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MORBIDITY AND MORTALITY WEEKLY REPORT

Perspectives in Disease Prevention and Health Promotion

Final Results: Medicare Influenza Vaccine Demonstration — Selected States, 1988–1992

Pneumonia and influenza (P&I) are the sixth leading cause of death in the United States (1), and persons aged \geq 65 years and persons with chronic conditions (e.g., lung or heart disease, diabetes, or cancer) are at greatest risk for P&I. During major epidemics, hospitalization rates for persons at highest risk may increase twofold to fivefold (2). However, only 30% of persons aged \geq 65 years responding to CDC's National Health Interview Survey for 1989 reported having received the influenza vaccine during the previous year (3). In 1988, the Health Care Financing Administration (HCFA) and CDC began a congressionally mandated 4-year demonstration project to evaluate the cost-effectiveness to Medicare of providing influenza vaccine to Medicare beneficiaries. This report presents final results of the Medicare Influenza Vaccine Demonstration conducted during 1988–1992.

Using intervention and comparison areas in Arizona, Illinois, Massachusetts, Michigan, New York, North Carolina, Ohio, Pennsylvania, and Texas and the entire state of Oklahoma (total Medicare population: approximately 2 million), the demonstration sought to 1) increase the provision of annual influenza vaccination among Medicare beneficiaries and 2) measure the accrued benefits of vaccination in terms of reduced morbidity and mortality and the difference in the cost to Medicare of health services use. Levels of vaccination coverage were assessed at baseline and annually at all sites. The cost-effectiveness indices were calculated using morbidity and mortality data from the demonstration and published studies and compared with cost-effectiveness of other Medicare benefits.

In intervention areas, influenza vaccine was supplied without cost to Medicare providers by local health departments using computerized vaccine monitoring and distribution systems. Providers were reimbursed for administration of vaccine. Before the 1990–91 and 1991–92 influenza seasons, the HCFA sent letters to all Medicare beneficiaries living in the intervention areas urging them to be vaccinated. The letters contained specific program information and a local telephone number for obtaining information. In addition, intervention sites undertook varied activities directed to both providers and patients to promote and distribute vaccine to Medicare beneficiaries (*4*).

Vaccination Coverage

The number of doses of vaccine administered during the 4-year demonstration and the percentage of the Medicare population vaccinated in the intervention areas increased from 477,316 (26%) during 1989–90 (the first full year of the project) to 995,884 (51%) during 1991–92. Because some Medicare beneficiaries received influenza vaccines from sources not reimbursed by Medicare, annual surveys were conducted to accurately estimate vaccine coverage in each intervention and comparison site. For 1991–92, the overall vaccine coverage estimate for the 10 intervention sites was 59%, compared with 46% overall vaccine coverage in the comparison sites with no enhanced vaccine delivery or promotion activities. Four intervention sites exceeded 60% vaccination coverage. The increase in influenza vaccination coverage in comparison sites was approximately the same as that in the rest of the United States during this period (CDC, unpublished data, 1993).

Vaccine Effectiveness

Three case-control studies of influenza vaccine effectiveness in preventing hospitalization for pneumonia were conducted during the demonstration. In aggregate, these studies estimated that influenza vaccine was 31%–45% effective in preventing hospitalization for any pneumonia during the 1989–90, 1990–91, and 1991–92 influenza seasons (5–7; HCFA, unpublished data, 1993).

Cost-Effectiveness

Simulation models were used to calculate Medicare hospital payment savings by incorporating a range of vaccination rates (from 35% to 60% or an increase from the 30% baseline rate of 5%–30%) and a range of influenza vaccine effectiveness estimates in reducing pneumonia hospitalizations and deaths (from 5% to 70%). Total net costs to Medicare were calculated by subtracting savings in hospital payments from

Vaccine Demonstration — Continued

mated cost of a year of life gained through cervical cancer screening is \$1600-\$2900 (9).

Because of these generally favorable results, influenza vaccine was made a covered benefit for all Medicare part B beneficiaries on May 1, 1993.

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Editorial Note: The Medicare Influenza Vaccine Demonstration increased annual influenza vaccine coverage and measured both health and economic benefits of influenza vaccine for Medicare. The perspective of the payer used in this study was important in securing coverage for this benefit; however, it differs from cost-effectiveness studies of prevention strategies that usually use a societal perspective and include all direct costs, not just those of the payer. In this study, only the costs paid by Medicare were included. Other costs, such as those incurred by patients for travel or by providers for patient's visits or vaccine administration above the amount paid by Medicare, were not included.

In the last year of the demonstration, influenza vaccination levels exceeded the national health objective for the year 2000 of 60% vaccine coverage among noninstitutionalized persons aged \geq 65 years (objective 20.11) (*10*) in four of 10 intervention sites and overall vaccination levels in the demonstration (59%) nearly reached this objective. Vaccination rates were well beyond the rate of 40% shown to incur zero net

	Cost per		Basis for cal savings per be			
Category	beneficiary	Category	Severe season	Mild season		
Vaccine Administration and	\$0.80	Cost per P&I [†] admission	\$5308.00	\$5308.00		
claims processing	1.15	No. P&I admissions	0.016	0.015		
Distribution	0.28	Effectiveness	40%	20%		
Outreach Adverse medical	0.20	Vaccination rate above baseline	10%	10%		
outcomes	<0.01	Probability of				
Total	\$2.43	severe/mild season	40%	60%		
		Total savings in hospital payments (10-year annual avera	ge)[§] \$1.37	\$0.95		
		Total savings per beneficiary [¶]	\$	2.32		

TABLE 1. Cost to Medicare of influenza vaccine delivery and savings to Medicare, based
on severe and mild influenza seasons* — Medicare Influenza Vaccine Demonstration,
1988–1992

*A severe influenza season was defined as one with pneumonia and influenza (P&I) morbidity and mortality substantially above expected thresholds; a mild season was defined as one in which P&I morbidity and mortality did not exceed expected thresholds.

[†]Pneumonia and influenza.

[§]Savings are calculated as the product of the cost per P&I admission, the number of P&I admissions per beneficiary, the effectiveness of the vaccine, the vaccination rate above base-line, and the probability of a severe or mild season.

[¶]Sum of savings based on probability of a mild or severe season.

Vaccine Demonstration — Continued

costs in the cost-effectiveness analysis and would generate savings for Medicare if achieved nationally.

The demonstration's success in vaccine delivery resulted from focused interventions to overcome common barriers to adult vaccination, including the absence of a comprehensive vaccine delivery system, limited reimbursement mechanisms, and lack of vaccination programs where adults congregate. No statutory requirements mandating vaccination of Medicare beneficiaries were necessary to implement this program (4). The results of the cost-effectiveness analysis varied because of the variability of influenza from season to season in causing disease outcomes and the difficulty of attributing these outcomes to influenza. Nonetheless, provision of influenza vaccine was cost-effective for Medicare and may be cost-saving, depending on the effectiveness of the vaccine and the level of vaccination coverage.

Health-care providers such as physicians, hospitals, skilled-nursing facilities, home health agencies, and public health departments can now bill Medicare for reimbursement for the cost of influenza vaccine and the cost of its administration. The procedure codes for billing are 90724 and Q0124, respectively. Additional information for health-care providers in each state is available from the state's Medicare intermediary or carrier.

Implementation of this benefit should substantially improve influenza vaccine coverage among all Medicare beneficiaries, and thus reduce the high levels of morbidity and mortality attributed to influenza. However, both the public and health-care providers need to be educated about the major health burden of influenza-related illness and the necessity of vaccination to prevent it.

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Tuberculosis Among Pregnant Women — New York City, 1985–1992

From 1985 through 1992, the number of reported tuberculosis (TB) cases increased 20% in the United States (1). During 1985–1990, TB cases increased 44% among persons aged 25–44 years and 27% among children (aged <15 years) (2), indicating that TB may be an increasing problem among reproductive-aged women (3,4). To determine the prevalence of active TB during pregnancy, the medical records from 1985 through 1992 of two public hospitals in New York City were reviewed. This report summarizes the results of the survey.

The populations served by these two hospitals are largely inner-city, indigent, and minority populations with a high prevalence of both TB and human immunodeficiency virus (HIV) infection. Active TB was defined as a positive culture for tubercle bacilli (sputum, urine, or spinal fluid specimens), regardless of smear findings for acid-fast bacilli. Sixteen pregnant women with active TB (12 from one hospital) were identified; TB was diagnosed in five among 40,388 births (12.4 per 100,000 births) at these hospitals during 1985–1990, and in 11 among 11,595 births (94.8) during 1991–1992.

Five of the 16 women had received prenatal care before TB diagnosis: two, after a positive skin test and further evaluation, and three, after admission to the emergency department with TB-related symptoms. The 11 remaining women had received no prenatal care before TB diagnosis; these women's pregnancies were confirmed when they were admitted to the emergency department with symptoms associated with TB.

Of the 16 women, TB was diagnosed in one during the first trimester of her pregnancy; in seven, during the second trimester; and in eight, during the third trimester. A Mantoux tuberculin intradermal test was positive for six of the 15 women who were tested. Ten of the 16 women had pulmonary TB; six had extrapulmonary TB (two had tuberculous meningitis; one, mediastinal; one, renal; one, gastrointestinal; and one, pleural).

Seven of 11 women tested for HIV were HIV positive. Seven of the 16 women were drug users (defined as current use of cocaine or heroin). Six of the seven women who were HIV positive were drug users or were described by their physicians as injecting-drug users (IDUs): two women were cocaine users, three were IDUs, and one was both a cocaine user and IDU. Six of the seven women who were HIV positive and five of the six women who were drug users had received no prenatal care at the time their TB was diagnosed.

Thirteen of the 16 patients were successfully treated with isoniazid (INH), ethambutol (EMB), and rifampin (RIF). Two women with TB of the central nervous system received pyrazinamide (PZA). One woman with pulmonary TB (cavitary) received additional PZA because of persistent positive sputum cultures after 5 months of therapy with INH, EMB, and RIF. The remaining 10 women became asymptomatic on initial therapeutic regimens: eight had negative repeat cultures, and two required invasive biopsies and were not recultured.

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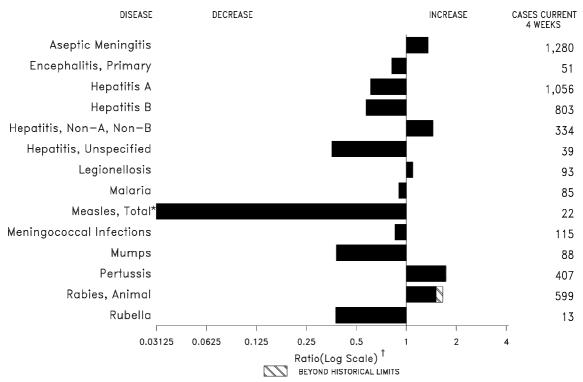


FIGURE I. Notifiable disease reports, comparison of 4-week totals ending August 7, 1993, with historical data — United States

*The large apparent decrease in reported cases of measles(total) reflects dramatic fluctuations in the historical baseline. (Ratio (log scale) for week thirty-one is 0.02966).

[†]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.

	Cum. 1993		Cum. 1993
AIDS* Anthrax Botulism: Foodborne Infant Other Brucellosis Cholera Congenital rubella syndrome Diphtheria Encephalitis, post-infectious Gonorrhea Haemophilus influenzae (invasive disease) [†]	67,732 8 15 2 56 15 6 9 8 223,223 754	Measles: imported indigenous Plague Poliomyelitis, Paralytic [§] Psittacosis Rabies, human Syphilis, primary & secondary Syphilis, congenital, age < 1 year [¶] Tetanus Toxic shock syndrome Trichinosis Tuberculosis	29 177 3 32 15,411 677 19 140 8 11,670
Hansen Disease Leptospirosis Lyme Disease	99 21 3,256	Tularemia Typhoid fever Typhus fever, tickborne (RMSF)	74 187 196

TABLE I. Summary — cases of specified notifiable diseases, United States, cumulative, week ending August 7, 1993 (31st Week)

*Updated monthly: last update July 31, 1993. [†]Of 695 cases of known age, 228 (33%) were reported among children less than 5 years of age. [§]No cases of suspected poliomyelitis have been reported in 1993; 10 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. [¶]Reports through first quarter of 1993.

	r												
	AIDS*	Aseptic Menin-	Enceph	Post-in-	Gond	orrhea			/iral), by	type Unspeci-	Legionel-	Lyme	
Reporting Area		gitis	Primary	fectious			A	В	NA,NB	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	67,732	5,213	353	98	223,223	292,951	12,526	7,166	2,760	363	671	3,256	
NEW ENGLAND Maine	3,232 94	123 16	11 1	5	4,876 52	6,043 56	282 8	315 9	316 -	9	25 4	795 4	
N.H. Vt.	67 14	17 17	- 3	2	43 16	75 15	13 3	54 5	250 2	2	2	31 3	
Mass. R.I.	1,818 219	52 21	5 2	3	1,737 228	2,212 434	154 53	195 16	57 7	7	15 4	79 126	
Conn.	1,020	-	-	-	2,800	3,251	51	36	-	-	-	552	
MID. ATLANTIC Upstate N.Y.	15,598 2,373	371 168	29 22	6 3	25,705 4,747	31,339 6,431	652 212	824 238	193 113	4 1	132 39	1,788 1,048	
N.Y. City	8,289	100	1	-	6,768	10,640	177	121	1	-	3	3	
N.J. Pa.	2,991 1,945	- 99	6	3	4,435 9,755	4,464 9,804	178 85	230 235	56 23	3	18 72	349 388	
E.N. CENTRAL	5,419	699	92 21	20	43,111	55,176	1,349	841	409	9	182	25	
Ohio Ind.	938 634	231 94	31 11	4 8	12,356 4,556	16,256 5,027	183 457	136 135	31 8	1	95 36	17 4	
III. Mich.	1,939 1,379	133 228	18 26	2 6	12,862 10,013	18,285 13,051	323 130	142 262	34 308	2 6	8 36	2 2	
Wis.	529	13	6	-	3,324	2,557	256	166	28	-	7	-	
W.N. CENTRAL Minn.	2,428 511	303 51	16 7	-	11,802 1,521	15,561 1,761	1,523 271	387 42	90 3	10 4	46 1	89 47	
lowa Mo.	141 1,374	59 76	1	-	602 6,706	1,009 8,571	26 966	15 279	5 64	1 5	6 11	6 7	
N. Dak. S. Dak.	1 22	8 7	3 3	-	29 164	53 102	56 12	-	-	-	1	2	
Nebr. Kans.	135 244	7 95	2	-	476 2,304	982 3,083	131 61	11 40	8 10	-	22 5	4 23	
S. ATLANTIC	244 14,279	93 1,239	2 64	40	2,304 60,419	90,319	754	1,361	357	47	122	442	
Del. Md.	253 1,630	32 113	3 14	-	823 9,610	1,047 8,925	8 106	107 172	72 7	- 5	9 28	218 77	
D.C.	896 1,049	24	24	-	3,034	3,924	5 93	30 91	-	20	13 3	2	
Va. W. Va.	46	118 13	10	4	7,192 369	10,587 516	9	26	22 16	-	1	32 _3	
N.C. S.C.	790 933	105 17	12	-	14,638 6,191	14,917 6,692	40 9	185 25	40	- 1	15 12	57 4	
Ga. Fla.	1,854 6,828	75 742	1	- 36	4,660 13,902	27,454 16,257	63 421	120 605	51 149	- 21	23 18	27 22	
E.S. CENTRAL	1,796	333	16	5	25,943	28,015	153	749	531	1	29	13	
Ky. Tenn.	213 731	121 82	9 5	4	2,726 7,852	2,846 9,187	74 31	52 631	9 508	-	11 13	3 8	
Ala. Miss.	531 321	87 43	1 1	- 1	9,296 6,069	9,094 6,888	32 16	63 3	4 10	1	2 3	2	
W.S. CENTRAL	6,957	591	26	2	26,475	31,748	1,204	962	157	110	20	26	
Ark. La.	267 921	30 41	1 1	-	5,128 6,915	4,689 8,978	31 46	35 127	2 61	2 2	2 2	1	
Okla. Tex.	590 5,179	1 519	6 18	- 2	2,120 12,312	3,214 14,867	81 1,046	168 632	53 41	7 99	11 5	13 12	
MOUNTAIN	2,948	319	16	4	6,448	7,294	2,446	350	186	55	48	13	
Mont. Idaho	22 52	-7	-	1	42 106	63 65	57 110	4 29	2	- 1	5 1	- 1	
Wyo. Colo.	31 985	5 82	- 6	-	55 1,932	32 2,659	11 617	16 48	55 34	32	5 5	8	
N. Mex.	240	57	3	2	559	531	219	135	58	2	3	-	
Ariz. Utah	992 197	110 15	5 1	-	2,440 204	2,556 161	849 519	54 33	10 21	8 11	9 6	2	
Nev.	429	42	1	1	1,110	1,227	64	31	6	1	14	2	
PACIFIC Wash.	15,075 1,008	1,236 -	83 1	16 -	18,444 2,318	27,456 2,453	4,163 463	1,377 130	521 115	118 7	67 9	65 1	
Oreg. Calif.	575 13,233	- 1,158	- 78	- 16	1,048 14,417	982 23,312	59 3,111	22 1,201	10 385	- 108	- 52	1 62	
Alaska Hawaii	47 212	11 67	3 1	-	320 341	424 285	477 53	7 17	9 2	- 3	6	- 1	
Guam	-	2	-	-	38	48	2	2	-	1	-	-	
P.R. V.I.	1,950 34	31	-	-	296 70	119 63	53	219 2	34	2	-	-	
Amer. Samoa C.N.M.I.	-	- 2	-	-	30 50	26 51	13	- 1	-	- 1	-	-	
0.11.111.1.		۷	-	-	50	51	-	1	-	I	-	-	

TABLE II. Cases of selected notifiable diseases, United States, weeks ending August 7, 1993, and August 1, 1992 (31st Week)

N: Not notifiable U: Unavailable *Updated monthly; last update July 31, 1993. C.N.M.I.: Commonwealth of Northern Mariana Islands

	Measles (Rubeola) Menin-															
				-			Menin- gococcal	Mu	mps		Pertussi	s	Rubella			
Reporting Area	Malaria	Indig	enous	Impo	orted*	Total	Infections									
	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	608	-	177	4	29	2,072	1,558	15	1,055	118	2,060	1,211	4	134	125	
NEW ENGLAND		-	47	-	4	54	91 5	-	8	9 1	463 9	95	-	1	6	
Maine N.H.	1 6	-	-	-	-	2 13	5 12	-	-	-	213	4 29	-	1	1 -	
Vt. Mass.	1 19	-	30 8	-	1 2	- 14	4 50	-	- 2	3 5	51 148	3 40	-	-	-	
R.I.	2	-	-	-	1	21	1	-	2	-	3	-	-	-	4	
Conn.	15	-	9	-	-	4	19	-	4	-	39	19	-	-	1	
MID. ATLANTIC Upstate N.Y.	96 35	-	7	-	3 1	194 110	192 88	2	80 27	13 7	245 97	63 30	3 2	39 8	10 7	
N.Y. City N.J.	24 27		2 5	-	- 2	48 36	19 30		- 8	-	7 26	9 24	-	15 11	- 3	
Pa.	10	-	- 5	-	-		55	2	45	6	115	- 24	1	5	-	
E.N. CENTRAL	31	-	12	-	1	43	238	-	147	17	315	136	-	2	9	
Ohio Ind.	9 3	-	5	-	-	6 20	73 40	-	57 3	16	158 35	29 17	-	1	-	
III.	14	-	3	-	-	10	65	-	35	-	33	21	-	-	8	
Mich. Wis.	5	-	4	-	1	4	41 19	-	49 3	1	21 68	6 63	-	- 1	1	
W.N. CENTRAL	18	-	1	-	2	11	100	-	31	24	162	102	-	1	7	
Minn. Iowa	4 1	-	-	-		10 1	6 16	-	1 7	19 1	83 2	33 3	-	-	- 2	
Mo.	5	-	1	-	-	-	38	-	18	-	48	42	-	1	1	
N. Dak. S. Dak.	2 2	-	-	-	-	-	3 3	-	4	- 2	3 5	10 5	-	-	-	
Nebr.	3	-	-	-	-	-	8	-	1	2	8	5	-	-	-	
Kans. S. ATLANTIC	1 180	-	- 17	-	2 3	- 119	26 298	- 7	- 343	2 25	13 239	4 81	-	- 8	4 12	
Del.	2	-	-	-	-	1	11	-	4	1	7	3	-	2	-	
Md. D.C.	19 5	-	-	-	2	16	33 5	4	62	5	79 2	14 1	-	2	4	
Va.	17	-	-	-	1	14	26	-	16	3	27	6	-	-	-	
W. Va. N.C.	2 88	-	-	-	-	24	11 55	2	11 195	2 3	11 38	4 14	-	-	1	
S.C.	1 9	-	-	-	-	29	26	-	14 14	-	8	8	-	-	2	
Ga. Fla.	37	-	17	-	-	- 35	65 66	- 1	27	- 11	12 55	8 23	-	4	- 5	
E.S. CENTRAL	19	-	1	-	-	459	96	-	36	4	93	20	-	-	1	
Ky. Tenn.	2 7	-	-	-	-	442	19 22	-	- 11	- 3	8 46	- 5	-	-	- 1	
Ala.	6	-	1	-	-	-	32	-	20	1	36	13	-	-	-	
Miss.	4	-	-	-	-	17	23 131	- 2	5 153	-	3 67	2 157	-	-	-	
W.S. CENTRAL Ark.	14 2	-	2	-	3	1,073	14	-	4	11 3	6	7	-	16	6	
La. Okla.	1 4	-	1		-	- 11	25 18	:	12 8	- 8	6 36	2 24	-	1 1	-	
Tex.	7	-	1	-	3	1,062	74	2	129	-	19	124	-	14	6	
MOUNTAIN	22	-	2	-	-	18	128	1	38	5	177	212	-	5	5	
Mont. Idaho	2 1	-	-	-	-	-	11 9	-	- 5	4	1 44	3 23	-	- 1	- 1	
Wyo. Colo.	- 13	-	- 2	-	-	1 14	2 21	-	2 9	-	1 61	- 26	-	-	-	
N. Mex.	5	-	- 2	-	-	14	21	N	9 N	1	25	20 44	-	-	-	
Ariz. Utah	-	-	-	-	-	2	62 12	-	6 3	-	29 16	91 24	-	1 2	2 1	
Nev.	1	-	-	-	-	-	7	1	13	-	-	1	-	1	1	
PACIFIC	184	-	88	4	13	101	284	3	219	10	299	345	1	62	69	
Wash. Oreg.	18 4	-	-	-	-	10 3	48 21	N	9 N	- 1	24 9	98 21	-	- 2	6 1	
Calif. Alaska	157 1	-	77	- 1 [†]	4 1	51 9	194 13	3	188 5	8	254 3	205 4	-	35 1	41	
Hawaii	4	-	- 11	3†	8	28	8	-	5 17	1	3 9	4 17	1	24	21	
Guam	1	U	2	U	-	10	1	U	6	U	-	-	U	-	1	
P.R. V.I.	-	-	224	-	-	293	6	-	2 3	-	2	9	-	-	-	
Amer. Samoa	-	U	1	U	- 1	- ว	-	U	-	U	2	6	U	-	-	
C.N.M.I.	-	U	-	U	1	2	- internationa	U	12	U	-	1	U	-	-	

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending August 7, 1993, and August 1, 1992 (31st Week)

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

Reporting Area		ohilis Secondary)	Toxic- Shock Syndrome	Tuber	culosis	Tula- remia	Typhoid Fever	Typhus Fever (Tick-borne) (RMSF)	Rabies, Animal
	Cum. 1993	Cum. 1992	Cum. 1993	Cum. 1993	Cum. 1992	Cum. 1993	Cum. 1993	Cum. 1993	Cum. 1993
UNITED STATES	15,411	20,416	140	11,670	13,002	74	187	196	4,931
NEW ENGLAND	248	393	10	270	223	-	18	2	823
Maine N.H.	3 25	2 28	2 2	7 9	17 3	-	- 1	-	53
Vt. Mass.	1 94	1 190	1 4	3 149	3 98	-	- 12	- 2	19 309
R.I.	9	21	1	34	23	-	-	-	-
Conn.	116	151	-	68	79	-	5	-	442
MID. ATLANTIC Upstate N.Y.	1,473 125	2,963 225	26 14	2,805 299	3,168 388	1 1	43 8	16 1	1,932 1,433
N.Y. City N.J.	773 202	1,665	1	1,646 454	1,885 529	-	26	- 10	323
Pa.	373	386 687	11	454 406	529 366	-	6 3	5	323 176
E.N. CENTRAL	2,320	3,098	38	1,179	1,300	3	20	9	53
Ohio Ind.	707 196	468 155	17 1	191 125	195 101	1 1	5 1	6	4 4
III.	796	1,391	5	551	663	-	9	1	7
Mich. Wis.	374 247	610 474	15	258 54	289 52	1	4 1	2	7 31
W.N. CENTRAL	962	813	9	257	306	25	2	9	221
Minn.	50	50	2 5	35	85	-	-	1	29
lowa Mo.	32 774	33 627	5	36 126	24 135	- 10	2	3 3	36 7
N. Dak. S. Dak.	- 1	1	-	5 10	4 14	- 11	-	2	47 32
Nebr.	10	21	-	14	13	1	-	-	7
Kans.	95	81	2	31	31	3	-	-	63
S. ATLANTIC Del.	4,158 80	5,631 134	16 1	2,031 29	2,389 25	2	26 1	95 2	1,204 94
Md.	238	410	-	232	172	-	5	9	354
D.C. Va.	228 368	249 476	- 4	100 270	78 179	-	- 3	- 5	11 221
W. Va. N.C.	8 1,170	12 1,431	- 3	49 293	53 305	- 1	-	4 47	50 51
S.C.	613	752	-	293	242	-	-	47	99
Ga. Fla.	707 746	1,132 1,035	2 6	444 365	535 800	- 1	1 16	16 5	282 42
E.S. CENTRAL	2,295	2,612	6	797	873	4	3	20	59
Ky.	187	89	2	231	234	-	-	5	10
Tenn. Ala.	650 510	728 980	1 2	144 286	235 233	3 1	1 2	11 2	49
Miss.	948	815	1	136	171	-	-	2	-
W.S. CENTRAL Ark.	3,251 504	3,523 544	2	1,385 120	1,302 103	29 18	2	41 1	348 18
La.	1,499	1,487	-	-	107	-	1	1	4
Okla. Tex.	241 1,007	177 1,315	2	167 1,098	95 997	8 3	- 1	38 1	54 272
MOUNTAIN	136	238	9	278	341	6	6	4	90
Mont. Idaho	1	7 1	- 1	15 8	- 14	2	-	-	15 5
Wyo.	5	3	-	2	-	2	-	4	11
Colo. N. Mex.	36 19	36 27	2	8 35	30 47	- 1	5	-	9 5
Ariz.	59	117	1	126	156	-	1	-	38
Utah Nev.	4 12	6 41	4 1	17 67	51 43	1	-	-	1 6
PACIFIC	568	1,145	24	2,668	3,100	4	67	-	201
Wash.	34 50	58	4	149	176	1	4	-	-
Oreg. Calif.	478	26 1,052	20	69 2,260	78 2,656	2 1	61	-	184
Alaska Hawaii	4 2	4 5	-	30 160	41 149	-	- 2	-	17
Guam	2	3	-	28	42	-	۲ -	-	-
P.R.	334	191	-	152	135	-	-	-	28
V.I. Amer. Samoa	31	39	-	2 2	3	-	-	-	-
C.N.M.I.	3	5	-	19	38	-	-	-	-

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending August 7, 1993, and August 1, 1992 (31st Week)

U: Unavailable

	All Causes, By Age (Years)								All Causes, By Age (Years)						
Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	P&I [†] Total	Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	P&I [†] 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.	581 162 36 11 21 54 21 17 55 55 35	399 985 7 18 366 18 15 19 28 33 33 5 33 26	90 29 5 4 2 10 1 2 6 5 3 9 5	58 19 4 1 4 2 1 3 5 1 9 4	22 10 2 - 3 - 1 3 - 3 3	12 6 - - 1 - 1 - 1 - 1 - 1	42 24 3 - 4 - 1 1 3 - 1 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.	1,263 151 218 77 109 110 50 63 57 48 150 206 24	734 91 132 44 76 57 27 36 32 35 88 95 21	283 30 52 21 18 32 8 17 12 5 32 54 2	159 23 24 8 9 15 7 4 12 3 19 34 1	43 5 7 2 3 4 - 3 7 12	43 2 3 4 4 3 4 6 1 2 4 10	42 4 10 2 4 - 1 3 4 3 6 5
Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Erie, Pa.§	53 2,446 55 35 100 22 23 42	38 1,516 35 26 68 10 14 31	8 520 13 5 25 5 2 5	4 291 2 3 3 2 7 4	62 5 1 3 4 -	3 57 - 1 1 2	3 104 2 1 3 3	E.S. CENTRAL Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala. Nashville, Tenn.	702 89 52 77 72 169 85 50 108	462 36 43 49 111 53 35 73	141 14 11 20 10 34 22 9 21	68 9 3 10 8 18 8 2 10	15 1 2 2 5 1 3 1	16 3 2 3 1 1 3 3	45 3 2 3 19 4 2 9
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.	37 1,338 84 20 297 73 7 128 23 24 93 24 93 24 93 24 93 24 93	23 795 32 11 188 46 4 93 20 22 67 13 18 U	6 293 20 4 65 16 3 23 2 3 2 19 9 2 U	7 181 23 37 7 - - 3 1 1 U	1 34 1 2 5 - 1 - U	35 5 1 5 3 - 3 1 - U	3 47 8 4 13 4 - 3 - 4 5 1 1 U	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.	1,007 59 43 59 191 52 119 U 63 117 189 43 92	620 31 28 26 106 36 72 U 38 69 118 25 71	211 12 11 10 43 8 23 U 15 22 39 14 14	104 9 4 3 27 5 13 U 5 11 21 2 4	49 3 12 3 7 U 3 7 9 2 3	20 4 3 4 U 2 5 2	43 5 4 2 3 5 U 2 - 9 4 7
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Dayton, Ohio Dayton, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Garand Rapids, Micl Indianapolis, Ind. Madison, Wis. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio W.N. CENTRAL	2,206 59 26 573 97 148 171 103 212 46 60 12	1,293 423 18 230 68 90 105 77 116 32 45 5 32 45 532 114 22 78 37 32 34 633 53	449 8 7 117 25 19 41 11 11 11 3 8 35 7 25 7 25 7 16	246 4 112 8 21 14 27 3 4 3 2 27 3 4 3 2 23 3 8 6 46	155 4 89 2 6 5 4 9 4 1 4 1 1 4 1 2 3	63 1 25 2 6 2 1 9 - - 3 4 1 5 - 1 - 1 1 2 1	113 7 14 9 3 10 7 10 4 3 - 4 11 2 10 1 1 6 9 2 43	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. San Diego, Calif. San Francisco, Calif.	b. 41 98 118 20 183 17 76 2,025 18 117 28 73 91 626 22 139 152 132 152 152	9 73 23 47 58 371 14 95 96 88 84	$\begin{array}{c} 135\\ 12\\ 7\\ 16\\ 20\\ 4\\ 41\\ 16\\ 15\\ 367\\ 5\\ 24\\ 16\\ 13\\ 112\\ 3\\ 25\\ 322\\ 20\\ 39\\ \end{array}$	67 7 2 16 1 1 5 1 6 7 256 3 9 1 6 11 100 3 14 15 30	28 3 - 1 3 - 9 1 6 5 79 - 7 - 2 6 1 2 4 6 6 5	18 2 1 2 1 1 4 - 3 4 3 6 1 4 - 2 3 8 - 1 3 1 2	34 3 5 3 12 5 4 94 3 4 5 12 21 2 4 7 3
W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn. Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	117 29 34 98 53	533 84 25 25 64 38 92 56 75 47 27	22 3 7 20 11 21 12 21	40 7 2 8 4 8 5 8 1 3	23 3 - 3 - 4 3 6 3 1	21 1 3 - 3 2 4 6 1	43 5 1 8 6 10 3 - 8 1	San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash. TOTAL	149 35 149 50 83 11,711 [¶]	98 27 102 36 59	27 4 22 7 14	20 4 15 4 6	2 5 1 2 476	2 5 2 2 286	15 2 4 6 5 5

TABLE III. Deaths in 121 U.S. cities,* week ending August 7, 1993 (31st 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.

[†]Pneumonia and influenza.

⁹Because 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.

Tuberculosis — Continued

Women's Health and Fertility Br, Div of Reproductive Health, National Center for Chronic Disease Prevention and Health Promotion, CDC.

Editorial Note: The findings in this report document an increase in active TB among pregnant inner-city women in two hospitals in New York City. Many of these women had TB diagnosed after presentation with TB-related symptoms. These findings underscore the need for TB screening in high-risk communities. Because of their high rate of TB and their inadequate use of prenatal and general health care, special attention should be given to minority urban populations and some populations of recent immigrants from countries with high prevalences of TB (2,5).

HIV infection is an important risk factor for the development of clinical TB in an adult coinfected with *Mycobacterium tuberculosis* (6). Thus, screening for TB should focus on populations at high risk for HIV infection and acquired immunodeficiency syndrome, including IDUs and persons already infected with HIV.

TB-related symptoms can mimic the physiologic changes that occur during pregnancy (i.e., increased respiratory rate and fatigue). Consequently, pregnant women in high-risk groups and women from areas with a high prevalence of both HIV infection and TB should be routinely asked about contact with infectious TB patients, and tuberculin skin testing should always be considered for these women. Because prenatal or peripartum care is often the only contact many high-risk women have with the healthcare system, screening for TB and HIV counseling and testing should be offered at this time.

The most appropriate method of screening for TB infection is the tuberculin skin test (Mantoux technique). Pregnancy does not measurably alter the response to a tuberculin test; subsequent investigation of tuberculin reactors, and persons with symptoms of TB, should facilitate the diagnosis and treatment of TB in pregnant women.

Because approximately 10% of immunocompetent and 40% of HIV-infected persons with active TB are negative by the tuberculin skin test, a negative result should never rule out the possibility of active disease (3,6-8). Factors such as age, poor nutrition, immunosuppression by disease or drugs, viral infections, and overwhelming TB can decrease tuberculin reactivity (3). Anergy to tuberculin has been reported among adults with HIV infection; therefore, a thorough investigation to detect active TB should be undertaken for all persons with clinical features compatible with TB, regardless of the results of the tuberculin skin test (7), and for all pregnant women at risk for or with known HIV infection.

To rule out active TB, routine chest roentgenogram with proper shielding of the abdomen should be performed after the 12th week of gestation for women with a positive tuberculin skin test (3,7). A chest roentgenogram should be performed sooner if the woman has symptoms suggestive of pulmonary TB, even if the tuberculin skin test is negative (3,4). Moreover, a comprehensive and systematic diagnostic approach, including appropriate examination of specimens for mycobacteria, should be followed for all patients with HIV infection and pulmonary disease (7). A complete review of systems and physical examination should be conducted to exclude extrapulmonary TB.

The Advisory Council for the Elimination of Tuberculosis recommends initial treatment for nonpregnant patients with four drugs: INH, RIF, PZA, and EMB or streptomycin (SM) (1). For pregnant women, this regimen is modified to exclude SM be-

Tuberculosis — Continued

cause it may cause congenital ototoxicity, and PZA, because the risk for teratogenicity has not been determined (1,3,9). Pregnant women with drug-susceptible organisms can be treated safely with INH, RIF, and EMB (1,3), but treatment must be continued for 9 months (1,3). If resistance to other drugs is probable and susceptibility to PZA is likely, the risks and benefits of PZA should be weighed carefully, and its use should be considered.

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Emerging Infectious Diseases

Update: Hantavirus Disease — United States, 1993

Since the recognition of acute hantavirus-associated respiratory disease in the United States in May 1993, laboratory evidence of acute hantavirus infection has been confirmed in 30 persons in the southwestern United States; 20 (67%) of these persons have died. Of those 30 persons, 23 resided in the four-corners region (14 in New Mexico, six in Arizona, and three in Colorado). Previously reported cases outside the four-corners states occurred in a Nevada resident (1) and a Texas resident (2), neither of whom had traveled to the four-corners area, and a resident of another state who had traveled to and presumably was infected in the four-corners area (3). This report summarizes the other four confirmed cases and describes two cases under investigation; all of these cases occurred outside the four-corners area during July 1992–August 1993.

Confirmed Cases

Louisiana. During June 1993, a 58-year-old Louisiana bridge inspector who had not traveled to the four-corners area died following an illness characterized by bilateral interstitial infiltrates and hypoxemia. Polymerase chain reaction (PCR) evidence of hantavirus infection was found in lung tissue, and nucleotide sequence analysis of

Hantavirus Disease — Continued

viral genetic material PCR-amplified from the lung suggests the presence of a previously unrecognized hantavirus most closely related to but distinct from both the Prospect Hill virus and the virus circulating in the four-corners area.

Nevada. In August 1993, a 51-year-old central Nevada resident rapidly developed bilateral interstitial infiltrates and hypoxemia over 12 hours following a 6-day illness characterized initially by fever, myalgia, nausea, and vomiting, which progressed to coughing and shortness of breath. The patient, who developed high-titered immuno-globulin M (IgM) antibodies to hantavirus, had not traveled to the four-corners area. As of August 11, the patient remained hospitalized.

California. Two cases have been confirmed in California. In the first, in July 1993, a 27-year-old field biologist, who was working on the eastern slope of the California Sierra Nevada mountain range, had acute onset of an illness characterized by 2 days of fever, myalgia, and headache. The patient developed rapidly progressive bilateral interstitial infiltrates and hypoxemia and died the following day. Hantavirus infection was confirmed by IgM serology, PCR, and a positive immunohistochemical stain for hantavirus antigen on lung tissue. The second case was in a 29-year-old ranch worker on the California coast who died of rapidly progressive respiratory failure during September 1992, following 3 days of fever, myalgia, and cough. Recent immunohistochemical staining of preserved autopsy tissues revealed hantavirus antigen. Neither person had recently traveled to the four-corners area.

Other Investigations

CDC is assisting state health departments in other investigations, including 1) a California man who had serologic evidence of past hantavirus infection following recovery from a hantavirus-compatible illness during April 1993 and 2) a 16-year-old Oregon youth in whom hantavirus antigen was identified by immunohistochemical staining of lung tissue saved from autopsy in July 1992. The California man, but not the Oregon teenager, had traveled to a four-corners state during the month before onset of illness.

Reported by: J Bertman, MD, Mono County Health Dept, Bidgeport; H Meyers, MD, Orange County Health Dept, Santa Ana; A Chovil, Santa Barbara County Dept of Health, Santa Barbara; R Jackson, MD, GW Rutherford, III, MD, State Epidemiologist, California Dept of Health Svcs. C Ward, MD, TB Callister, MD, H Hayes, Nye Regional Medical Center, Tonopah; LM Oksenholt, DO, D Jones, MD, S Parker, MD, Reno; D Nelson, AF DiSalvo, MD, State Health Laboratory, D Kwalick, MD, State Health Officer, Div of Health, Nevada State Dept of Human Resources. K Hedberg, MD, D Fleming, MD, State Health Div, Oregon Dept of Human Resources. KJ Steier, DO, Dept of Medicine, EA Conway Medical Center, Louisiana State Univ, Monroe; L McFarland, DrPH, State Epidemiologist, Office of Public Health, Louisiana Dept of Health and Hospitals. Div of Field Epidemiology, Epidemiology Program Office; National Institute for Occupational Safety and 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: Newly recognized cases of acute illness with evidence of hantavirus infection in Louisiana, Nevada, and California, along with previously recognized cases in Nevada and Texas, further demonstrate that hantavirus-associated respiratory illness is not confined to the four-corners area of the southwestern United States. Distinctive hantavirus nucleotide sequences have been identified from a person with acute illness in Louisiana; this information, together with confirmation of human disease in areas of Texas (2) and Louisiana outside the known range of *Peromyscus*

Hantavirus Disease — Continued

maniculatus (4)—the implicated reservoir in the four-corners area—suggests the existence of an additional hantavirus with a different rodent reservoir in the south central United States (3,5,6). The continued occurrence of hantavirus disease underscores the importance of minimizing risk for exposure to rodents and their excreta. Interim recommendations for hantavirus infection risk reduction have been developed (7). This document contains specific recommendations for reducing rodent shelter and food sources in and around the home, recommendations for eliminating rodents inside the home and preventing them from entering the home, precautions for prevention measures for persons who have occupational exposure to wild rodents, and precautions for campers and hikers. Investigations of cases of recognized and suspected human hantavirus disease and potential rodent reservoirs are ongoing.

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Notice to Readers

Announcement of Meeting on Research Case Definition for Chronic Fatigue Syndrome

CDC will sponsor a meeting to address the research case definition for chronic fatigue syndrome (CFS) on September 27, 1993, in Atlanta. The meeting will be open to public health officials, researchers, and the public. The purpose of the meeting is to review data from population and clinical studies related to use of the CFS research case definition.

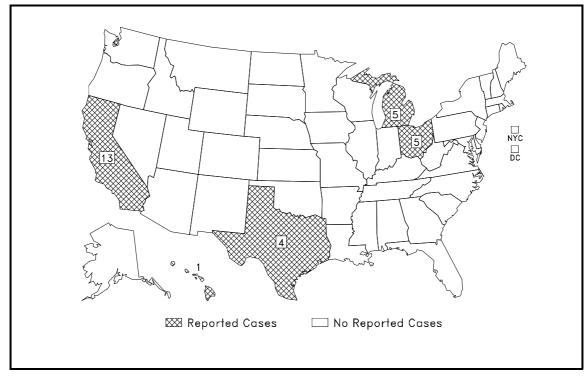
Additional information is available from CDC's CFS Research Program, Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases, Mailstop A-15, 1600 Clifton Road, NE, Atlanta, GA 30333; telephone (404) 639-1338; fax (404) 639-3163.

Erratum: Vol. 42, No. 29

In the article "Schistosomiasis in U.S. Peace Corps Volunteers—Malawi, 1992," on page 567 in the editorial note, the first paragraph, second sentence, should read "*S. mansoni* and *S. japonicum* primarily affect the *gastrointestinal* tract; chronic infection can lead to hepatosplenomegaly, variceal bleeding, and cirrhosis."

Erratum: Vol. 42, No. 23

In the article "Mortality Trends and Leading Causes of Death Among Adolescents and Young Adults—United States, 1979–1988," in Table 1 on page 460, the percentage change in other injury death rates for 15–19-year-olds should be -36.5, and the percentage change in other injury death rates for 20–24-year-olds should be -35.1.



Reported cases of measles, by state — United States, weeks 26–30, 1993

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