210	261	247	338
Ohio Ind.	852 429	16,430 6,182	16,437 6,511	1,425 132	652 297	85 179	129 161	8 6	18 8	124 59	159 36
III. Mich.	1,736 825	15,964 13,285	18,586 13,541	217 279	450 215	94 310	230 287	33 163	71 164	13 24	31 59
Wis.	280	4,299	5,819	135	179	41	75	-	-	27	53
W.N. CENTRAL Minn.	1,266 285	14,719 2,133	16,922 2,463	1,417 144	910 168	454 44	499 46	95 2	69 14	90 5	79 2
lowa Mo.	71 564	1,151 8,493	1,094 9,341	50 1,014	47 464	33 316	24 374	11 55	9 18	17 44	28 26
N. Dak. S. Dak.	6 15	20 127	32 166	23 49	5 31	4	2	8 1	1	4	4
Nebr.	84	697	1,030	34	104	22	24	6	11	10	13
Kans. S. ATLANTIC	241 14,155	2,098 75,291	2,796 80,057	103 938	91 936	33 1,043	29 1,568	12 251	16 335	7 167	5 285
Del. Md.	241 2,250	1,670 7,471	1,451 14,100	7 166	21 136	1,010 2 198	12 261	1	1 17	2 27	31 65
D.C.	827	3,483	5,411	19	17	15	40	-	1	4	6
Va. W. Va.	1,082 86	7,962 507	9,983 603	159 17	133 15	89 41	101 31	14 43	20 24	15 4	6 3
N.C. S.C.	816 766	18,048 9,333	20,525 9,929	88 38	100 31	224 37	216 25	46 17	50 7	31 29	19 12
Ga. Fla.	1,784 6,303	11,257 15,560	U 18,055	55 389	26 457	63 374	516 366	15 111	171 44	23 32	100 43
E.S. CENTRAL	1,763	31,492	35,469	1,193	474	624	887	753	716	43	71
Ky. Tenn.	221 709	3,684 10,215	3,736 11,460	32 956	124 212	52 488	66 761	22 729	23 678	10 24	8 36
Ala. Miss.	484 349	12,798 4,795	12,079 8,194	69 136	77 61	84	60	2	15	6 3	12 15
W.S. CENTRAL	4,691	23,326	36,232	3,018	2,340	1,245	957	508	249	12	35
Ark. La.	209 785	2,350 8,475	5,075 9,072	343 94	151 120	36 152	22 133	4 132	7 139	1 2	6 12
Okla. Tex.	206 3,491	1,496 11,005	3,696 18,389	661 1,920	228 1,841	376 681	105 697	323 49	46 57	3 6	11 6
MOUNTAIN	1,716	6,563	7,566	3,006	3,535	586	503	341	348	88	73
Mont. Idaho	17 38	55 93	/1 68	100 244	18 266	19 64	18 67	12 41	10 64	4 2	14 1
Wyo. Colo.	12 523	42 2,176	66 2,632	90 409	23 388	16 93	21 78	137 54	128 58	8 33	4 15
N. Mex. Ariz.	137 545	763 2,522	760 2,474	620 869	857 1,395	223 90	161 53	39 35	43 17	4 9	3 9
Utah Nev.	112 332	131 781	195 1,300	555 119	404 184	54 27	60 45	9 14	15 13	13 15	6 21
PACIFIC	9,642	19,977	24,452	6,980	6,588	1,556	1,903	615	668	108	56
Wash. Oreg.	717 347	1,957 224	2,214 764	593 1,486	850 764	137 62	177 117	152 29	193 32	20	10
Calif.	8,328	16,821	20,224	4,738	4,758	1,335 9	1,573	395	438	83	44
Alaska Hawaii	60 190	544 431	695 555	40 123	176 40	13	12 24	1 38	5	5	2
Guam P.R.	۔ 1,925	58 446	97 382	2 81	22 47	1 452	4 270	- 177	- 133	1	1
V.I. Amer. Samoa	27	6 19	25 25	- 6	3	2	7	-	1	-	-
C.N.M.I.	-	23	25 41	15	6	7	1	-	-	-	-
N: Not notifiable	11.11	navailable	i no rong	orted cases	C		mmonwo	alth of No	rthern Ma	riana Islan	de

TABLE II. Cases of selected notifiable diseases, United States, weeks endingSeptember 30, 1995, and October 1, 1994 (39th Week)

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, National Center for Prevention Services, last update September 28, 1995.

							Measle	es (Rube						
Reporting Area		me ease	Mal	aria	Indig	enous	Impo	orted*	То	otal		gococcal ctions	Mu	mps
	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	1995	Cum. 1995	1995	Cum. 1995	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994
UNITED STATES	6,197	9,264	883	807	-	241	1	24	265	860	2,275	2,083	603	1,097
NEW ENGLAND	1,559	2,185 18	37 5	59 4	-	6	1	2	8	27 5	104 8	100 19	10 4	19
Maine N.H.	16 19	22	1	3	-	-	-	-	-	1	19	8	1	3 4
Vt. Mass.	8 141	14 145	1 12	3 27	-	- 1	-	-1	- 2	3 7	8 37	2 44	- 2	-3
R.I. Conn.	279 1,096	312 1,674	4 14	7 15	-	5	- 1	- 1	5 1	7 4	- 32	- 27	1 2	2 7
MID. ATLANTIC	3,750	5,549	236	160	-	7	-	5	12	212	269	223	93	93
Upstate N.Y. N.Y. City	1,904 153	3,560 14	52 117	43 58	-	1 2	-	- 3	1 5	17 14	82 35	73 28	24 13	27 7
N.J. Pa.	770 923	1,074 901	50 17	35 24	-	4	-	2	6	173 8	74 78	49 73	12 44	13 46
E.N. CENTRAL	64	463	83	88	-	- 7	-	3	- 10	102	315	307	104	186
Ohio Ind.	42 14	32 15	9 14	14 12	-	1	-	-	1	17 1	92 59	88 40	33 4	50 7
III.	3	23	32	40	-	-	-	2	2	56	71	99	31	84
Mich. Wis.	5	5 388	15 13	19 3	-	4 2	-	1	5 2	25 3	56 37	46 34	36	35 10
W.N. CENTRAL Minn.	180 117	214 106	19 4	35 11	-	2	-	-	2	170	149 24	134 12	30 2	56 4
lowa	10	13	1	5	-	-	-	-	-	7	26	18	-	13
Mo. N. Dak.	34	84	6 1	11 1	-	1	-		1	160	61 1	65 1	22 1	34 4
S. Dak. Nebr.	- 1	- 3	2 3	-4	-	-	-	-	-	- 2	5 12	8 10	- 4	- 1
Kans.	18	8	2	3	-	1	-	-	1	1	20	20	1	-
S. ATLANTIC Del.	431 7	648 101	187 1	162 3	-	10	-	1	11	62	409 6	305 5	88	157
Md.	267	209	52	57	-	-	-	1	1	4	30	26 4	20	44
D.C. Va.	1 47	6 116	15 41	12 23	-	-	-	-	-	3	3 51	55	20	38
W. Va. N.C.	22 47	17 69	2 15	- 9	-	-	-	-	-	37 3	8 64	12 42	- 16	3 35
S.C. Ga.	16 12	7 109	1 22	4 29	-	- 2	-	-	- 2	- 3	53 80	21 66	9 8	7 8
Fla.	12	14	38	25	-	8	-	-	8	12	114	74	15	22
E.S. CENTRAL Ky.	40 8	37 21	20 2	29 10	-	-	-	-	-	28	144 46	148 33	13	18
Tenn. Ala.	20 7	10	7	9	-	-	-	-	-	28	37 32	28 57	- 4	6 5
Miss.	5	-	3	9	-	-	-	-	-	-	32 29	30	4 9	5
W.S. CENTRAL Ark.	84 5	97 8	39 3	36 3	-	21 2	-	3	24 2	16 1	281 22	247 38	39 3	196 5
La.	4	1	4	6	-	17	-	1	18	1	39	31	9	23
Okla. Tex.	36 39	54 34	1 31	4 23	U -	- 2	U -	- 2	4	- 14	26 194	24 154	- 27	23 145
MOUNTAIN	7	12	46	25	-	67	-	1	68	163	158	140	24	132
Mont. Idaho	-	- 3	3 1	- 2	-	-	-	-	-	-	2 7	6 15	1 3	-7
Wyo. Colo.	3	3 1	- 22	1 11	-	- 26	-	-	- 26	- 19	7 42	6 27	- 1	2 3
N. Mex. Ariz.	1	3	4	3	-	30 10	-	1	31 10	- 1	31 48	13 48	N 2	N 94
Utah	1	1	5	4	-	-	-	-	-	134	14	18	11	14
Nev. PACIFIC	2 82	1 59	4 216	2 213	-	1 121	-	- 9	1 130	9 80	7 446	7 479	6 202	12 240
Wash.	10 4	1	16 9	23 14	-	16	-	4 1	20 1	3	73 70	75 106	10 N	14 N
Oreg. Calif.	68	52	179	162	-	105	-	3	108	61	291	291	174	207
Alaska Hawaii	-	-	2 10	2 12	-	-	-	- 1	- 1	10 4	8 4	2 5	13 5	3 16
Guam	-	-	-	-	U	-	U	-	-	228	3	-	3	6
P.R. V.I.	-	-	1	4	Ū	11 -	- U	-	11 -	11 -	23	6 -	2 2	2 4
Amer. Samoa C.N.M.I.	-	-	- 1	- 1	U U	-	U U	-	-	- 29	-	-	-	2 2

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending September 30, 1995, and October 1, 1994 (39th Week)

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

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

Reporting Area Pertussis			Rubella		Sypl (Prima Secon	ary &	Tubero	ulosis	Rab Anii			
Reporting Area	1995	Cum. 1995	Cum. 1994	1995	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994
UNITED STATES	90	2,849	2,924	1	117	208	11,213	16,003	14,466	16,433	5,298	5,764
NEW ENGLAND	22	379	341	-	34	128	139	169	380	379	1,195	1,415
Maine N.H.	1 3	27 31	15 66	-	1 1	-	2 1	4 4	12 15	23 13	39 120	- 118
Vt.	-	59	38	-	-	-	-	-	3	6	141	107
Mass. R.I.	8	238 2	190 5	-	7	124 2	46 3	73 12	208 38	194 35	360 260	541 40
Conn.	10	22	27	-	25	2	87	76	104	108	275	609
MID. ATLANTIC	12	250	458	-	12	6	626	1,070	3,011	3,356	995	1,529
Upstate N.Y.	9	126 21	191 84	-	4 7	5	43 287	143 471	368	424	367	1,136
N.Y. City N.J.	-	13	84 12	-	1	- 1	129	172	1,600 573	1,944 577	- 275	214
Pa.	3	90	171	-	-	-	167	284	470	411	353	179
E.N. CENTRAL	10	270	432	-	4	9	1,981	2,377	1,442	1,564	68	52
Ohio Ind.	3 1	111 19	116 47	-	-	-	664 214	905 194	201 176	261 142	10 12	4 12
III.	4	67	89	-	1	1	743	802	711	777	3	17
Mich. Wis.	2	61 12	47 133	-	3	8	228 132	229 247	299 55	338 46	35 8	11 8
W.N. CENTRAL	_	180	133		_	2	596	935	438	432	265	167
Minn.	-	88	51	-	-	-	34	37	100	97	19	14
lowa Mo.	-	43	9 35	-	-	- 2	37 490	49 796	48 166	44 194	90 19	68 18
N. Dak.	-	43	35 4	-	-	-	490	190	3	194	24	10
S. Dak.	-	11	15	-	-	-	-	1	20	21	72	29
Nebr. Kans.	-	8 22	8 10		-	-	9 26	11 40	20 81	16 52	5 36	- 28
S. ATLANTIC	28	276	257	-	26	15	2,864	4,172	2,503	2,975	1,625	1,533
Del.	-	10	2	-	-	-	11	22	42	32	74	44
Md. D.C.	1	28 5	58 7		-	-	137 84	229 172	241 82	240 95	265 11	419 2
Va.	-	15	30	-	-	-	476	614	167	255	320	313
W. Va. N.C.	- 26	- 110	4 58	-	- 1	-	9 862	8 1,292	56 333	61 366	94 366	61 130
S.C.	- 20	20	12	-	1	-	456	622	234	287	100	142
Ga.	- 1	24 64	24	-	1 23	2 13	544 285	635	323	526	204	296
Fla.	I		62		23			578	1,025	1,113	191 222	126
E.S. CENTRAL Ky.	-	253 11	119 58	-	-	-	2,908 156	2,917 156	1,133 232	1,148 242	222 23	156 19
Tenn.	-	204	18	-	-	-	658	796	294	378	72	34
Ala. Miss.	-	35 3	31 12	N	N	N	507 1,587	522 1,443	319 288	314 214	118 9	99 4
W.S. CENTRAL	4	227	151	-	7	13	1,426	3,465	1,743	2,079	527	517
Ark.	-	28	22	-	-	-	82	388	113	204	21	25
La. Okla.	2 U	15 14	10 22	- U	-	- 4	774 54	1,346 123	6 146	11 193	25 31	55 30
Tex.	2	170	97	-	7	9	516	1,608	1,478	1,671	450	407
MOUNTAIN	13	421	377	-	5	5	201	206	464	421	145	123
Mont. Idaho	- 1	3 81	6 44	-	-	-	4	3 1	10 12	9 11	41 3	15 3
Wyo.	-	1	-	-	1	-	4	-	2	7	22	17
Colo. N. Mex.	9 3	77 86	185 20	-	-	-	95 33	107 18	37 64	51 43	9 5	11 6
Ariz.	-	149	20 97	-	3	-	33	39	234	168	45	51
Utah	-	19	23	-	1	4	4	10	24	38	14	12
Nev.	-	5	2	-	-	1	28	28	81	94	6	8
PACIFIC Wash.	1	593 206	657 91	1	29 2	30	472 11	692 29	3,352 181	4,079 207	256 5	272 15
Oreg.	1	27	86	-	1	4	7	30	33	90	-	9
Calif. Alaska	-	319	465	1	23	22	453 1	627 3	2,951 59	3,543 51	247 4	215 33
Hawaii	-	41	15	-	3	4	-	3	128	188	-	-
Guam	U	-	2	U	-	1	5	3	35	62	-	-
P.R. V.I.	- U	12	2	- U	-	-	231 2	242 25	165	150	44	67
Amer. Samoa	U	-	1	U	-	-	-	1	3	4	-	-
C.N.M.I.	U	-	-	U	-	-	4	1	13	25	-	-

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending September 30, 1995, and October 1, 1994 (39th Week)

U: Unavailable -: no reported cases

	All Causes, By Age (Years)				P&I [†]	2&I [†]	All Causes, By Age (Years)						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. New Bedford, Mass. New Haven, Conn. Providence, R.I. Somerville, Mass. Springfield, Mass. Waterbury, Conn.	29 54 5 49 42	389 84 17 15 18 31 18 15 21 19 36 5 36 34	94 33 5 1 7 7 3 3 3 5 9 - 4 4	58 15 2 6 2 1 2 2 7 8 3	13 4 3 - 2 - 2 1 - -	15 5 2 1 - - 2 1 1 - 1 1	31 7 1 3 1 5 6 3	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. E.S. CENTRAL	1,394 178 280 88 121 135 35 89 62 51 156 192 7 815	841 104 157 55 76 79 24 57 46 38 107 92 6 532	301 38 59 16 25 35 4 22 13 7 29 53 -	180 24 51 14 17 17 3 3 4 10 36 1 72	43 6 2 3 2 1 3 3 1 6 10 - 25	29 6 7 1 - 2 3 4 - 1 4 1 4 1 - 17	70 25 1 5 1 2 2 6 3 14 5 55
Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Erie, Pa.§	57 2,426 52 21 113 28 21 42	40 1,629 43 15 80 17 16 35	10 410 4 6 16 4 4 3	5 302 3 - 13 6 1 3	1 49 1 - 4 1 - 1	1 36 - - - -	4 124 4 1 4 2 1 2	Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala. Nashville, Tenn.	139	532 79 53 39 54 126 83 31 67	31 19 20 24 34 17 6 17	20 2 7 2 19 6 3 13	25 6 1 3 2 5 1 1 6	2 2 1 4 5 - 3	55 4 5 7 6 15 5 3 10
Jersey City, N.J. New York City, N.Y. Newark, N.J. Paterson, N.J. Philadelphia, Pa. Pittsburgh, Pa.§ Reading, Pa. Rochester, N.Y. Schenectady, N.Y. Scranton, Pa.§ Syracuse, N.Y. Trenton, N.J. Utica, N.Y. Yonkers, N.Y.	31	17 808 28 U 257 64 11 89 15 15 59 20 16 21	12 212 16 U 71 15 4 13 2 2 12 5 7 2	1 185 13 U 48 1 - 3 5 11 - 1	1 24 2 U 8 2 - 5 - - - - - -	14 U 14 2 - 3 - 1	49 2 20 6 2 15 1 5 4 2 4	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.	1,345 70 46	853 51 34 32 106 52 63 243 39 46 115 21 51	293 8 6 12 42 22 12 87 18 22 39 5 20	141 10 6 25 4 17 38 5 10 13 4 7	25 - 1 9 2 2 5 1 2 2 1 2 2 1	33 1 2 6 2 1 5 3 3 2 5	86 2 3 6 5 7 9 29 5 14 5 1
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Dayton, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Grand Rapids, Mict Indianapolis, Ind. Madison, Wis. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans. Kansas City, Mo.	242 55 119 39 47 37 84 59 733 52 38 30 107	$\begin{array}{c} 1,486\\ 23\\ 33\\ 267\\ 132\\ 93\\ 117\\ 91\\ 129\\ 3117\\ 41\\ 6\\ 399\\ 160\\ 43\\ 93\\ 21\\ 300\\ 32\\ 58\\ 47\\ 519\\ 32\\ 58\\ 47\\ 519\\ 32\\ 32\\ 58\\ 47\\ 60\\ 72\\ 519\\ 32\\ 32\\ 31\\ 31\\ 24\\ 60\\ 72\\ 72\\ 72\\ 72\\ 72\\ 72\\ 72\\ 72\\ 72\\ 72$	457 996 322 427 55 105 20 44 7 122 82 10 105 31 164	213 2 26 200 15 11 3 27 1 22 3 12 2 8 17 59 4 4 3 10	66 1 222 3 7 4 3 10 1 - 1 2 1 2 1 2 5 - 2 5	666 1 222 5 6 2 2 6 2 2 6 2 2 6 2 2 6 2 2 5 6 6 2 2 2 7 5 6 6 2 2 7 5 6 6 2 2 7 5 6 6 2 2 5 6 6 2 2 5 6 6 2 2 5 6 6 2 2 7 5 6 6 2 2 7 6 6 2 2 7 6 6 2 2 7 6 6 2 2 7 7 7 7 7 7 7 7 7 7 7 7 7	124 1 21 10 3 20 8 7 2 - - 2 2 12 2 12 2 12 2 12 3 9 5 6 1 36 4 1 - 6 2	MOUNTAIN Albuquerque, N.M. 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. Pasadena, Calif. Pasadena, Calif. Pasadena, Calif. San Joego, Calif. San Joego, Calif. San Jose, Calif. San Jose, Calif. San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash.	111 138 21 139 30 78 120 1,624 18 90 14 79 81 239 24 124 124 124 124 149	528 87 299 71 89 1,057 84 21 44 86 1,057 9 51 12 2 60 48 143 160 48 143 105 89 113 105 31 74	$165 \\ 26 \\ 9 \\ 24 \\ 36 \\ 5 \\ 23 \\ 16 \\ 314 \\ 3 \\ 23 \\ 23 \\ 24 \\ 314 \\ 3 \\ 23 \\ 24 \\ 314 \\ 49 \\ 7 \\ 28 \\ 49 \\ 25 \\ 38 \\ 25 \\ 38 \\ 25 \\ 38 \\ 25 \\ 30 \\ 6 \\ 15 \\ 15 \\ 15 \\ 15 \\ 15 \\ 15 \\ 15 $	76 10 8 14 10 3 15 4 3 9 164 6 5 9 12 31 6 8 11 22 16 8 11 22 16 4 13 5 5	30 5 1 2 10 6 6 49 3 1 15 6 4 5 3 5 1 3 1 1	14 2 1 2 1 2 1 2 3 3 9 - 8 - 4 5 3 2 7 7 - 1 2 2 7 - 2 1 2 - 2 3 3 9 - 8 - - 4 5 - - 5 - - - - - - - - - - - - -	47 2 3 11 8 2 7 5 7 5 127 5 17 111 6 6 8 15 04 7 5 2
Lincoln, Nebr. Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	33 140 83 115 52 83	27 106 61 82 46 50	4 19 12 15 2 18	1 8 7 12 2 8	- 3 3 6 - 5	1 4 - 2 2	2 13 5 2 2 1		12,027 [¶]				325	261	700

TABLE III. Deaths in 121 U.S. cities,* week ending September 30, 1995 (39th Week)

*Mortality data in this table are voluntarily reported from 121 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.
 ¹Pneumonia and influenza.
 §Because of changes in reporting methods in these 3 Pennsylvania cities, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks.
 ¹Total includes unknown ages.
 U: Unavailable -: no reported cases

Vol. 44 / No. 39

MMWR

Lower Respiratory Tract Illness - Continued

Five of the six case-patients were male; patients ranged in age from 30 to 37 years. Five resided in Berkeley County and one in adjacent Charleston County. All had onset of symptoms during July 19–23 and reportedly had recently used illicit drugs. Five had been hospitalized. Predominant clinical features included nonproductive cough, fever (maximum: 103 F [39.4 C]), pleuritic chest pain, and progressively severe shortness of breath. Laboratory findings included a polymorphonuclear leukocytosis (range: 9100 mm³–23,600 mm³) in all six patients and severe hypoxia (PO₂ range: 49–81) in five. All patients had bibasilar or diffuse pulmonary infiltrates on chest radiographs. Of the five patients who were hospitalized, three developed respiratory failure within 1–4 days of admission and within 7–10 days of onset of symptoms; two died.

Diagnostic studies included sputum gram stain and cultures, blood cultures, serologic tests, and pathologic examination of tissue obtained by open lung biopsy or postmortem. These tests were negative for common bacterial pathogens and for *Mycoplasma* sp, *Chlamydia* sp, *Legionella* sp, *Pneumocystis carinii*, *Mycobacterium tuberculosis*, human immunodeficiency virus, respiratory syncytial virus, cytomegalovirus, adenoviruses, Epstein-Barr virus, and influenza virus. Fungal cultures of sputum from the two deceased patients and open lung biopsy from another were positive for molds believed to be contaminants; final identification is pending. Histopathologic analysis of lung tissue from the two deceased patients indicated diffuse alveolar damage with microemboli and minimal inflammatory cell infiltrate. Tissue from the open lung biopsy of the third patient indicated severe organizing fibrinous pneumonia with bronchiolitis obliterans and diffuse alveolar damage. Analyses of blood (five patients) and lung tissue (three patients) specimens and of samples taken from drug paraphernalia (i.e., homemade "pipes" of one patient) were negative for potential toxins.

Although interviews with patients, relatives, and acquaintances suggested several potential exposures (e.g., rodents and herbicides), the only exposure common to all patients was nonparenteral use of drugs during the week before onset of symptoms. Five reported use of crack cocaine and one reported smoking marijuana. Several of the patients were acquainted; however, investigation has not detected a single event attended by all the patients or a common source for the crack cocaine. The ongoing investigation includes following up all possible drug-related contacts of the patients and continued surveillance.

Reported by: L Lettau, MD, S Miller, MD, D Handshoe, MD, J Chambers, MD, Trident Health District, Charleston; L Bell, MD, E Brenner, MD, J Gibson, MD, State Epidemiologist, South Carolina Dept of Health and Environmental Control. Div of Environmental Health and Hazard Effects, National Center for Environmental Health; Div of Field Epidemiology, Epidemiology Program Office, CDC.

Editorial Note: The clinical features of the cluster of cases of acute lower respiratory illness in South Carolina are similar to those reported previously in crack cocaine users who have sustained inhalational injuries. These cases have been characterized by a variety of clinical syndromes including pulmonary edema, interstitial pneumonitis, obliterative bronchitis, and pulmonary hemorrhage (1–6). However, the cluster of cases in South Carolina is the first known outbreak of acute, severe respiratory tract illnesses associated with crack cocaine. Potential explanations for the cases in South Carolina include an idiosyncratic reaction to crack cocaine or the effects of a contaminant or adulterant introduced during the preparation or smoking of the crack.

Lower Respiratory Tract Illness — Continued

The temporal and geographic clustering of the cases and the similarity of their clinical features suggest a common exposure to a unique yet unidentified toxin or microbiologic agent associated with inhalational drug abuse. Cases of similar severe illnesses should be reported to the Division of Disease Control, South Carolina Department of Health and Environmental Control, telephone (803) 737-4165.

References

- 1. Ettinger NA, Albin RJ. A review of the respiratory effects of smoking cocaine. Am J Med 1989; 87:664–8.
- 2. Meisels IS, Like J. The pulmonary effects of free-base cocaine: a review. Cleve Clin J Med 1993;60:325–9.
- 3. Haim DY, Lippmann ML, Goldberg SK, Walkenstein MD. The pulmonary complications of crack cocaine: a comprehensive review. Chest 1995;107:233–40.
- 4. Forrester JM, Steele AW, Waldron JA, Parsons PE. Crack lung: an acute pulmonary syndrome with a spectrum of clinical and histopathologic findings. Am Rev Respir Dis 1990;142:462–7.
- 5. O'Donnell AE, Mappin FG, Sebo TJ, Tazelaar H. Interstitial pneumonitis associated with "crack" cocaine abuse. Chest 1991;100:1155–7.
- 6. Patel RC, Dutta D, Schonfeld SA. Free-base cocaine use associated with bronchiolitis obliterans organizing pneumonia. Ann Intern Med 1987;107:186–7.

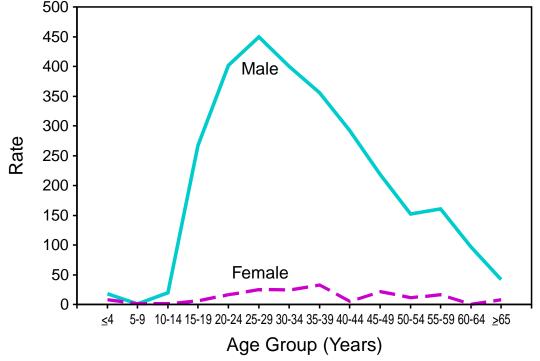
Patterns of Homicide — Cali, Colombia, 1993–1994

In Colombia, as in the United States, homicide occurs disproportionately among urban residents (1,2). Homicide rates in the city of Cali, Colombia (1994 population: 1,776,436), increased fivefold from 1985 through 1992, reaching levels of 100 per 100,000 persons. Because of this increase, in 1992 the city of Cali established the Development, Security, and Peace Program (DESEPAZ) to implement a series of strategies to prevent violence and improve security among the residents of Cali. An important element of this program was the establishment of a surveillance system to enable characterization of patterns and determinants of homicide to provide information to decision makers for formulating policies and programs. This report summarizes findings from this system for January 1993–May 1994.

Data about homicide are collected daily and reviewed weekly by a committee with representatives from the police department, the public health service, the district attorney's office, the ombudsman's office, the National Institute of Legal Medicine and Forensic Sciences, and the department of transportation, under the coordination of an epidemiologist assigned by the mayor's office (3). The list of cases is developed by comparing reports from the different sources represented on the committee. Tabulation and mapping of homicides are distributed weekly to the participating groups and to the news media.

In 1993, a total of 1829 homicides occurred in Cali, representing a crude rate of 104 homicides per 100,000 residents. In 1994, a total of 866 homicides occurred during January–May, compared with 625 for the same 5-month period in 1993. In 1993, age group-specific rates were highest among men aged 25–29 years (450 per 100,000) (Figure 1). Overall, the risk for homicide among males was 16-fold higher than that among females (209 per 100,000 and 13 per 100,000, respectively).

Homicide — *Continued*





*Per 100,000 population.

Homicides were clustered in specific areas of the city: more than one half (54%) occurred in 59 neighborhoods in which 37% of the population resided. Homicide rates varied inversely with the socioeconomic status of the neighborhood of residence of victims and was greatest in three areas: an inner-city location near downtown Cali (rate: 254 per 100,000), a large area of immigrant settlements on the east side (rate: 245 per 100,000), and an area on the west side of the city (rate: 167 per 100,000). In most (89%) cases, homicides occurred in the same neighborhood in which the victim resided.

Of the 2695 total homicides during January 1993–May 1994, for 2461 (91%) no suspect was charged with the murder. For 641 (24%) of all homicides, the suspected perpetrator was identified as a "hit man" (i.e., a hired assassin) (458 [71%]), gang member (91 [14%]), or a relative/acquaintance (62 [10%]). The circumstance was determined for 835 (31%) homicides; of these, the most frequently cited categories were assaults (309 [37%]) and brawls (261 [31%]). Most (2544 [94%]) homicides occurred in streets or other public places; 6% occurred in homes or other residential settings. A total of 2134 (79%) homicides were committed using firearms. During 1994, blood alcohol concentration (BAC) was determined for 98% of the decedents; the BAC was >0.15 g/dL in 23% of the persons.

Homicides occurred more commonly (43%) during weekend days (Friday, Saturday, and Sunday)—especially during weekends coinciding with biweekly paydays. The hour of death was known for 2631 homicides; of these, 51% occurred between 9 p.m. and 6 a.m.

Homicide — Continued

Reported by: V Espitia, MS, Office for the Epidemiology of Violence, Municipality of Cali; R Guerrero, MD, Pan American Health Organization, Cali, Colombia. A Concha, MD, National Center for Injury Surveillance and Research; C Sanchez, MD, VM Cardenas, MD, Colombian Field Epidemiology Training Program, National Institute of Health. Div of Violence Prevention, National Center for Injury Prevention and Control, CDC.

Editorial Note: The data summarized in this report are from the first population-based homicide surveillance system established in Latin America. These findings are being used to develop and evaluate the policies and programs of DESEPAZ, which include efforts to enhance public security by enforcing state and city regulations and using the Mayor's office to issue decrees and laws that further enhance security (4). For example, in response to the relation between homicide and alcohol use, the mayor restricted the hours during which alcoholic beverages could be sold. Similarly, the high proportion of homicides committed with guns prompted institution of prohibitions on carrying guns in public during high-risk weekends, holidays, and election days. In addition, the city initiated efforts to educate members of the community through the news media, schools, and families about resolving conflicts without violence.

To address homicide in areas of the city where the risk has been highest and the socioeconomic status of residents lowest, social and economic development projects are being implemented to provide housing, primary health care, and job opportunities for residents, especially persons in high-risk groups (e.g., prostitutes, street children, and members of youth gangs). Other actions include improving the quality of the relationship between the police and the community, the modernization of the judicial system, and the promotion and protection of human rights. The homicide surveillance data are being used to evaluate the impact of these policies and programs on homicide rates.

Violence is a problem that affects urban areas throughout the Americas. In 1991, the average homicide rate for the 15 largest cities in Colombia (excluding Medellin [rate: 435 per 100,000]) was 61 per 100,000 (1). In comparison, in the United States in 1991, the crude homicide rate was 32 per 100,000 for cities with populations \geq 1 million (2). The Pan American Health Organization with assistance from CDC and other organizations is working to promote the application of public health surveillance, analysis, and evaluation methods to assist countries in this region in reducing the problem.

References

- Gaitan Daza F, Diaz Moreno J. The violence in Colombia: some basic explanations. In: Concha Eastman A, Carrion F, Cobo G, eds. Cities and violence in Latin America. Urban management series, vol 2. Quito, Ecuador: United Nations Urban Management Program, Regional Office for Latin America and the Caribbean, 1994:75–97.
- 2. US Department of Justice. Crime in the United States, 1991. Washington, DC: US Department of Justice, Federal Bureau of Investigation, 1992.
- Eugenia Espitia V, Guerrero Velasco R, Concha Eastman A. Homicide surveillance in Cali, January to December of 1993: the application of epidemiology to public administration. In: Concha Eastman A, Carrion F, Cobo G, eds. Cities and violence in Latin America. Urban management series, vol 2. Quito, Ecuador: United Nations Urban Management Program, Regional Office for Latin America and the Caribbean, 1994:155–66.
- 4. Guerrero Velasco R, Concha Eastman A, Alvarez A, Cobo G, de Roux G, Alzate S. Development, Security, and Peace Program (DESEPAZ): strategies of the Cali mayor's office for addressing the insecurity and the violence. In: Concha Eastman A, Carrion F, Cobo G, eds. Cities and

Homicide — *Continued*

Violence in Latin America. Urban management series, vol 2. Quito, Ecuador: United Nations Urban Management Program, Regional Office for Latin America and the Caribbean, 1994: 119–53.

Notice to Readers

National Surveillance for Infectious Diseases, 1995

On October 13, CDC will release the annual *Summary of Notifiable Diseases, United States, 1994* (1). A notifiable disease traditionally has been considered to be a condition for which regular, frequent, and timely information about individual cases is necessary for the prevention and control of the disease. During 1994, a total of 49 infectious diseases were considered notifiable at the national level and were reported to CDC; 41 were reported weekly, and eight were reported monthly.

During November 30–December 2, 1994, the Council of State and Territorial Epidemiologists (CSTE) and CDC held a conference to review the status of national infectious disease surveillance. At this conference, 10 diseases were proposed for deletion from the list of nationally notifiable diseases: amebiasis, aseptic meningitis, primary encephalitis (except for arboviral encephalitis), postinfectious encephalitis, granuloma inguinale, unspecified hepatitis, leptospirosis, lymphogranuloma venereum, rheumatic fever, and tularemia. In addition, nine diseases were proposed for addition to the list during 1995: genital Chlamydia trachomatis infections, coccidioidomycosis (for regional surveillance), cryptosporidiosis, hantavirus disease, (postdiarrheal) hemolytic uremic syndrome, pediatric infection with human immunodeficiency virus, invasive group A streptococcal infections, streptococcal toxicshock syndrome, and drug-resistant Streptococcus pneumoniae invasive disease. These changes were approved by a vote of the full membership of CSTE in March 1995. However, these conditions are currently not reportable in all states, and the mechanism for reporting may not involve clinicians or consist of weekly reports of individual cases (i.e., traditional notification methods); rather, some may be reported directly by laboratories or in summary form on a monthly basis.

As of October 1, 1995, 52 infectious diseases were designated as notifiable at the national level (Table 1).

Reported by: Council of State and Territorial Epidemiologists. Div of Surveillance and Epidemiology, Epidemiology Program Office, CDC.

Editorial Note: In 1878, Congress authorized the U.S. Marine Hospital Service (the precursor to the Public Health Service [PHS]) to collect morbidity reports on cholera, smallpox, plague, and yellow fever from U.S. consuls overseas; this information was used to implement quarantine measures to prevent the introduction and spread of these diseases into the United States. In 1879, Congress approved appropriations explicitly for the collection and publication of reports of these notifiable diseases. The authority for weekly reporting and publication was expanded by Congress in 1893 to include information from states and municipal authorities. To increase the uniformity of the data, Congress enacted a law in 1902 directing the Surgeon General to provide forms for the collection and compilation of data and for the publication of reports at the national level. Surveillance for Infectious Diseases — Continued

TABLE 1. Infectious diseases designated as notifiable at the national level* — United
States, 1995

Acquired immunodeficiency	Haemophilus influenzae,	Rabies, human
syndrome	invasive disease	Rocky Mountain spotted
Anthrax	Hansen disease (Leprosy)	fever
Botulism [†]	Hantavirus disease [†]	Rubella
Brucellosis	Hepatitis A	Salmonellosis [†]
Chancroid [†]	Hepatitis B	Shigellosis [†]
Chlamydia trachomatis,	Hepatitis, C/non-A, non-B	Streptococcal disease,
genital infections [†]	HIV infection, pediatric [†]	invasive, Group A [†]
Cholera	Legionellosis	Streptococcus pneumoniae,
Coccidioidomycosis [†]	Leprosy	drug-resistant [†]
Congenital rubella syndrome	Lyme disease	Streptococcal toxic-shock
Congenital syphilis	Malaria	syndrome [†]
Cryptosporidiosis [†]	Measles	Syphilis
Diphtheria	Meningococcal disease	Tetanus
Encephalitis, California [†]	Mumps	Toxic-shock syndrome
Encephalitis, eastern equine [†]	Pertussis	Trichinosis
Encephalitis, St. Louis ^{† .}	Plague	Tuberculosis
Encephalitis, western equine [†]	Poliomyelitis	Typhoid fever
Escherichia coli 0157:H7 [†]	Psittacosis	Yellow fever [†]
Gonorrhea	Rabies, animal	

*Although varicella is not a nationally notifiable disease, the Council of State and Territorial Epidemiologists recommends reporting of cases of this disease to CDC.

[†]Not currently published in the weekly tables.

In 1912, state and territorial health authorities—in conjunction with PHS—recommended immediate telegraphic reporting of five infectious diseases and monthly reporting by letter of 10 additional diseases. The first annual summary of *The Notifiable Diseases* in 1912 included reports of 10 diseases from 19 states, the District of Columbia, and Hawaii. By 1928, all states, the District of Columbia, Hawaii, and Puerto Rico were participating in national reporting of nearly 30 specified conditions. At their meeting in 1950, the State and Territorial Health Officers authorized a conference of state and territorial epidemiologists whose purpose was to determine which diseases should be reported to PHS. In 1961, CDC assumed responsibility for the collection and publication of data on nationally notifiable diseases.

The list of nationally notifiable diseases is revised periodically. Diseases may be added to the list as new pathogens emerge and deleted as their incidence declines. Public health officials at state health departments and CDC collaborate in determining which diseases should be nationally notifiable; CSTE, in conjunction with CDC, makes recommendations annually for additions and deletions to the list of nationally notifiable diseases. However, reporting of nationally notifiable diseases to CDC by the states is voluntary. Reporting is currently mandated (by state legislation or regulation) only at the state level. Therefore, the list of diseases that are considered notifiable varies slightly by state. All states generally report the internationally quarantinable diseases (i.e., cholera, plague, and yellow fever) in compliance with the World Health Organization's International Health Regulations.

The Summary of Notifiable Diseases, United States, 1994 contains summary tables of the official statistics for the reported occurrence of nationally notifiable diseases during 1994. Data are presented in tables by month, geographic location, and patient

Surveillance for Infectious Diseases — Continued

age and race/ethnicity and in maps and graphs for many conditions. New features included in this year's annual summary include written highlights of important developments in the reported occurrences of selected nonnotifiable diseases, data from the Public Health Laboratory Information System, and short statements under each map or graph that underscore their important public health messages. Tables presenting historical notifiable diseases data since 1945 and a table on deaths associated with specific notifiable diseases reported to CDC's National Center for Health Statistics also are included.

Reference

1. CDC. Summary of notifiable diseases, United States, 1994. MMWR 1995;43(no. 53).

Monthly Immunization Table

To track progress toward achieving the goals of the Childhood Immunization Initiative (CII), CDC publishes monthly a tabular summary of the number of cases of all diseases preventable by routine childhood vaccination reported during the previous month and year-to-date (provisional data). In addition, the table compares provisional data with final 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).

	No. cases, August	Total o January		No. cases among children aged <5 years [†] January-August			
Disease	1995	1994	1995	1994	1995		
Congenital rubella							
syndrome (CRS)	0	2	4	2	4		
Diphtheria	0	2	0	1	0		
Haemophilus influenzae§	71	763	798	215	185		
Hepatitis B [¶]	746	7486	6491	78	54		
Measles	16	833	255	197	92		
Mumps	33	992	554	157	108		
Pertussis	506	2454	2241	1454	1,301		
Poliomyelitis, paralytic**	0	1	0	0	0		
Rubella	15	202	113	22	16		
Tetanus	5	22	18	0	1		

Number of reported cases of diseases preventable by routine childhood vaccination — United States, August 1994 and 1994–1995*

* Data for 1994 and 1995 are provisional.

[†]For 1994 and 1995, age data were available for \geq 92% cases.

[§]Invasive disease; *H. influenzae* serotype is not routinely reported through the National Electronic Telecommunications System for Surveillance. Of 185 cases among children aged <5 years, serotype was reported for 49 cases, and of those, 28 were type b, the only serotype of *H. influenzae* preventable by vaccination.

¹Because 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.

**One case with onset in July 1994 has been confirmed; this case was vaccine-associated. An additional six suspected cases are under investigation. In 1993, three of 10 suspected cases were confirmed; two of the confirmed cases were vaccine-associated, and one was imported. The imported case occurred in a 2-year-old Nigerian child brought to the United States for care of his paralytic illness; no poliovirus was isolated from the child.

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

Data in the weekly *MMWR* are provisional, based on weekly reports to CDC by state health departments. The reporting week concludes at close of business on Friday; compiled data on a national basis are officially released to the public on the following Friday. Address inquiries about the *MMWR* Series, including material to be considered for publication, to: Editor, *MMWR* Series, Mailstop C-08, CDC, 1600 Clifton Rd., N.E., Atlanta, GA 30333; telephone (404) 332-4555.

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

Director, Centers for Disease Control	Editor, <i>MMWR</i> Series						
and Prevention	Richard A. Goodman, M.D., M.P.H.						
David Satcher, M.D., Ph.D.	Managing Editor, <i>MMWR</i> (weekly)						
Deputy Director, Centers for Disease Control	Karen L. Foster, M.A.						
and Prevention	Writers-Editors, <i>MMWR</i> (weekly)						
Claire V. Broome, M.D.	David C. Johnson						
Director, Epidemiology Program Office	Darlene D. Rumph-Person						
Stephen B. Thacker, M.D., M.Sc.	Caran R. Wilbanks						
☆U.S. Government Printing Office: 1996-733-175/27018 Region IV							