



MORBIDITY AND MORTALITY WEEKLY REPORT

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Epidemiologic Notes and Reports

Lead Intoxication Associated with Chewing Plastic Wire Coating — Ohio

In December 1991, a venous blood lead level (BLL) of 50 μ g/dL was detected in a 46-year-old Ohio man during a routine pre-employment examination. He was referred to a university-based pharmacology and toxicology clinic for further evaluation; clinic physicians investigated the case. Although a repeat BLL obtained 1 month later was 51 μ g/dL, he reported no exposure to known sources of lead during the interim. However, he reported numbness of his fingers and palms, tinnitus, and a possible decrease in his ability to perform basic arithmetical calculations.

A comprehensive occupational and environmental history obtained at the time of the second BLL test revealed no apparent source of his lead exposure. Although he had been employed for approximately 20 years as a microwave technician during military service and while employed at a television station, he reported no history of exposure to lead from soldering or welding. He had no activities or hobbies associated with exposure to lead or lead products, no previous bullet or birdshot wounds, and he denied drinking illicitly distilled alcohol or using lead additives in his car.

His residence was built in 1974 (after lead was banned from use in residential paint)*, and household water was obtained from a well. In January 1992, blood lead testing of family members revealed levels of 5 μ g/dL for his wife and <5 μ g/dL for his 17-year-old child. His only medication was ranitidine[†], which he had used for the previous 1½ years for "indigestion." He reported occasional cigarette smoking.

Although results of a neurologic examination were normal, neuropsychiatric testing on March 13 demonstrated mild memory deficits, as evidenced by abnormalities on verbal and figural memory tests. Because of these abnormalities, beginning March 13, he was treated for 19 days with dimercaptosuccinic acid (DMSA), an oral chelating agent, and on April 4, his BLL had decreased to 13 μ g/dL. However, BLLs on May 15 and July 23 were 49 μ g/dL and 56 μ g/dL, respectively.

^{*16} CFR §1303.2. Ban of lead-containing paint and certain consumer products bearing lead-containing paint.

[†]Ranitidine alters gastric acidity, which theoretically can influence gastrointestinal absorption of lead.

Lead Intoxication — Continued

During a July 1992 follow-up clinic visit, he mentioned that for approximately 20 years he had habitually chewed on the plastic insulation that he stripped off the ends of electrical wires. Samples of the copper wire with white, blue, and yellow plastic insulation were obtained and analyzed for lead content. The clear plastic outer coating (present on all colors of wire) and the copper wire contained no lead; however, the colored coatings contained 10,000–39,000 μg of lead per gram of coating.§ On receipt of these results, he was instructed immediately to discontinue chewing the wire coating.

In January 1993, when his BLL was 24 μ g/dL, he reported subjective improvement in his symptoms; follow-up neuropsychiatric testing is pending.

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Editorial Note: This report likely represents the first documented case of lead poisoning following ingestion of lead as a consequence of chewing on plastic wire coatings. Plastic coatings previously have been associated with lead exposure in the burning of lead-containing plastics during repair of a storage tank (1), the production of plastics (2,3), and the manufacture and use of stabilizers and pigments in the plastics industry (4). Although lead exposure also can occur among workers who burn the plastic coating off copper wire to recycle the copper, lead intoxication by this route has not been reported (5).

Lead compounds may be employed in the production of colored plastics (in which lead chromates are used as pigment) and in the manufacture of polyvinyl chloride (PVC) plastics (in which 2%–5% lead salts [including lead oxides, phthalate, sulfate, or carbonate, depending on the desired quality of the final product] are used as stabilizers). Although environmental regulation has reduced considerably the amount of lead used in the United States in the manufacture of PVC plastics, manufacturers of electrical wire and cable continue to produce PVC stabilized and/or pigmented with lead compounds (6).

More than 573,400 U.S. workers are employed in occupations involving electrical work. Among these workers, potential for excessive exposure to lead may result from inhalation of fumes generated during lead soldering (7). Because the plastic coating from wires is usually removed by mechanical stripping, ingestion of lead from these plastic coatings is probably uncommon. Nonetheless, the findings in this report remind occupational and other health-care providers of the need to be aware of this potential source of lead exposure. In addition, workers should be warned of the potential hazard of chewing plastic coatings or other plastic products that may contain lead.

References

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[§]Samples were analyzed using graphite furnace atomic absorption spectroscopy, following dissolution of the plastic coating in tetrahydrofuran.

Lead Intoxication — Continued

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Current Trends

Arboviral Diseases — United States, 1992

During 1992, health departments from 23 states reported to CDC 45 cases of arboviral encephalitis in humans and 97 in horses. An additional four states reported detection of arboviral activity in bird and mosquito populations. Unlike 1990 and 1991, when three St. Louis encephalitis (SLE) epidemics and an eastern equine encephalitis (EEE) epizootic occurred, during 1992, no focal outbreaks of arboviral disease were reported. This report summarizes information regarding arboviral encephalitis in the United States during 1992.

SLE. During 1992, 14 sporadic SLE cases occurred in Texas (12 cases) and California (two) (1)—a substantial decrease from 1990 and 1991 (247 and 78, respectively), when SLE cases were at their highest level since 1976.

LaCrosse encephalitis (LAC). During 1992, 29 cases of LAC encephalitis were reported from Illinois (seven cases), Ohio (six), West Virginia (six), Wisconsin (four), Minnesota (three), and North Carolina (three). This is the lowest number of LAC cases reported since surveillance began in 1964.

EEE and Western equine encephalitis (WEE). During 1992, Florida and Massachusetts each reported one case of EEE. Because of isolation of EEE virus from *Aedes albopictus* during 1991 in Florida, human case surveillance was intensified at five regional medical centers. From May through September 1992, 357 cerebrospinal fluid samples were collected from persons with symptoms suggestive of meningitis or encephalitis. None had EEE-specific immunoglobin M antibody. In 1992, 88 cases of EEE in horses were reported from Florida (54 cases), Georgia (nine), Virginia (nine), Mississippi (four), South Carolina (four), North Carolina (three), Texas (two), Arkansas (one), Kentucky (one), and Michigan (one). Although no cases of WEE were reported in humans, nine cases of WEE in horses were reported during 1992: Idaho (two cases), Missouri (two), Oklahoma (two), Colorado (one), South Dakota (one), and Utah (one).

Enzootic arbovirus activity. In 1992, 28 states conducted arboviral surveillance using virus isolation or antigen detection in captured mosquitoes or viral-specific antibody assays in sentinel or wild birds. Enzootic arboviral activity was reported from 16 states: EEE (Delaware, Florida, Georgia, Massachusetts, Michigan, New Jersey, North Carolina, Ohio, and South Carolina), SLE (Arizona, California, Illinois, Michigan, and Texas), WEE (Arizona, California, Colorado, Nevada, and Utah), and LAC (Illinois).

Reported by: WG Hlady, MD, Florida Dept of Health and Rehabilitative Svcs. Participating state epidemiologists, veterinarians, and vector-control coordinators. Arbovirus Diseases Br, Div of Vector-Borne Infectious Diseases, National Center for Infectious Diseases, CDC.

Arboviral Diseases — Continued

Editorial Note: An increased number of EEE cases had been anticipated in 1992 for two reasons: 1) in 1991, EEE virus had been isolated from *Aedes albopictus* (2), a more anthropophilic mosquito vector; and 2) in 1991, an EEE epizootic occurred in the Southeast (1). Although arboviral infections are often underreported, the results of intensified surveillance in Florida suggest that human EEE infection did not increase in 1992.

The last nationwide arboviral epidemic (1975 and 1976) resulted in 2194 cases of SLE in 35 states and was preceded by a modest increase in human SLE cases in 1974. Because early recognition of arboviral activity allows for early institution of preventive measures, surveillance of virus activity in mosquito, avian, equine, and human populations has been emphasized.

During 1990 and 1991, moderate increases in arboviral encephalitis cases were noted with outbreaks in Arkansas, Florida, and Texas (3,4). Despite changes in the arboviral surveillance system to encourage a greater number of states to report regularly, only 45 cases of human arboviral encephalitis were reported—the lowest number of cases reported since the early 1960s. Most arboviral encephalitis cases were reported from midwestern states. Serosurveys indicate that arboviral infections have a wide geographic distribution in the United States, and that cases are often underreported.

Because early identification of cases is important in reducing the risk for arboviral disease through vector-control practices and changes in human activity patterns, health-care providers should consider arboviruses in the differential diagnosis of viral meningoencephalitis, obtain appropriate specimens for serologic testing, and promptly report cases to state health departments.

References

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International Notes

Mortality Among Newly Arrived Mozambican Refugees — Zimbabwe and Malawi, 1992

An estimated 1.3 million persons have fled Mozambique since 1986 because of civil war in that country. More than 1 million refugees have sought asylum in Malawi and approximately 230,000 in Zimbabwe (Figure 1); of the combined total, an estimated 130,000 (10%) fled during January–September 1992. The rate of exodus accelerated during 1992 because of a severe drought that affected most of southern Africa. During August–September 1992, the Bureau for Refugee Programs of the U.S. Department of State and CDC, in collaboration with the Office of the United Nations High Commissioner for Refugees, assessed the impact of the drought on the health status of refugees in the region through observations of refugee conditions and examinations

of data in refugee camps in Zimbabwe and Malawi. This report summarizes the findings of the assessment.

In Zimbabwe, most newly arriving refugees were placed in Chambuta camp (in south Zimbabwe); the population in this camp increased from 6700 in January to its capacity of 25,000 in August. In Malawi, refugees were placed in Lisungwe Camp, which opened in November 1991; the population of this camp reached 65,000 by the end of August 1992. From July through September, the number of new arrivals each month in Lisungwe ranged from 6000 to 20,000. Because of limited space in Lisungwe in September, approximately 16,000 Mozambican refugees were detained at border posts and temporary reception centers in other camps in Malawi with inadequate shelter, sanitation, and water.

Crude Mortality

In Chambuta, detailed records on deaths were compiled by health center staff. During August 1–20, 1992, the crude mortality rate (CMR) was 3.5 deaths per 10,000 population per day. Although age-specific data were not available, most deaths were reported anecdotally to have occurred in children aged <5 years. During the first 4 weeks after refugees arrived in camp, daily death rates increased from 7.3 per 10,000 population to 8.2, after which rates were inversely related to duration of stay. However, the CMR for refugees who had resided in the camp for more than 6 months was three times the CMR (0.5 per 10,000 per day) reported by the United Nations Children's Fund (UNICEF) for nondisplaced persons in Mozambique (1).

(Continued on page 475)

FIGURE 1. Location of camps that received refugees — Malawi and Zimbabwe, 1992

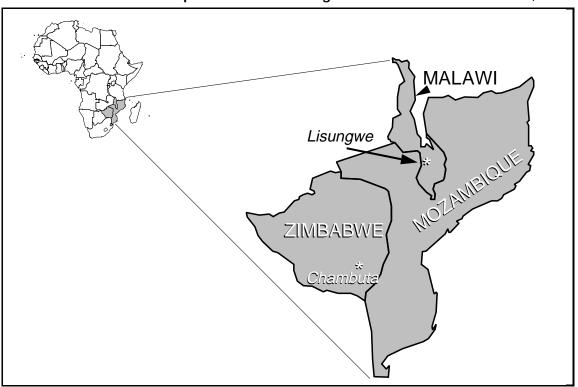


FIGURE I. Notifiable disease reports, comparison of 4-week totals ending June 19, 1993, with historical data — United States

DISEASE	DECREA	SE		INCREASE	CASES CURRENT 4 WEEKS
Aseptic Meningitis			1		470
Encephalitis, Primary					22
Hepatitis A					919
Hepatitis B					834
Hepatitis, Non—A, Non—B					249
Hepatitis, Unspecified					40
Legionellosis					66
Malaria					54
Measles, Total*					31
Meningococcal Infections					126
Mumps					146
Pertussis					240
Rabies, Animal					560
Rubella					7
0.0312	5 0.0625		0.5 1 Log Scale) † D HISTORICAL LIMITS	2	4

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

TABLE I. Summary — cases of specified notifiable diseases, United States, cumulative, week ending June 19, 1993 (24th Week)

	Cum. 1993		Cum. 1993
AIDS* Anthrax Botulism: Foodborne Infant Other Brucellosis Cholera Congenital rubella syndrome Diphtheria Encephalitis, post-infectious Gonorrhea Haemophilus influenzae (invasive disease)† Hansen Disease	51,608 - 6 12 2 36 14 5 - 80 171,571 602 88	Measles: imported indigenous Plague Poliomyelitis, Paralytic [§] Psittacosis Rabies, human Syphilis, primary & secondary Syphilis, congenital, age < 1 year Tetanus Toxic shock syndrome Trichinosis Tuberculosis Tularemia	17 110 3 - 26 - 12,187 - 15 113 8 9,201 42
Leptospirosis Lyme Disease	15 1,952	Typhoid fever Typhus fever, tickborne (RMSF)	153 64

[†]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 thehatched area begins is based on the mean and two standard deviations of these 4-week totals.

^{*}Updated monthly; last update June 5, 1993.

†Of 550 cases of known age, 187 (34%) were reported among children less than 5 years of age.

§No cases of suspected poliomyelitis have been reported in 1993; 4 cases of suspected poliomyelitis were reported in 1992; 6 of the 9 suspected cases with onset in 1991 were confirmed; the confirmed cases were vaccine associated.

TABLE II. Cases of selected notifiable diseases, United States, weeks ending June 19, 1993, and June 13, 1992 (24th Week)

		Ju	ne 19,		_								
	4100+	Aseptic Menin-	Enceph				He	oatitis (\	Legionel-	Lyme			
Reporting Area	AIDS*	gitis	Primary	Post-in- fectious	Gond	rrhea	Α	В	NA,NB	Unspeci- fied	losis	Disease	
	Cum. 1993	Cum. 1993	Cum. 1993	Cum. 1993	Cum. 1993	Cum. 1992	Cum. 1993	Cum. 1993	Cum. 1993	Cum. 1993	Cum. 1993	Cum. 1993	
UNITED STATES	51,608	3,101	232	80	171,571	222,224	9,574	5,355	2,115	280	505	1,952	
NEW ENGLAND Maine	2,166 59	66 9	5 1	5	3,188 39	4,643 39	245 8	231 9	204	7	18 4	255 2	
N.H.	63	8	-	2	30	58	14	51	190	1	2	20	
Vt. Mass.	14 1,188	7 34	1 3	3	14 1,309	13 1,694	3 137	3 125	2 8	6	9	1 46	
R.I. Conn.	104 738	8	-	-	163 1,633	356 2,483	49 34	14 29	4	-	3 -	51 135	
MID. ATLANTIC	11,379	306	9	6	18,882	23,361	574	688	153	4	107	1,395	
Upstate N.Y. N.Y. City	1,938 6,197	119 104	1 1	3	3,841 4,260	4,871 7,914	170 177	189 121	90 1	1 -	32 3	1,036 3	
N.J. Pa.	2,072 1,172	83	- 7	3	3,273 7,508	3,212 7,364	152 75	184 194	43 19	3	15 57	129 227	
E.N. CENTRAL	4,160	411	72	15	32,802	41,597	928	530	349	7	130	16	
Ohio Ind.	662 502	131 49	25 4	3 7	8,850 3,453	12,898 3,905	150 392	111 83	29 5	1	72 21	12 1	
III. Mich.	1,442 1,083	86 135	16 24	5	11,300 6,829	12,989 9,896	269 111	112 219	20 275	2 4	4 25	1 2	
Wis.	471	10	3	-	2,370	1,909	6	5	20	-	8	-	
W.N. CENTRAL Minn.	2,163 431	184 45	10 5	-	7,978 320	12,156 1,369	1,221 205	324 31	93 3	5 4	31 1	35 4	
Iowa Mo.	130 1,270	42 41	1	-	602 4,964	803 6,624	16 792	12 243	4 68	1	5 9	5 7	
N. Dak.	-	5	2	-	23	41	42	-	-	-	1	1	
S. Dak. Nebr.	20 100	7 2	2	-	123 170	82 743	10 109	7	9	-	12	1	
Kans.	212	42	- 42	-	1,776	2,494	47	31	9	-	3	17	
S. ATLANTIC Del.	10,888 208	741 6	43 3	32	48,139 612	71,078 808	590 5	954 67	254 59	35 -	87 6	180 83	
Md. D.C.	1,216 548	68 19	10	-	7,487 2,567	6,729 3,483	84 3	130 14	5	4	22 12	29 2	
Va. W. Va.	731 38	76 6	14 7	3	5,457 265	8,445 423	63	71 18	20 15	11	2	18 2	
N.C.	453	58	8	-	11,260	11,313	30	148	30		12	26	
S.C. Ga.	673 1,562	5 43	1	-	4,710 4,660	5,238 22,472	7 47	18 33	20	1 -	10 12	1	
Fla.	5,459	460	-	29	11,121	12,167	348	455	105	19	10	19	
E.S. CENTRAL Ky.	1,396 161	153 63	9 4	4 4	19,970 2,099	21,742 2,242	116 64	538 45	420 5	1 -	21 8	6 2	
Ténn. Ala.	528 463	21 41	4 1	-	6,096 7,124	6,992 7,278	19 23	438 52	407 3	- 1	10 1	2 2	
Miss.	244	28	-	-	4,651	5,230	10	3	5	-	2	-	
W.S. CENTRAL Ark.	5,311 227	265 14	19	-	20,469 3,893	20,657 3,878	811 26	693 28	96 2	76	14	10 1	
La.	727	23	-	-	5,161	3,175	36	89	35	1	2	-	
Okla. Tex.	423 3,934	1 227	4 15	-	1,719 9,696	2,226 11,378	49 700	109 467	22 37	6 69	8 4	5 4	
MOUNTAIN	2,599	178	11	3 1	4,921	5,634 49	1,945	264 4	146	48	48 5	3	
Mont. Idaho	15 43	6	-	-	22 80	59	53 93	22	-	1	1	-	
Wyo. Colo.	28 868	3 38	3	-	41 1,501	24 2,138	10 469	13 30	45 21	- 28	5 4	2	
N. Mex. Ariz.	212 881	37 63	3	2	444 1,839	419	163 676	111 40	49 9	2 7	3	-	
Utah	185	6	1	-	154	1,874 106	445	20	18	10	7	1	
Nev. PACIFIC	367 11,546	25 797	- 54	- 15	840 15,222	965 21,356	36 3,144	24 1,133	4 400	- 97	14 49	- 52	
Wash. Oreg.	764 502	-	-	-	1,791 904	1,950 712	339 52	96 20	89 8	7	7	1	
Calif.	10,149	752 4	51	15	12,071	18,116	2,313 394	1,002	297 4	88	37	50	
Alaska Hawaii	12 119	41	2 1	-	212 244	339 239	394 46	6 9	2	2	5	1	
Guam P.R.	- 1,561	2 27	-	-	38 209	36 72	2 35	2 172	- 21	1 2	-	-	
V.I. Amer. Samoa	33	-	-	-	55 12	53 20	10	2		-	-	-	
C.N.M.I.	-	2	-	-	41	20 29	-	-	-	1	-	-	

N: Not notifiable

U: Unavailable

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

^{*}Updated monthly; last update June 5, 1993.

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending June 19, 1993, and June 13, 1992 (24th Week)

			Measle	s (Rube	eola)		Menin-		·		•					
Reporting Area	Malaria	Indig	enous	,	orted*	Total	gococcal Infections	Mu	mps	Pertussis			Rubella			
	Cum. 1993	1993	Cum. 1993	1993	Cum. 1993	Cum. 1992	Cum. 1993	1993	Cum. 1993	1993	Cum. 1993	Cum. 1992	1993	Cum. 1993	Cum. 1992	
UNITED STATES	6 424	1	110	_	17	1,900	1,270	16	854	80	1,215	712	2	100	99	
NEW ENGLAND Maine	29 1	1	46	-	4	41	78 5	1	7	41	331 8	66 2	-	1 1	5	
N.H.	4	-	-	-	-	9	11	-	-	27	190	20	-	-	-	
Vt. Mass.	1 10	-	30 7	-	1 2	8	4 41	-	2	-	42 57	33	-	-	-	
R.I. Conn.	2 11	1	9	-	1	20 4	1 16	1	2 3	14	2 32	11	-	-	4 1	
MID. ATLANTIC Upstate N.Y.	80 28	-	6	-	2 1	205 106	158 71	1 1	60 22	-	177 73	73 23	1 1	28 4	11 8	
N.Y. City	24	-	2	-	-	40	19	-	-	-	12	9	-	17	-	
N.J. Pa.	20 8	-	4	-	1 -	54 5	21 47	-	8 30	-	21 71	18 23	-	6 1	2 1	
E.N. CENTRAL Ohio	29 6	-	1	-	-	32 5	170 53	2 1	128 53	5 5	166 108	54 15	-	2 1	7	
Ind.	4	U	-	U	-	19 5	27	ΰ	3 27	Ŭ	24 15	12 9	U	-	- 7	
III. Mich.	14 5	-	1	-	-	2	51 38	1	45	-	16	2	-	1	-	
Wis. W.N. CENTRAL	13	-	1	-	2	1	1 78	-	24	4	3 84	16 49	-	- 1	- 5	
Minn. Iowa	3	-	-	-	-	5 1	2 15	-	7	4	43	15 1	-	-	-	
Mo.	3 2	-	1	-	-	-	30 3	-	12	-	21	20 7	-	1	1	
N. Dak. S. Dak.	2		-		-	-	3	-	4	-	2 1	3	-	-	-	
Nebr. Kans.	1 1	U -	-	U -	2	-	4 21	U -	1	U -	5 11	2 1	U -	-	4	
S. ATLANTIC Del.	122 1	-	20 3	-	3	112 1	257 10	3	275 4	9	121 1	60	-	7 2	7	
Md. D.C.	13 5	-	-	-	2	15	23 4	1	49	5 1	41 2	12	-	1	4	
Va.	8	-	-	-	1	11	20	-	14	1	10	4	-	-	-	
W. Va. N.C.	2 68	-	-	-	-	24	10 44	1	6 157	-	6 20	2 1 <u>4</u>	-	-	-	
S.C. Ga.	3	-	-	-	-	29 -	20 60	-	13 9	-	5 5	7 6	-	-	-	
Fla. E.S. CENTRAL	22 9	-	17 1	-	-	32 432	66 80	1	23 32	2 3	31 50	15 12	-	4	3 1	
Ky. Tenn.	5	-	-	-	-	415	16 16	-	- 9	- 2	3 30	5	-	-	1	
Ala.	2	-	1	-	-	-	29	-	18	1	16	7	-	-	-	
Miss. W.S. CENTRAL	2 11	-	1	-	-	17 975	19 102	- 5	5 121	- 1	1 32	100	-	- 12	6	
Ark. La.	2	-	1	-	-	-	12 24	1	4	-	2	6	-	1	-	
Okla. Tex.	4 5	-	-	-	-	11 964	9 57	4	2 104	1	12 13	13 81	-	1 10	- 6	
MOUNTAIN	12	-	2	-	-	12	110	-	35	2	80	105	-	4	3	
Mont. Idaho	2	-	-	-	-	-	10 7	-	- 5	-	- 15	1 14	-	- 1	- 1	
Wyo. Colo.	- 7	-	2	-	-	1 11	2 15	-	2	2	1 28	21	-	-	-	
N. Mex. Ariz.	3	-	-	-	-	-	3 61	N	N 6	-	19 10	26 37	-	- 1	- 1	
Utah	-	-	-	-	-	-	5 7	-	3 11	-	7	5 1	-	1 1	i	
Nev. PACIFIC	119	-	32	-	6	- 85	237	4	172	- 15	- 174	193	1	45	- 54	
Wash. Oreg.	13 3	-	-	-	-	10	35 19	- N	8 N	1 1	19 3	51 13	-	- 1	6 1	
Calif. Alaska	101	-	22	-	1	42 9	165 10	4	145 5	13 -	142 3	121	-	23 1	34	
Hawaii	2	-	10	-	5	24	8	-	14	-	7	8	1	20	13	
Guam P.R.	1	U -	2 122	U -	-	10 222	1 6	U -	6 1	U -	- 1	9	U -	-	1 -	
V.I. Amer. Samoa	-	-	1	-	-	-	-	-	3	-	- 2	- 6	-	-	-	
C.N.M.I.	-	-	-	-	1	-	-	-	11	-	-	1	-	-	-	

^{*}For measles only, imported cases include both out-of-state and international importations. N: Not notifiable U: Unavailable † International § Out-of-state

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending June 19, 1993, and June 13, 1992 (24th Week)

		philis	Toxic-			Tula-		Typhus Fever	Rabies,
Reporting Area		Secondary)	Shock Syndrome	Tuber	culosis	remia	Typhoid Fever	(Tick-borne) (RMSF)	Animal
	Cum. 1993	Cum. 1992	Cum. 1993	Cum. 1993	Cum. 1992	Cum. 1993	Cum. 1993	Cum. 1993	Cum. 1993
UNITED STATES	12,187	15,814	113	9,201	9,408	42	153	64	3,664
NEW ENGLAND	184 2	309	7 1	193 7	148 13	-	13	2	609
Maine N.H.	21	23	2	4	-	-	-	-	29
Vt. Mass.	1 83	1 149	3	3 116	3 64	-	11	2	15 217
R.I. Conn.	7 70	16 120	1	28 35	13 55	-	2	-	348
MID. ATLANTIC	1,138	2,220	22	2,034	2,278	-	44	4	1,332
Upstate N.Y. N.Y. City	103 541	183 1,214	12 1	189 1,226	298 1,319	-	9 26	1 -	989 -
N.J. Pa.	167 327	316 507	9	318 301	383 278	-	6 3	2 1	198 145
E.N. CENTRAL	1,931	2,291	36	968	922	3	14	5	33
Ohio Ind.	554 168	334 111	15 1	140 100	149 80	1 1	5 1	4	3
III. Mich.	729 297	1,021 457	5 15	487 202	460 197	- 1	4 4	1	4 2
Wis.	183	368	-	39	36	-	-	-	24
W.N. CENTRAL Minn.	747 14	637 42	8 2	203 26	219 50	12 -	2	6	169 21
Iowa Mo.	32 619	20 476	4	18 113	21 93	3	2	4	31 5
N. Dak.	-	1	-	2	3	-	-	-	36
S. Dak. Nebr.	1 7	- 17	-	9 8	14 13	7 -	-	2	19 2
Kans.	74	81	2	27	25	2	-	-	55
S. ATLANTIC Del.	3,307 63	4,421 109	12 1	1,589 18	1,749 23	1 -	18 1	21 1	980 75
Md. D.C.	177 184	342 205	-	173 80	120 57	-	3 -	1 -	290 6
Va. W. Va.	304 3	378 9	2	176 40	125 29	-	1	2	188 40
N.C. S.C.	913	1,079	3	212 190	228	-	-	11 1	37
Ga.	516 569	594 903	- -	352	188 397	-	1	1	80 222
Fla. E.S. CENTRAL	578 1,718	802 2,078	6 4	348 645	582 683	1 3	12 2	4 6	42 45
Ky.	143	66	2	176	180	-	-	3	7
Tenn. Ala.	499 392	561 837	1 1	144 219	164 192	2 1	2	1 -	38
Miss.	684	614	-	106	147	-	-	2	-
W.S. CENTRAL Ark.	2,594 451	2,674 415	1 -	943 82	915 71	17 10	2	18 -	294 15
La. Okla.	1,105 177	1,173 116	- 1	- 151	87 62	4	1	- 18	- 55
Tex.	861	970	-	710	695	3	1	-	224
MOUNTAIN Mont.	105 1	195 3	7 -	200 5	241	1 -	4	2	45 9
ldaho Wyo.	4	1 1	1	6 1	12	- 1	-	2	6
Colo. N. Mex.	31 17	28 19	1	8 18	17 39	-	3	-	1 3
Ariz.	45	97	1	108	111	-	1	-	25
Utah Nev.	2 5	5 41	3 1	11 43	33 29	-	-	-	1
PACIFIC	463	989	16	2,426	2,253	5	54	-	157
Wash. Oreg.	25 47	49 23	2	118 50	137 46	1 2	4	-	-
Calif. Alaska	387 2	910 3	14 -	2,118 19	1,924 36	2	48	-	141 16
Hawaii	2	4	-	121	110	-	2	-	-
Guam P.R.	1 258	2 130	-	28 64	34 120	-	-	-	- 22
V.I. Amer. Samoa	26	28	-	2	3	-	-	-	
C.N.M.I.	2	4	-	16	14	-	-	-	-

U: Unavailable

TABLE III. Deaths in 121 U.S. cities,* week ending June 19, 1993 (24th Week)

All Causes, By Age (Years) Post All Causes, By Age (Years) Post All Causes, By Age (Years) Post															
Daniel Marie		All Cau	ses, By	/ Age (\	'ears)		P&I	B	All Causes, By Age (Years)						
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. Springfield, Mass. Springfield, Mass. Materbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J.	39 44 8 42 27 54 2,716 55 25 100 30	384 84 222 8 23 32 18 11 33 36 6 6 28 23 40 1,811 19 76	7 2 6 8 3 5 2 10 4 1 9 1 6 4 92 9 4 1 16 8 8 8 8 9 16 16 16 16 16 16 16 16 16 16 16 16 16	51 12 5 1 2 9 1 1 1 4 4 - 3 1 7 259 4 - 3	11 2 2 - 1 - - 4 - - 1 1 69 - 2 3 2	21 14 1 - 1 - 1 2 1 - 81 1 - 2 3	43 17 4 2 1 1 2 2 1 5 2 6 115 2	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 Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala.	143 U 7 773 129 54 96 66 204 51	579 U 116 63 66 67 42 41 34 43 103 U 4 49 77 38 63 39 133 34	158 U 32 23 25 18 7 12 6 7 27 U 1 153 23 10 21 15 39 13	90 U 27 4 12 16 11 7 1 3 8 U 1 64 15 6 7 6 7 6 15 3	24 U 4 4 4 1 1 1 1 2 3 3 U	20 U 4 1 3 2 2 4 1 2 U 1 31 8 3 2 11	47 U 18 4 2 - 6 3 2 4 8 U - 50 3 4 5 3 22 4
Elizabeth, N.J. Erie, Pa.§ 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. E.N. CENTRAL	72 25 494 97 10 127 29 27 110 34 28	7 36 36 829 36 17 339 69 8 91 23 20 82 21 25 20	3 6 15 247 16 3 89 16 - 24 4 5 14 10 2 1	3 6 9 157 3 - 44 5 2 7 2 - 7 3 1 2	- 1 29 8 3 14 2 - 4 - 1 - -	1 45 8 -7 5 -1 1 -7 	1 47 5 27 6 - 5 - 1 9 3	Montgomery, Ala. Nashville, Tenn. W.S. CENTRAL Austin, Tex. Baton Rouge, La. Corpus Christi, Tex. Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La. San Antonio, Tex. Shreveport, La. Tulsa, Okla. MOUNTAIN	51 122 1,354 72 30 55 200 44 76 367 67 126 183 36 98	36 79 849 51 21 37 116 28 56 212 36 82 118 25 67 535	9 23 247 17 3 11 33 10 7 76 13 19 37 5 16	3 9 169 4 4 3 31 3 8 56 10 17 4 12 64	2 5 58 1 2 11 1 5 17 3 8 8 1 1	1 6 31 1 2 9 2 6 5 3 1 2	2 7 66 5 1 4 4 25 4 12 3 1
Akron, Ohio Canton, Ohio Canton, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Gary, Ind. Grand Rapids, Micl Indianapolis, Ind. Madison, Wis. Milwaukee, Wis. Peoria, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans. Kansas City, Kans. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	60 40 274 115 149 126 240 43 655 13 45 34 134 43 50 52 104 62 820 94 20 22 134 20	1,261 34 122 76 92 96 87 149 37 51 18 30 105 27 102 31 33 50 598 61 11 100 156 61 90 45 36	13 58 58 22 28 22 25 46 28 27 43 31 19 12 7 6 18 17 7 34 16 17 13	52 7 17 10 8 30 2 2 18 3 10 - 8 3 6 1 2 6 3 8 4 9 2 6	36 33 4 33 26 -21 101 -116 2 174 113 -21 -14	4 11 67 78 87 74 99 22 11 	91	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. Los Angeles, Calif. Pasadena, Calif. Portland, Oreg. Sacramento, Calif. San Diego, Calif. San Francisco, Cali San Jose, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash. TOTAL	73 97 149 30 162 29 1 101 133 1,792 14 78 27 98 84 379 44 129 130 186	44 26 63 93 20 103 1,178 103 1,178 50 22 67 51 245 26 27 85 133 82 126 25 73 34 52	14 10 15 35 9 33 9 19 20 309 3 12 2 18 64 8 15 26 21 31 45 4 40 9	8 5 10 10 - 17 1 9 4 201 - 10 1 8 9 49 5 11 10 23 34 19 4 12 4	7 1 4 7 1 6 - 4 3 55 2 1 1 1 1 4 2 2 6 4 4 10 6 6 7 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	2 5 4 - 4 3 3 44 1 1 4 5 3 3 3 4 4 3 5 6 6 1 - 2 2 2 1 3 3 4 0	1 2 12 7 4 7 11 9 99 3 1 5 166 13 3 8 7 17 2 14 1 5 3 3 1 624

^{*}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

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.

Total includes unknown ages.

U: Unavailable.

In Lisungwe, mortality data were collected by Malawian health surveillance assistants and compiled by Médecins Sans Frontières (MSF)/France, a private voluntary organization. From January through September 1992, the average daily CMR ranged from 1.0 to 3.6 per 10,000 population. For children aged <5 years, the daily death rate peaked in June at 5.0 per 10,000 population. CMRs in Lisungwe were compared with those in Chifunga, a neighboring camp with comparable environmental conditions and a similar surveillance system but that had not received new arrivals during 1992. From January through September, the monthly CMR in Lisungwe was 4.5 times higher than that in Chifunga.

Cause-Specific Mortality

In both Chambuta and Lisungwe, diarrhea (including cholera-associated), dehydration, malnutrition, and measles accounted for 75% of all reported deaths. In both camps, diarrhea-specific death rates were substantial (1.5 and 1.6 per 10,000 per day in Chambuta and Lisungwe, respectively), and coverage rates for household latrines were low: in August 1992, latrines were present in approximately 22% of households in Chambuta and 13% of households in Lisungwe. The daily measles-specific death rate was higher in Chambuta (0.9 per 10,000) than in Lisungwe (0.1 per 10,000).

Prevention Effectiveness

The costs were determined for programs to prevent deaths associated with measles and diarrheal disease in Chambuta and Lisungwe. The cost for measles vaccine provided by UNICEF and administered using disposable syringes was 30¢ U.S. per 0.5-cc dose delivered (2). Assuming that 8170 children aged <15 years arrived during January–July 1992*, the estimated cost of vaccinating all eligible children in Chambuta (new arrivals aged 6 months–15 years and children reaching the age of 6 months while in the camp) during January–September 1992 would have been \$2451, plus \$708 for the cost of two full-time health workers to administer vaccine. During June and July, measles caused 113 deaths in this largely unvaccinated population. Assuming a vaccination rate of 90% and a two-dose schedule for children aged <9 months (resulting in a vaccine efficacy of 85%), the cost of averting 86 of the 113 measles deaths would have been approximately \$37 per death.

The presence of a latrine in the residential setting reduces diarrhea-associated morbidity and mortality by approximately 36% (3). In Malawian camps, the cost of an installed latrine, using refugee-donated labor, is \$8. Thus, the cost of providing a latrine to each household in Lisungwe from January through August would have been \$54,309 and would have averted 54 deaths and 1408 (36%) of 3911 reported episodes, an investment of approximately \$1004 for each death averted and \$38 for each diarrheal episode averted. Assuming that the CMR remained constant through 1992 (based on the mean January–August CMR), then declined to 0.5 per 10,000 per day

^{*}There were approximately 19,000 new arrivals during this period. Based on the demographics of a neighboring camp (Tongo Garra) for which information was available, an estimated 43% of the population would be <15 years of age on arrival and >6 months of age by July 1992.

of the population would be <15 years of age on arrival and >6 months of age by July 1992.

†The measles-specific death rate during July 25–August 13, 1992, was 0.9 per 10,000 per day, equivalent to 2.7 per 1000 per month. The mean camp population was assumed to be 19,000 in June and 23,000 in July, based on camp administrative data.

[§]Based on estimated mid-month populations and reported CMRs. Based on June data, it was assumed that 45% of all deaths were from some form of diarrheal disease. None of the malnutrition-associated deaths were assumed to be preventable through sanitation. Of the estimated 7803 families in Lisungwe at the end of August, 6789 (87%) did not have a latrine.

and that the fraction of deaths attributable to diarrhea remained constant over time, the cost per death averted would be \$85 over the 5-year expected duration of the latrine[¶]. This analysis does not consider the other social and health-related benefits associated with latrine availability (3,4).

In both Zimbabwe and Malawi, the severe drought diminished food supplies available for established resident populations and strained medical and social programs for citizens of both countries. Because of the problems these conditions posed for the Malawian and Zimbabwean governments and for international and nongovernmental relief organizations, recommended measures included 1) accelerating efforts to ensure that every child aged 6 months–15 years is vaccinated against measles on arrival in a camp; 2) increasing resources for family latrine construction; and 3) providing refugees in reception centers with adequate soap, water, buckets, latrines, and shelter.

Reported by: Office of the United Nations High Commissioner for Refugees; Regional Medical Office, Ministry of Health; Médecins Sans Frontières, Blantyre, Malawi. Office of the United Nations High Commissioner for Refugees; Ministry of Health, Harare, Zimbabwe. Bur for Refugee Programs, Washington, DC. Technical Support Div, International Health Program Office, CDC.

Editorial Note: In Africa, an estimated 5 million refugees have fled war and civil conflict in their homelands. In addition, more than 10 million persons are "internally displaced" in countries such as Liberia, Mozambique, Somalia, and Sudan. The high death rates and the major causes of death among refugees newly arrived from Mozambique are consistent with rates reported for other refugee populations in Africa during the early phase of displacement (5).

Diarrheal diseases and measles are particular health risks for refugee populations in Africa. Enteric pathogens may be spread in refugee camps because of exposure to human excrement resulting from insufficient availability of latrines, water supplies, and other sanitation resources (i.e, buckets and soap). In addition, the crowded conditions of refugee camps may promote the transmission of measles and other contagious diseases (6).

The prompt and complete vaccination of susceptible children against measles may be difficult in the setting of massive influxes of new refugees. For example, in Chambuta, many new arrivals may not have been screened or vaccinated because camp health staff were often overwhelmed by such influxes and could not arrange for vaccination coverage. In Lisungwe, most new arrivals aged 6 months–12 years were vaccinated against measles, but deaths may have occurred among persons who had been infected in Mozambique and had entered the camp while already incubating measles.

Cost estimates in this report indicate that targeting prevention efforts to refugee populations can be highly cost effective. In the camps in Malawi and Zimbabwe, the estimated cost per death averted was 10–100 times less than World Bank estimates for averting measles and diarrhea-associated deaths through country-wide programs (7). To ensure that cost-effective services can be readily provided, even during fluctuating and acute emergencies, refugee health programs should incorporate detailed contin-

This assumes that 1) the mean January–September CMR of 0.7 per 10,000 per day continued through the end of 1992; and 2) the 45,000 efugees who had arrived through August (and who were provided with latrines) had a constant CMR of 0.5 per 10,000 per day during the years 1993–1996, based on the CMR reported in the nearby stableChifunga camp during 1992 and among nonrefugee Mozambicans in 1989 (1).

gency plans and emphasize the importance of basic preventive services, such as those described in this report (i.e., vaccination programs and latrine construction).

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Emerging Infectious Diseases

Update: Outbreak of Hantavirus Infection — Southwestern United States, 1993

Since May 1993, the New Mexico Department of Health, the Arizona Department of Health, the Colorado Department of Health, the Utah Department of Health, the Indian Health Service, and CDC, with the assistance of the Navajo Nation Division of Health, have been investigating an outbreak of illness associated with hantavirus infection (1,2). This report updates information regarding the relation between illness and infection with a previously unrecognized hantavirus.

Through June 21, laboratory evidence of hantavirus infection had been confirmed in 12 patients meeting the case definition (2); of these, nine (75%) persons have died. Of the 12 cases, nine occurred in New Mexico, two in Arizona, and one in Colorado. Ten (83%) cases occurred in persons aged 20–40 years. Similar illnesses in an additional 20 persons, eight of whom died, are being investigated for possible hantavirus infection. As of June 21, cases of acute illness associated with hantavirus infection had been documented only in persons residing in Arizona, Colorado, and New Mexico.

The laboratory evidence of hantavirus infection in the 12 case-patients includes demonstration of antibody to hantavirus antigens (eight case-patients), immunohistochemical evidence of hantavirus antigen in autopsy specimens (five case-patients), and amplification of hantavirus-specific RNA sequences by polymerase chain reaction (PCR) performed on RNA extracted from autopsy specimens (three case-patients). Hantavirus-related antigens were immunohistochemically detected in formalin-fixed lung and kidney tissue using a monoclonal antibody that cross-reacts with conserved hantavirus nucleoprotein epitopes (3). Immunostaining did not occur when a battery of other hantavirus-specific monoclonal and polyclonal antibodies was used for case-

Hantavirus Infection — Continued

patients. Immunostaining did not occur with any of the monoclonal antibodies for tissue from seven persons who died from other illnesses.

Since June 6, 191 animals of 12 species have been collected from peridomestic settings in areas where cases have occurred and tested for evidence of hantavirus antibodies at CDC. Hantavirus antibodies were present in 32 (30%) of 107 deer mice (*Peromyscus maniculatus*), one (9%) of 11 piñon mice (*P. truei*), and one (2%) of 48 chipmunks (*Eutamias dorsalis*).

Hantavirus sequences from nine (75%) of 12 antibody-positive *Peromyscus* rodents have been amplified using PCR. Nucleotide sequence analysis of amplified DNA products from three PCR-positive humans and six PCR-positive *Peromyscus* rodents are closely related and provide a direct genetic link between the hantavirus sequence in the rodents and in the human case-patients.

Reported by: F Koster, MD, H Levy, MD, G Mertz, MD, A Qushing, MD, S Young, PhD, K Foucar, MD, J McLaughlin, PhD, B Bryt, MD, Univ of New Mexico School of Medicine, T Merlin, MD, Lovelace Medical Center, Albuquerque; R Zumwalt, MD, P McFeeley, MD, K Nolte, MD, New Mexico Office of the Medical Investigator; MJ Burkhardt, MPH, Secretary of Health, N Kalishman, MD, M Gallaher, MD, R Voorhees, MD, M Samuel, DrPH, M Tanuz, G Simpson, MD, L Hughes, PhD, E Umland, MD, G Oty, MS, L Nims, MS, CM Sewell, DrPH, State Epidemiologist, New Mexico Dept of Health. R Levinson, MD, F Yerger, MD, B Allan, MD, Scottsdale; P Rubin, Phoenix; L Sands, DO, K Komatsu, MPH, C Kioski, MPH, K Fleming, MA, J Doll, PhD, C Levy, MS, TM Fink, P Murphy, B England, MD, M Smolinski, MD, B Erickson, PhD, WSlanta, G Gellert, MD, State Epidemiologist, Arizona Dept of Health Svcs. P Shillam, MSPH, RE Hoffman, MD, State Epidemiologist, Colorado Dept of Health. S Lanser, MPH, CR Nichols, MPA, State Epidemiologist, Utah Dept of Health. L Hubbard-Pourier, MPH, Div of Health, Navajo Nation, Window Rock, Arizona. J Cheek, MD, A Craig, MD, R Haskins, MPH, B Muneta, MD, B Tempest, MD, M Carroll, MD, LA Shands, MPH, JP Sarisky, MPH, RE Turner, L White, P Bohan, MS, Indian Health Svc. Div of Field Epidemiology, Epidemiology Program Office; National Center for Environmental Health; Div of Bacterial and Mycotic Diseases, Div of Vector-Borne Infectious Diseases, Scientific Resources Program, and Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases, CDC.

Editorial Note: The patterns of cross-reactivity in the human convalescent and rodent serum specimens, the pattern of immunohistochemical reactivity, and the clinical syndrome in which adult respiratory distress syndrome is a prominent feature of the disease (1,2) suggest that a previously unrecognized hantavirus is responsible for this outbreak. Additional studies of sequences from the viral genome and studies of the virus, once it is isolated, will be necessary for further characterization of the agent.

The high prevalence of hantavirus antibodies in the deer mice and the similarity of PCR products in the deer mice and human case-patients suggest that this species may be involved in hantavirus transmission to humans. Studies of other rodent species have been initiated. *P. maniculatus* is distributed in all parts of the United States, except in the Southeast (4). Although serologic evidence of hantavirus infection was detected during 1985 in 11 (5%) of 218 *P. maniculatus* rodents collected from California, Colorado, and New Mexico (5), additional studies are needed to determine the current distribution of hantavirus infection in *Peromyscus* species in the United States.

Previously described transmission of hantavirus infection has been associated with exposure to rodent excreta and saliva; evidence thus far suggests rodents also are likely the primary source of infection for the hantavirus associated with this outbreak. Reports concerning previously identified hantaviruses have not documented personto-person transmission of these viruses, nor has there been evidence of

Hantavirus Infection — Continued

person-to-person transmission in the current outbreak. Nonetheless, an investigation of contacts of case-patients, including health-care workers, has been initiated along with other studies of risk factors for infection in this outbreak.

The findings implicating a hantavirus in this outbreak and knowledge regarding modes of hantavirus transmission support the previous recommendation that restriction of travel to areas affected by this outbreak is not considered necessary. However, activities that may disrupt rodent burrows or result in contact with rodents or aerosolization of rodent excreta should be avoided (1).

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Notice to Readers

Availability of Streptomycin and Para-Aminosalicylic Acid — United States

Since April 1992, CDC has distributed streptomycin to more than 1000 patients with active tuberculosis under an Investigational New Drug (IND) agreement until licensed, domestic production of streptomycin could be reestablished in the United States. In April 1993, the Food and Drug Administration issued a license allowing Pfizer Inc. to produce and distribute streptomycin. Beginning July 6, 1993, CDC will no longer accept new requests from clinicians to place their patients on streptomycin. Such requests should be directed to Richard Vastola, Roerig Streptomycin Program, Pfizer Pharmaceuticals, Inc., 235 E. 42nd Street, New York, NY 10017; telephone (800) 254-4445. CDC will continue to resupply any patients enrolled in the IND protocol before July 6, 1993, until they have completed their course of streptomycin therapy. Until further notice, CDC will continue to supply para-aminosalicylic acid under a separate IND agreement.

Additional information concerning streptomycin or para-aminosalicylic acid is available from CDC's Drug Service, Scientific Resources Program, National Center for Infectious Diseases, telephone (404) 639-3670.

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