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MORBIDITY AND MORTALITY WEEKLY REPORT 4

Current Trends

Update: Mortality Attributable to HIV Infection/AIDS Among Persons Aged 25–44 Years — United States, 1990 and 1991

During the 1980s, human immunodeficiency virus (HIV) infection emerged as a leading cause of death in the United States (1). This report updates national trends in deaths caused by HIV infection during 1990 and 1991 and indicates that HIV infection/ acquired immunodeficiency syndrome (AIDS) continues to cause an increasing proportion of all deaths.

Data presented in this report were obtained from death certificates filed in all 50 states and the District of Columbia. Cause of death was reported by attending physicians, medical examiners, and coroners; demographic characteristics were recorded by funeral directors. Data for 1991 are provisional (2); 1990 is the latest year for which final and more detailed mortality data are available (3).

In 1991, 29,850 U.S. residents died from HIV infection; of these, 3% were aged <25 years; 74%, 25–44 years; and 23%, ≥45 years. HIV infection was the ninth leading cause of death overall, accounting for 1% of all deaths, and the third leading cause of death among persons aged 25–44 years, accounting for 15% of deaths in this age group (Table 1). In 1990, HIV infection was the second leading cause of death among men aged 25–44 years and the sixth leading cause of death among women in this age group (accounting for 17% and 5% of deaths, respectively) (Table 2). In 1991, the proportion of deaths caused by HIV infection in these two groups increased to 19% and 6%, respectively.

While death rates from most other leading causes of death declined or remained relatively stable for men and women aged 25–44 years, the death rate for HIV infection steadily increased (Figures 1 and 2). In 1991, the death rate for HIV infection for men aged 25–44 years was seven times that for women in this age group; however, since 1985, proportionate increases in the rate were greater for women than for men.

For men aged 25–44 years, the proportion of deaths caused by HIV infection in 1990 was 22% for Hispanics, 19% for blacks (non-Hispanic), 15% for whites (non-Hispanic), 7% for Asians/Pacific Islanders (non-Hispanic), and 3% for American Indians/Alaskan

HIV Infection/AIDS Mortality — Continued

Natives (non-Hispanic) (Table 3). HIV death rates* varied substantially by race/ethnicity: for men aged 25–44 years, rates for black, Hispanic, American Indian/Alaskan Native, and Asian/Pacific Islander men were approximately three times, twice, one third, and one fourth, respectively, the rate for white men (Table 3).

* In determining death rates by race/ethnicity, data were excluded from four states (Connecticut, Louisiana, New Hampshire, and Oklahoma) because information concerning Hispanic ethnicity was available for less than 85% of deaths. The criteria used in this report for determining which states were excluded from analysis of mortality data by Hispanic ethnicity differ somewhat from those used by CDC's National Center for Health Statistics; therefore, numbers of deaths in Table 3 differ from those published in Table 17 of reference *3*.

TABLE 1. Percentage of deaths caused by HIV infection, rank of HIV infection among all causes of death,* and death rate for HIV infection, by year of death and age group — United States, 1987–1991[†]

		All a	ges			Aged 25–44 yrs						
		HIV	-related	d death		HIV	-related	death	s			
Year	Total deaths	Deaths	(%)§	Death %) [§] Rank rate)		Total deaths	Deaths	(%)	Rank	Death rate		
1987 1988 1989 1990 1991	2,123,323 2,167,999 2,150,466 2,148,463 2,165,000	13,468 16,602 22,082 25,188 29,850	(0.6) (0.8) (1.0) (1.2) (1.4)	15 15 11 10 9	5.5 6.8 8.9 10.1 11.8	131,164 136,591 141,443 143,653 147,340	9,820 12,220 16,322 18,748 22,050	(7.5) (8.9) (11.5) (13.1) (15.0)	6 4 3 3 3	12.7 15.5 20.3 23.3 26.8		

*Based on the proportion of deaths from each of the cause categories used by CDC'sNational Center for Health Statistics to rank the 15 leading causes of death β).

[†]Data for 1991 are provisional; data for earlier years are final.

[§]Percentage of deaths caused by HIV infection among total deaths in the age group.

[¶]Deaths caused by HIV infection per 100,000 population.

			Men			Women							
		Н	IV-relate	ed death	าร		HIV-related deaths						
Year	Total deaths	Deaths	(%) [§]	Rank	Death rate [¶]	Total deaths	Deaths	(%)	Rank	Death rate			
1987 1988	91,082 95,419	8,867 10,935	(9.7) (11.5)	5 3	23.2 28.1	40,082 41,172	953 1,285	(2.4) (3.1)	8 8	2.5 3.3			
1989 1990 1991	99,482 101,519 104,380	14,646 16,717 19,380	(14.7) (16.5) (18.6)	2 2	37.0 41.7 47.3	41,961 42,134 42,960	1,676 2,031 2,670	(4.0) (4.8) (6.2)	6 6	4.2 5.0 6.4			

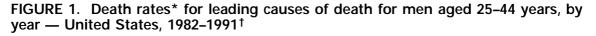
TABLE 2. Percentage of deaths caused by HIV infection, rank of HIV infection among all causes of death,* and death rate for HIV infection for persons aged 25–44 years, by sex and year of death — United States, 1987–1991[†]

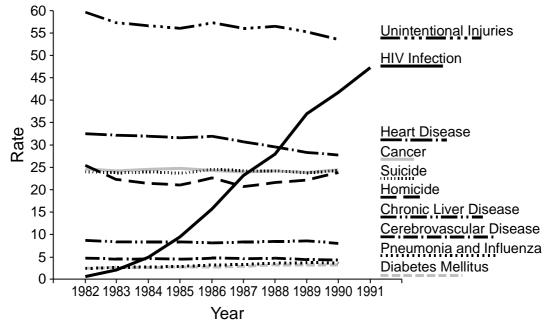
* Based on the proportion of deaths from each of the cause categories used by CDC'sNational Center for Health Statistics to rank the 15 leading causes of death (3). The rank could not be determined for 1991 because provisional sex- and age-specific data on deaths from other causes were unavailable for comparison.

[†]Data for 1991 are provisional; data for earlier years are final.

[§]Percentage of deaths caused by HIV infection among total deaths in the age and sex group. [¶]Deaths caused by HIV infection per 100,000 population.

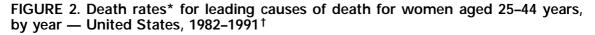
HIV Infection/AIDS Mortality - Continued

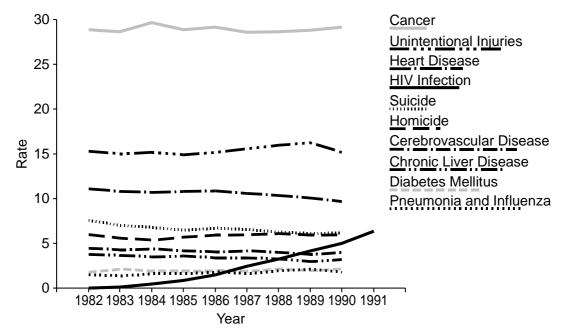




* Per 100,000 population.

[†]National vital statistics based on underlying cause of death, using final data for 1982–1990 and provisional data for HIV infection for 1991.





*Per 100,000 population.

[†]National vital statistics based on underlying cause of death, using final data for 1982–1990 and provisional data for HIV infection for 1991.

			Men			Women							
		ŀ	HV-relate	ed death	S	HIV-related deaths							
Race/ Ethnicity	Total deaths	Deaths	(%) †	Rank	Death rate [§]	Total deaths	Deaths	(%)	Rank	Death rate			
Non-Hispanic White Black Asian/	60,710 22,860	9,170 4,352	(15.1) (19.0)	2 2	31.4 105.0	25,354 10,584	434 1,160	(1.7) (11.0)	9 3	1.5 24.4			
Pacific Islander American Indian/	1,255	90	(7.2)	6	7.4	711	6	(0.8)	12	0.5			
Alaskan Native	894	24	(2.7)	7	10.0	390	3	(0.8)	11	1.2			
Hispanic Puerto Rican Cuban Mexican Other	1,897 554 4,985 1,187	728 223 659 231	(38.4) (40.3) (13.2) (19.5)	1 1 3 3	 	644 106 1,287 366	192 10 31 21	(29.8) (9.4) (2.4) (5.7)	1 5 8 4	 			
Unspecified nationality Total	1,722 10,345	382 2,223	(22.2) (21.5)	2 2	 59.1	459 2,862	58 312	(12.6) (10.9)	3 3	8.8			

TABLE 3. Percentage of deaths caused by HIV infection, rank of HIV infection among all causes of death,* and death rate for HIV infection among persons aged 25–44 years, by sex and race/ethnicity — United States, 1990

*Based on the proportion of deaths from each of the cause categories used by CDC's National Center for Health Statistics to rank the 15 leading causes of death.

[†]Percentage of deaths caused by HIV infection among total deaths in the age, sex, and racial/ethnic group.

[§]Deaths caused by HIV infection per 100,000 population, excluding data from four states (Connecticut, Louisiana, New Hampshire, and Oklahoma) because information concerning Hispanic ethnicity wasavailable for less than 85% of deaths. The criteria used in this report for determining which states were excluded from analysis of mortality data by Hispanic ethnicity differ somewhat from those used by CDC's National Center for Health Statistics; therefore, numbers of deaths differ from those published in Table 17 of reference 3. Death rates could not be determined by national origin for Hispanics, because information on national origin was available for less than 85% of their deaths in 28 states and age-specific population data were unavailable by national origin.

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HIV Infection/AIDS Mortality — Continued

For women aged 25–44 years, HIV infection accounted for 11% of deaths in 1990 for both black and Hispanic women; however, the HIV death rate for black women was nearly three times that for Hispanic women (Table 3). Both the proportions of deaths caused by HIV infection and the HIV death rates were substantially higher for black and Hispanic women than for women of white and other racial/ethnic groups.

Among Hispanics aged 25–44 years, the proportion of deaths caused by HIV infection in 1990 varied widely by national origin (including ancestry, not necessarily birthplace) (Table 3). In particular, among men of Cuban and Puerto Rican origin, HIV infection was the leading cause of death, accounting for approximately 40% of all deaths, while among men of Mexican origin, the proportion was lower (13%). In this age group, HIV infection was the leading cause of death among women of Puerto Rican origin—accounting for approximately 30% of all deaths—but caused a smaller proportion of deaths among women of Cuban origin (9%), Mexican origin (2%), and other Latin American origin (6%).

Reported by: Surveillance Br, Div of HIV/AIDS, National Center for Infectious Diseases; Mortality Statistics Br, Div of Vital Statistics, National Center for Health Statistics, CDC.

Editorial Note: The findings in this report underscore the role of HIV infection as a cause of death among men and women aged 25–44 years in the United States. Although deaths from all causes in this age group comprised only 7% of total U.S. deaths in 1991 (*2*), they impose a disproportionately high impact on society because of the loss of productive years of life and the loss of parents from families with young children. The impact of HIV infection on death patterns is even greater in many large cities than in the total U.S. population. For example, for persons aged 25–44 years in 1990, HIV was the leading cause of death among men in 64 (37%) of 172 cities with populations of at least 100,000 and among women in nine (5%) such cities (*4*).

In this report, the finding that rates of death for HIV infection were higher for blacks and Hispanics—particularly Hispanics of Puerto Rican origin—than for other racial/ ethnic groups is consistent with reported rates for the incidence of AIDS (*5,6*). Such comparisons of racial/ethnic groups may assist in targeting prevention efforts to groups at greatest risk. Differences in risk among racial/ethnic groups may reflect social, economic, behavioral, or other factors, rather than race/ethnicity directly (*7*). Further analyses are needed to better understand these associations.

The impact of HIV infection on U.S. mortality patterns is greater than indicated in this report. This analysis was based on the underlying cause of death recorded on death certificates; however, previous studies suggest that, for persons aged 25–44 years, deaths for which HIV infection is designated as the underlying cause represent 65%–85% of all HIV-related deaths among men and 55%–80% of those among women (8,9). In addition, provisional data for 1992 suggest that the number and proportion of deaths caused by HIV infection will increase beyond the levels described in this report (10). Increased prevention efforts to interrupt transmission of HIV are needed to decrease morbidity and mortality from HIV infection.

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HIV Infection/AIDS Mortality - Continued

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International Notes

Progress Toward Global Eradication of Poliomyelitis, 1988–1991

The report of the last case of smallpox from Somalia in 1977 demonstrated that an infectious disease could be eradicated globally. Because polioviruses have no animal reservoir and do not survive for long periods of time in the environment, and because lifelong immunity to paralytic poliomyelitis is conferred by existing, effective vaccines, poliomyelitis has been considered a candidate for eradication (1). In 1985, the Pan American Health Organization (PAHO) initiated a regional poliomyelitis eradication program. Based on the success of this program and high vaccination levels achieved worldwide by the Expanded Program on Immunization (EPI), in May 1988, the World Health Assembly of the World Health Organization (WHO) adopted a resolution to eradicate poliomyelitis globally by the year 2000. This report summarizes progress of the global poliomyelitis eradication initiative from 1988 through 1991*.

Global. Reported global vaccination coverage with three doses of oral poliovirus vaccine (OPV3) by age 1 year increased from 67% in 1988 to 84% in 1991 (Figure 1). During the same period, reported cases of poliomyelitis decreased 56%, from 32,286 to 14,176 (Figure 1). From 1988 through 1991, there were substantial decreases in the number of countries/territories reporting poliomyelitis cases (88 [45%] of 196 and 70 [34%] of 208, respectively) and the number of countries reporting more than 10 cases per year (56 [29%] and 38 [18%], respectively) (Figure 2). In addition, the number of countries reporting zero endemic cases increased from 107 (55%) to 129 (61%)[†].

African Region. Reported coverage with OPV3 increased from 44% to 57%, while reported cases of poliomyelitis decreased from 4546 to 2623; the number of countries in the region reporting poliomyelitis cases decreased from 37 (79%) of 47 to 25 (53%)

^{*}Based on surveillance data submitted to the EPI; because 1992 figures are povisional, 1991 data were used for global and regional disease incidence.

[†]The difference between the number of countries reporting poliomyelitis cases or zero cases and the total number of countries reflects those not submitting reports.

of 47. In 1991, the African Region reported 19% of the global total of poliomyelitis cases.

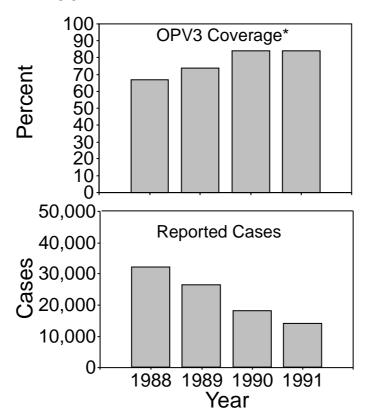
Region of the Americas. Reported coverage with OPV3 increased from 82% to 89%, while reported cases of poliomyelitis decreased from 340 to nine; the number of countries in the region reporting poliomyelitis cases decreased from 13 (28%) of 47 to two (4%) of 47. This region has reported no confirmed cases of poliomyelitis since September 1991 in Peru.

Eastern Mediterranean Region. Reported coverage with OPV3 increased from 69% to 80%, while reported cases of poliomyelitis decreased from 2332 to 2035; the number of countries in the region reporting poliomyelitis cases decreased from 17 (71%) of 24 to 15 (65%) of 23. In 1991, the Eastern Mediterranean Region reported 14% of the global total of poliomyelitis cases; 87% of the regional total were reported from Pakistan and Egypt. Despite OPV3 coverage of greater than 85%, small outbreaks also occurred in Oman (1988–1989) and Jordan (1991–1992); 51% of 118 persons with acute poliomyelitis in Oman and 53% of 32 persons with acute poliomyelitis in Jordan had received OPV3.

European Region. Reported coverage with OPV3 decreased from 86% to 82%, while reported cases of poliomyelitis increased from 206 to 313; the number of countries in the region reporting poliomyelitis cases increased from seven (23%) of 31 to 15 (33%)

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FIGURE 1. Reported coverage with three doses of oral poliovirus vaccine (OPV3) and poliomyelitis cases, by year — worldwide, 1988–1991



*Percentage of children who have received at least three doses of oral poliovirus vaccine by age 1 year.

CASES CURRENT DISEASE DECREASE INCREASE 4 WEEKS Aseptic Meningitis 562 Encephalitis, Primary 31 Hepatitis A 1,003 Hepatitis **B** 787 Hepatitis, Non-A, Non-B 271 Hepatitis, Unspecified 34 Legionellosis 74 Malaria 62 Measles, Total* 33 Meningococcal Infections 136 Mumps 163 Pertussis 234 Rabies, Animal 521 Rubella 5 2 0.03125 0.0625 0.125 0.25 0.5 1 4 Ratio(Log Scale) † \sum BEYOND HISTORICAL LIMITS

FIGURE I. Notifiable disease reports, comparison of 4-week totals ending June 26, 1993, with historical data — United States

*The large apparent decrease in 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 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) [†] Hansen Disease	51,608 7 12 2 36 14 5 - - - - - - - - - - - - - - - - - -	Measles: imported indigenous Plague Poliomyelitis, Paralytic [§] Psittacosis Rabies, human Syphilis, primary & secondary Syphilis, congenital, age < 1 year Tetanus Toxic shock syndrome Trichinosis Tuberculosis Tularemia	17 148 3 26 12,737 14 114 8 9,660 44
Leptospirosis Lyme Disease	17 1,916	Typhoid fever Typhus fever, tickborne (RMSF)	149 77

TABLE I. Summary — cases of specified notifiable diseases, United States, cumulative, week ending June 26, 1993 (25th Week)

*Updated monthly; last update June 5, 1993. [†]Of 573 cases of known age, 191 (33%) 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.

		Aseptic Menin-	Enceph		Com		Hep	oatitis (\	/iral), by t		Legionel-	Lyme
Reporting Area	AIDS*	gitis	Primary	Post-in- fectious	Gond	orrhea	Α	В	NA,NB	Unspeci- fied	Ĭosis	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	3,303	250	83	180,883	233,891	9,929	5,583	2,219	298	539	1,916
NEW ENGLAND	2,166	45	4	4	3,810	4,819	155	161	218	5	14	247
Maine N.H.	59 63	10 9	1	- 2	41 30	40 60	8 14	9 53	- 197	- 1	4 2	3 20
Vt.	14	7	2 1	-	14	13	3	3	2	-	- 5	1
Mass. R.I.	1,188 104	10 9	-	2	1,309 174	1,748 362	47 49	53 14	15 4	4	3	17 54
Conn.	738	-	-	-	2,242	2,596	34	29	-	-	-	152
MID. ATLANTIC Upstate N.Y.	11,379 1,938	313 123	11 3	6 3	20,335 4,037	24,550 5,070	587 179	725 200	162 93	4 1	114 35	1,331 969
N.Y. City	6,197	104	1	-	5,067	8,400	177	121	1	-	3	3
N.J. Pa.	2,072 1,172	- 86	-7	- 3	3,410 7,821	3,387 7,693	156 75	209 195	48 20	- 3	16 60	129 230
E.N. CENTRAL	4,160	435	76	15	34,957	43,922	976	572	358	8	143	17
Ohio	662	133	25	3	9,439	13,389	153	112	29	-	74	13
Ind. III.	502 1,442	65 87	6 16	7	3,695 11,381	4,039 14,409	412 293	110 123	6 21	1 2	30 5	1 1
Mich. Wis.	1,083 471	140 10	25 4	5	7,877 2,565	10,096 1,989	112 6	222 5	281 21	5	26 8	2
W.N. CENTRAL	2,163	192	4 11	-	8,945	12,498	1,250	337	21 97	5	35	37
Minn.	431	46	5	-	320	1,409	205	31	3	4	1	4
lowa Mo.	130 1,270	42 45	1	-	602 5,521	833 6,812	16 810	12 250	4 71	1	5 11	5 7
N. Dak.	-	5	2	-	23	43	43	-	-	-	1	1
S. Dak. Nebr.	20 100	7 4	3	-	150 476	84 754	10 113	- 8	- 9	-	14	-
Kans.	212	43	-	-	1,853	2,563	53	36	10	-	3	19
S. ATLANTIC Del.	10,888 208	811 8	45 3	36	49,609 642	73,121 838	616 6	984 73	271 63	44	91 7	208 106
Md.	1,216	70	10	-	7,726	7,010	85	135	6	5	22	30
D.C. Va.	548 731	19 79	- 15	- 3	2,681 5,651	3,535 8,672	3 64	14 73	- 20	- 18	12 2	2 19
W. Va.	38	7	7	-	279	439	3	18	16	-	1	2
N.C. S.C.	453 673	63 5	9	-	11,777 4,842	11,706 5,389	31 7	154 18	31	- 1	14 10	28 1
Ga.	1,562	53	1	-	4,660	23,200	57	35	21	-	12	-
Fla. E.S. CENTRAL	5,459 1,396	507 178	- 9	33 4	11,351 20,782	12,332 22,991	360 122	464 608	114 437	20 1	11 22	20 7
Ky.	161	70	4	4	2,193	2,345	64	46	5	-	8	2
Tenn. Ala.	528 463	27 49	4 1	-	6,326 7,418	7,345 7,843	23 25	505 54	424 3	- 1	11 1	3 2
Miss.	244	32	-	-	4,845	5,458	10	3	5	-	2	-
W.S. CENTRAL	5,311	298	19	-	21,411	23,806	924	753	101	81	15	10
Ark. La.	227 727	15 24	-	-	3,992 5,546	3,955 5,713	27 38	29 94	2 35	1	2	1
Okla.	423	1	4	-	1,738	2,340	53	115	26	6	9	5
Tex. MOUNTAIN	3,934 2,599	258 184	15 12	3	10,135 5,148	11,798 5,919	806 2,012	515 280	38 151	74 49	4 48	4
Mont.	15	-	-	1	22	51	54	4	-	-	5	-
ldaho Wyo.	43 28	6 3	-	-	87 41	59 25	95 10	23 13	- 45	1	1 5	- 2
Colo.	868	43	3	-	1,572	2,188	492	32	24	29	4	-
N. Mex. Ariz.	212 881	37 63	3 4	2	452 1,920	432 1,982	171 690	120 43	50 9	2 7	3 9	1
Utah Nev.	185 367	7 25	1 1	-	164 890	119	457 43	21 24	19 4	10	7 14	1
PACIFIC	307 11,546	25 847	63	- 15	15,886	1,063 22,265	43 3,287	24 1,163	4 424	- 101	57	- 55
Wash.	764	-	-	-	1,850	2,020	363	98	93	7	8	1
Oreg. Calif.	502 10,149	- 795	- 60	- 15	927 12,637	737 18,905	52 2,415	21 1,028	8 315	- 91	44	- 53
Alaska	12 119	4	2	-	221 251	359	410 47	6 10	6 2	- 3	- 5	- 1
Hawaii Guam	119	48 2	-	-	251	244 36	47	2	2	3 1	5	-
P.R.	1,561	27	-	-	209	84	36	180	22	2	-	-
V.I. Amer. Samoa	33	-	-	-	58 14	54 20	-10	2	-	-	-	-
C.N.M.I.	-	2	-	-	42	32	-	-	-	1	-	-
N: Not potifiable						o nu voalth (

TABLE II. Cases of selected notifiable diseases, United States, weeks endingJune 26, 1993, and June 20, 1992 (25th Week)

N: Not notifiable U: Unavailable *Updated monthly; last update June 5, 1993. C.N.M.I.: Commonwealth of Northern Mariana Islands

			Measle	s (Rube	eola)		Menin-									
Reporting Area	Malaria	Indig	enous		orted*	Total	gococcal Infections	Mu	mps	F	Pertussi	s		Rubella	3	
	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	5 439	3	148	1	17	1,951	1,300	32	888	44	1,233	783	2	107	106	
NEW ENGLAND Maine	22 1	-	42	-	2	44	56 5	-	5	4	295 8	67 2	-	1 1	6 1	
N.H.	5	-	-	-	-	12	12	-	-	4	192	20	-	-	-	
Vt. Mass.	1	-	30 3	-	1	- 8	4 16	-	-	-	42 19	1 33	-	-	-	
R.I. Conn.	2 11	-	- 9	-	1	20 4	1 18	-	2 3	-	2 32	- 11	-	-	4 1	
MID. ATLANTIC Upstate N.Y.	81 28	-	6	-	2 1	205 108	165 76	2 2	62 24	4 4	176 77	78 24	-	26 4	11 8	
N.Y. City	24	-	2	-	-	38	19	-	-	-	7	9	-	15	-	
N.J. Pa.	21 8	-	4	-	1	54 5	21 49	-	8 30	-	21 71	19 26	-	6 1	2 1	
E.N. CENTRAL Ohio	29 6	-	1	-	-	32 5	184 56	3 2	133 55	7 3	177 111	71 23	-	2 1	7	
Ind.	4	-	-	-	-	19	30	-	3	2	26	12	-	-	-	
III. Mich.	14 5	-	1	-	-	5 2	57 40	- 1	29 46	2	19 18	11 3	-	-	7	
Wis. W.N. CENTRAL	- 13	-	- 1	-	- 2	1 7	1 80	-	- 25	- 3	3 88	22 55	-	- 1	- 5	
Minn.	3	-	-	-	-	6	2	-	-	-	43	18	-	-	-	
lowa Mo.	1	-	-	-	-	1	15 32	-	7 13	- 3	1 24	1 22	-	1	1	
N. Dak. S. Dak.	2 2	-	-	-	-	-	3 3	-	4	-	3 1	7 4	-	-	-	
Nebr. Kans.	1 1	-	-	-	2	:	4 21	:	1	Ξ	5 11	2 1	-	:	4	
S. ATLANTIC	133	-	20	-	3	112	268	14	289	7	127	62	2	9	7	
Del. Md.	1 13	-	3	-	2	1 15	11 25	- 1	4 50	- 1	1 41	- 12	2	2 3	4	
D.C. Va.	5 8	-	-	-	- 1	- 11	4 24	- 2	- 16	- 1	2 11	- 4	-	-	-	
W. Va. N.C.	2 78	-	-	-	-	- 24	10 47	- 10	6 167	- 3	6 23	2 14	-	-	-	
S.C. Ga.	- 3	-	-	-	-	29	20 61	1	14	-	55	7	-	-	-	
Fla.	23	-	17	-	-	32	66	-	23	2	33	15	-	4	3	
E.S. CENTRAL Ky.	12	-	1	-	-	449 432	82 16	1	33	7	58 3	13	-	-	1	
Tenn. Ala.	7 3	-	- 1	-	-	-	17 30	1	10 18	3 4	33 20	5 7	-	-	1	
Miss.	2	-	-	-	-	17	19	-	5	-	2	, 1	-	-	-	
W.S. CENTRAL Ark.	11 2	-	1	-	-	999	114 12	11	132 4	-	32 2	105 6	-	12	6	
La. Okla.	- 4	-	1	-	-	- 11	24 10	- 5	11 7	-	5 12	13	-	1 1		
Tex.	5	-	-	-	-	988	68	6	110	-	13	86	-	10	6	
MOUNTAIN Mont.	13 2	-	2	-	-	12	111 10	-	35	7	87	118 1	-	4	4	
ldaho Wyo.	-	-	-	-	-	- 1	7	-	5 2	2	17 1	14	-	1	1	
Colo.	7 4	-	2	-	-	11	16	-	8	5	33	21	-	-	-	
N. Mex. Ariz.	4-	-	-	-	-	-	3 61	N	N 6	-	19 10	29 37	-	1	2	
Utah Nev.	-	-	-	-	-	-	5 7	-	3 11	-	7	15 1	-	1 1	1	
PACIFIC Wash.	125	3	74	1	8	91 10	240	1	174	5	193	214	-	52	59	
Oreg.	13 3	-	-		-	10	37 19	N	8 N	1	20 3	56 14	-	- 1	6 1	
Calif. Alaska	106	3	64	1 [†] -	3	47 9	166 10	1 -	147 5	4	160 3	136	-	30 1	36	
Hawaii	3	-	10	-	5	25 10	8	-	14	-	7	8	-	20	16	
Guam P.R.	1	U -	2 122	U -	-	10 244	1 6	U -	6 1	U -	-	- 9	U -	-	1	
V.I. Amer. Samoa	-	-	- 1	-	-	-	-	-	3	-	- 2	- 6	-	-	-	
C.N.M.I.	-	-	-	-	1	-	-	-	11	-	-	1	-	-	-	

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending June 26, 1993, and June 20, 1992 (25th 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	12,737	16,478	114	9,660	9,864	44	149	77	3,633
NEW ENGLAND	215	315	5	189	162	-	8	1	470
Maine N.H.	3 21	24	1 2	7 4	13	-	-	-	30
Vt. Mass.	1 86	1 151	- 1	3 112	3 74	-	- 6	- 1	16 76
R.I. Conn.	7 97	16 123	1	28 35	13 59	-	- 2	-	348
MID. ATLANTIC	1,243	2,320	23	2,108	2,389	-	44	6	1,386
Upstate N.Y.	103	187	12	189	300	-	9	1	1,036
N.Y. City N.J.	628 171	1,280 325	1	1,284 330	1,394 403	-	26 6	4	205
Pa.	341	528	10	305	292	-	3	1	145
E.N. CENTRAL Ohio	1,981 577	2,402 350	36 15	1,038 148	1,000 156	3 1	14 5	5 4	35 4
Ind. III.	177 732	114 1,098	1 5	113 528	82 489	1	1 4	- 1	- 4
Mich.	298	460	15	206	231	1	4	-	2
Wis. W.N. CENTRAL	197 783	380 655	- 8	43 214	42 232	- 13	- 2	- 6	25 176
Minn.	14	44	2	26	60	-	-	-	23
lowa Mo.	32 652	21 489	4	22 115	21 94	- 4	- 2	- 4	34 5
N. Dak. S. Dak.	- 1	1	-	2 10	3 14	-7	-	2	36 19
Nebr.	10	19	-	10	13	-	-	-	2
Kans. S. ATLANTIC	74	81	2 13	29	27	2 1	-	- 27	57
Del.	3,431 65	4,638 111	13	1,690 21	1,878 25	-	18 1	1	1,007 77
Md. D.C.	182 196	348 209	-	175 80	130 59	-	3	2	296 6
Va.	321	386	3	217	133	-	1	2	189
W. Va. N.C.	3 957	1,140	- 3	41 244	30 253	-	-	- 16	41 39
S.C. Ga.	538 569	633 951	-	204 360	193 423	-	- 1	1 1	84 233
Fla.	600	851	6	348	632	1	12	4	42
E.S. CENTRAL Ky.	1,812 146	2,161 66	4 2	667 183	706 185	3	2	8 3	45 7
Tenn.	522	605	1	144	164	2	-	3	-
Ala. Miss.	406 738	855 635	1	232 108	202 155	1	2	2	38
W.S. CENTRAL	2,694	2,802	2	946	929	17	2	22	299
Ark. La.	459 1,161	438 1,231	-	85	79 87	10	- 1	- 1	16 1
Okla. Tex.	187 887	123 1,010	2	151 710	68 695	4 3	- 1	21	58 224
MOUNTAIN	113	195	7		246	2	5	2	47
Mont. Idaho	1	3 1	- 1	219 5 6	- 12	-	-	-	9 1
Wyo.	4	1	-	1	-	1	-	2	6
Colo. N. Mex.	32 19	28 19	1	8 35	17 39	-	4	-	1 3
Ariz. Utah	50 2	97 5	1 3	108 11	112 37	- 1	1	-	25
Nev.	5	41	1	45	29	-	-	-	2
PACIFIC	465	990 49	16	2,589	2,322	5	54	-	168
Wash. Oreg.	27 47	24	2	127 53	143 49	1 2	4	-	-
Calif. Alaska	387 2	910 3	14	2,255 24	1,979 36	2	48	-	152 16
Hawaii	2	4	-	130	115	-	2	-	-
Guam P.R.	1 289	2 146	-	28 93	34 120	-	-	-	- 24
V.I.	28	31	-	2	3	-	-	-	-
Amer. Samoa C.N.M.I.	- 2	- 4	-	1 18	- 15	-	-	-	-

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending June 26, 1993, and June 20, 1992 (25th Week)

U: Unavailable

	A	All Cau	ises, By	y Age (Y		-1	P&I [†]			All Cau	uses, By	y Age (Y	/ears)		P&I [†]
Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn. Cambridge, Mass. Fall River, Mass. Hartford, Conn. Lowell, Mass. New Bedford, Mass. New Bedford, Mass. New Haven, Conn. Providence, R.I. Somerville, Mass. Springfield, Mass. Springfield, Mass. Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Elizabeth, N.J. Erie, Pa.§ Jersey City, N.J. New York City, N.Y. Newark, N.J. Paterson, N.J. Philadelphia, Pa. Reading, Pa. Reading, Pa. Rechester, N.Y. Scranton, Pa.§ Syracuse, N.Y. Trenton, N.J. Utica, N.Y.	44 49 6 22 61 2,691 33 17 92 355 25 40 53	$\begin{array}{c} 450\\ 106\\ 38\\ 24\\ 23\\ 34\\ 12\\ 22\\ 27\\ 37\\ 4\\ 38\\ 18\\ 52\\ 1,664\\ 19\\ 12\\ 633\\ 819\\ 35\\ 100\\ 307\\ 40\\ 307\\ 40\\ 307\\ 40\\ 307\\ 40\\ 12\\ 34\\ 18\\ 21\\ 34\\ 18\\ 21\\ 18\\ 18\\ 18\\ 18\\ 18\\ 18\\ 18\\ 18\\ 18\\ 1$	30 1 4 5 14 10 2 1 8 4 4 5 5 6 6 9 4 19 10 3 9 12 308 2 107 10 - 17 1 2 10 - 12 10 - 12 - 10 - 12 - 10 - 12 - 10 - - - - - - - - - - - - -	53 21 5 3 1 5 3 1 - - 3 324 1 5 324 1 5 324 1 5 324 1 5 324 1 5 32 7 53 5 2 3 1 - 4 6 1 2	14 6 4 - 1 - 1 - 2 63 2 - 3 1 - 1 2 24 3 1 22 - 1 - 2 - 1	9 7 1 1 - - - - - - - - - - - - - - - - -	56 21 72 32 12 22 5 9 121 22 23 46 7 29 3 12 12 22 3 46 7 12 12 26 - 1	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. 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.	160 126 28 785 91 92 89 75 159 94 59 126 1,536 60	$\begin{array}{c} 667\\ U\\ 167\\ 411\\ 76\\ 34\\ 300\\ 500\\ 355\\ 411\\ 107\\ 64\\ 22\\ 501\\ 107\\ 64\\ 22\\ 501\\ 107\\ 57\\ 400\\ 78\\ 935\\ 453\\ 332\\ 117\\ 553\\ 322\\ 117\\ 553\\ 322\\ 117\\ 553\\ 322\\ 117\\ 553\\ 322\\ 117\\ 553\\ 322\\ 117\\ 553\\ 322\\ 588\\ 199\\ 400\\ 966\\ 134\\ 70\\ 588\\ 588\\ 199\\ 96\\ 588\\ 109\\ 588\\ 109\\ 588\\ 109\\ 588\\ 109\\ 588\\ 109\\ 588\\ 109\\ 588\\ 109\\ 588\\ 109\\ 588\\ 109\\ 588\\ 109\\ 588\\ 109\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 588\\ 100\\ 100\\ 588\\ 100\\ 100\\ 588\\ 100\\ 100\\ 100\\ 100\\ 100\\ 100\\ 100\\ 1$	$\begin{array}{c} 236\\ U\\ 75\\ 122\\ 17\\ 15\\ 8\\ 4\\ 35\\ 266\\ 4\\ 169\\ 120\\ 299\\ 200\\ 13\\ 29\\ 286\\ 11\\ 7\\ 8\\ 322\\ 123\\ 800\\ 16\\ 23\\ 43\\ 14\\ 14\end{array}$	$\begin{array}{c} 126\\ U\\ 34\\ 6\\ 11\\ 22\\ 6\\ 5\\ 4\\ 3\\ 15\\ 18\\ 2\\ 66\\ 9\\ 9\\ 4\\ 6\\ 13\\ 12\\ 4\\ 9\\ 178\\ 10\\ 16\\ 6\\ 23\\ 6\\ 6\\ 50\\ 9\\ 20\\ 20\\ 5\\ 7\end{array}$	33 U 7 2 2 2 2 8 1 2 2 5 - 29 3 3 2 1 9 2 2 7 81 - 2 4 10 2 9 17 5 5 11 1 5	43 U 7 4 1 6 3 4 4 1 1 3 2 0 2 7 7 1 3 1 3 3 3 5 3 2 9 9 1 4 15 5 4 10 12	51 U2222 5312113 606556659310 89924433353 1592
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Cleveland, Ohio Cleveland, Ohio Dayton, 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. Rockford, 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. Minneapolis, Minn Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	2,289 44 48 590 800 146 197 128 214 35 62 22 40 117 56 727 400 33 19 111 29	$\begin{array}{c} 1,364\\ 32\\ 36\\ 217\\ 60\\ 88\\ 123\\ 91\\ 132\\ 25\\ 47\\ 12\\ 46\\ 116\\ 24\\ 47\\ 101\\ 355\\ 322\\ 9\\ 73\\ 45\\ 530\\ 311\\ 23\\ 17\\ 77\\ 19\\ 143\\ 65\\ 711\\ 53\\ 31\\ \end{array}$	483 9 8 146 12 37 47 47 7 8 39 8 25 5 11 8 24 7 112 7 8 24 7 112 7 8 24 7 112 7 8 24 7 112 7 8 24 7 8 24 7 7 8 24 7 7 8 24 7 7 8 8 25 5 1 8 8 25 7 7 8 8 8 8 8 9 8 8 8 8 9 8 8 8 8 9 8 8 8 8 9 8 8 8 9 8 8 8 9 8 8 9 8 8 8 9 8 8 8 9 8 8 8 9 8 8 8 8 9 8 8 8 8 9 8 8 8 9 8 8 8 9 8 8 8 8 9 8	2266 2 3 1066 6 14 14 7 21 3 7 7 5 4 1 7 7 5 4 1 7 2 8 1 7 7 5 4 1 7 7 5 4 1 7 7 5 4 1 7 7 5 4 1 6 6 6 6 14 14 7 7 7 5 4 1 6 6 6 6 14 14 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	117 127 13629 2-366 -21162 241217 101 -1101	99 1 49 1 4 7 4 7 4 5 - 2 - 3 8 5 - 1 1 7 1 1 6 - - - - - - - - - - - - -	123 3200 7266100221 102214 9914423101 14723101 412146 131866	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. Glendale, Calif. Glendale, Calif. Glendale, Calif. Glendale, Calif. Honolulu, Hawaii Long Beach, Calif. Dortland, Oreg. Sacramento, Calif. San Jego, Calif. San Jose, Calif. Sant a Cruz, Calif. Seattle, Wash. Tacoma, Wash. TOTAL	b. 56 104 106 24 180 19 83 104 1,798 31 4 1,798 31 75 25 82 79 432 20 146 170 125	494 55 40 63 18 105 56 74 1,177 22 56 21 60 58 258 9 97 114 76 799 117 31 94 352 7,782	19 30 1 37 1 18 313 3 11 3 10 88 7 29 29 23 33 23 6 20 8 8 12	85 11 7 12 3 23 1 9 8 192 4 192 14 4 192 14 6 61 2 11 15 12 32 16 2 17 4 1,295	23 3 1 - 2 10 2 4 64 1 3 - 5 2 15 - 6 7 4 5 3 1 7 4 5 3 1 7 4 1 8 448	20 3 7 7 1 - 5 - 4 - - 4 - - - - - - - - - - - - -	41 2 6 1 1 6 4 102 3 1 9 6 21 2 7 8 6 3 17 6 2 4 5 684

TABLE III. Deaths in 121 U.S. cities,* week ending June 26, 1993 (25th 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.

⁵Because 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.

of 45. In 1991, the European Region reported 2% of the global total of poliomyelitis cases; 68% of the regional total was from republics of the former Soviet Union.

Southeast Asian Region. Reported coverage with OPV3 increased from 57% to 93%, while reported cases of poliomyelitis decreased from 22,814 to 6581; the number of countries in the region reporting poliomyelitis cases (nine [82%] of 11) was unchanged. In 1991, the Southeast Asian Region reported 46% of the global total of poliomyelitis cases; 91% of the regional total was from India.

Western Pacific Region. Reported coverage with OPV3 increased from 89% to 95%, while reported cases of poliomyelitis increased from 2079 to 2615; the number of countries in the region reporting poliomyelitis cases decreased from six (17%) of 35 to five (14%) of 35. In 1991, the Western Pacific Region reported 18% of the global total of poliomyelitis cases; 98% of the regional total was from the People's Republic of China and Vietnam.

Reported by: Expanded Program on Immunization, World Health Organization, Geneva. Surveillance, Investigations, and Research Br, National Immunization Program; Respiratory and Enterovirus Br, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases, CDC.

Editorial Note: Since 1988, all six WHO regions have reported substantial progress toward poliomyelitis eradication, and poliomyelitis has apparently been completely eliminated from one region.[§] In the Region of the Americas, three major strategies

[§]In April 1993, Canada reported isolation of wild poliovirus type 3 from asymptomatic members of a religious group that objects to vaccination. This virus was likely imported because it was identical to a wild poliovirus type 3 that caused an outbreak among persons of a eligious community objecting to vaccination in the Netherlands in 1992–1993 (2).

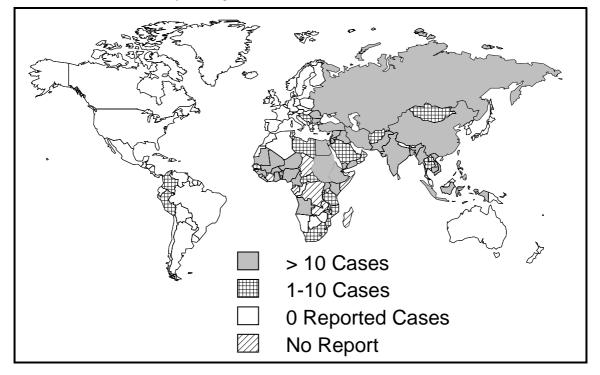


FIGURE 2. Incidence of poliomyelitis - worldwide, 1991

were used to eliminate poliomyelitis: 1) achievement of high vaccination coverage; 2) maintenance of sensitive systems of clinical and laboratory surveillance; and 3) implementation of supplementary vaccination activities, including national vaccination days biannually for all children below a specified age (usually age 5 years, regardless of prior vaccination status) and door-to-door vaccination campaigns in areas with a high incidence of poliomyelitis cases and/or low vaccination coverage (3).

In regions other than the Americas, vaccination strategies for poliomyelitis control have consisted primarily of routine vaccination. However, recent poliomyelitis outbreaks in highly vaccinated populations (4,5) and studies indicating suboptimal seroconversion to poliovirus types 1 and 3 following three doses of oral poliovirus vaccine in many tropical and subtropical regions suggest that routine vaccination alone may be insufficient to eliminate wild poliovirus infections and that supplementary activities, including national vaccination days, are necessary in countries where poliomyelitis is endemic (6).

In addition to the strategies used in the Region of the Americas, current global poliomyelitis eradication strategies include establishing and expanding polio-free zones and focusing additional resources on countries that are major exporters of wild poliovirus (7). The Global Poliomyelitis Eradication Plan of Action, endorsed by the EPI Global Advisory Group, emphasizes achieving effective surveillance of acute flaccid paralysis in all countries, initiating supplementary vaccination activities in all countries, and establishing a fully operational laboratory network in all WHO regions by 1995 with the goal of eliminating wild poliovirus transmission globally by the year 2000 (7).

Despite progress in increasing vaccination coverage and decreasing the incidence of poliomyelitis worldwide, there are at least five major barriers to global poliomyelitis eradication: 1) the presence of populations with suboptimal vaccination coverage, including unvaccinated subpopulations; 2) the failure of some countries and regions to identify poliomyelitis eradication as a priority activity (including the implementation of national vaccination days); 3) inadequate managerial skills to implement surveillance and vaccination programs effectively in certain countries; 4) suboptimal immunogenicity of oral poliovirus vaccine in many tropical and subtropical regions; and 5) inadequate commitment of financial resources at national and international levels (*3*).

The success of efforts to eradicate poliomyelitis in the Region of the Americas was based on the financial support of a broad coalition of national governments, international donor agencies (e.g., Rotary International, the United Nations Children's Fund, the Inter-American Development Bank, the Canadian Public Health Association, and the United States Agency for International Development), the Pan American Health Organization, and nongovernment community organizations. The creation of such coalitions—both regionally and globally—is of paramount importance in future efforts. In addition, success in global disease eradication requires that unaffected countries provide necessary assistance to geographic areas lacking adequate resources (1). The success of the global poliomyelitis eradication initiative will entail finding solutions to these financial, political, and technical challenges.

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Emerging Infectious Diseases

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

An outbreak of illness associated with hantavirus infection continues to be investigated by state health departments in New Mexico, Arizona, Colorado, and Utah; the Indian Health Service; and CDC, with the assistance of the Navajo Nation Division of Health (1-3). This report updates information regarding the outbreak and presents information on two cases that occurred in the 10 months preceding this outbreak.

Laboratory evidence of acute hantavirus infection has been confirmed in 15 patients who had onsets of illness from January 1 through June 30. Each of these patients has had one or more of the following: positive enzyme-linked immunosorbent assay (ELISA) serology with elevated immunoglobulin M titers indicating recent infection, seroconversion by ELISA, positive immunohistochemistry on formalin-fixed lung tissue, or amplification of hantavirus nucleotide sequences from frozen tissue. Of the 15 cases, 10 occurred in New Mexico, three in Arizona, and one in Colorado; 12 (80%) occurred among persons aged 20–40 years. Eleven patients died. Similar illnesses in an additional 23 persons, 10 of whom died, are being investigated for possible hantavirus infection.

Since June 6, a total of 668 rodents have been trapped in and around houses in 14 different rural sites. *Peromyscus maniculatus* (deer mouse) comprised 63% (range: 36%–88%) of all rodents trapped and 85% of those trapped in homes. Of the first 283 rodents tested, hantavirus antibodies were detected in 23%.

In June 1993, two persons were identified who had evidence of hantavirus infections in 1992. In November 1992, fever and acute respiratory distress occurred in a resident of the outbreak area. Recent serologic evaluation of an acute serum specimen obtained at the time of illness showed evidence of hantavirus infection. In August 1992, fever and myalgias followed by adult respiratory distress syndrome occurred in a person who resided outside the outbreak area; onset of illness was approximately 2 weeks after this person had returned home from a trip to the four-state area. The traveler had engaged in outdoor activities and was exposed to rodents and rodent excreta during both indoor and outdoor activities during the trip. A serum sample

Outbreak of Hantavirus Infection - Continued

tested in June 1993 showed elevated immunoglobulin G titers to hantavirus. Although a high immunoglobulin G titer in a single, recently obtained serum sample does not definitively establish the occurrence of a hantavirus infection at the time of illness, the serologic data and the clinical illness are strongly suggestive of hantavirus infection.

Reported by: F Koster, MD, H Levy, MD, G Mertz, MD, A Cushing, 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; 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, W Slanta, L Sands, DO, Acting 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 Naton, 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 identification of two persons with evidence of hantavirus infection that occurred in 1992 suggests that hantavirus infection has been present previously but was not recognized. Investigations are now in progress to identify whether changes in the local environment or other factors have been associated with the increased occurrence and/or transmission of this infection. Preliminary data from field investigations indicate that *P. maniculatus* is the likely reservoir of this virus. Alth ough the exact mechanism of hantavirus transmission to humans is unknown, potentially hazardous exposures include direct aerosolization of urine and other potentially infective rodent body fluids, secondary aerosolization of dried rodent excreta, contamination of food, and direct contact with virus-bearing rodents or their excreta or saliva.

Additional studies are under way to identify practical and effective means of preventing infection caused by hantaviruses. Residents and travelers in New Mexico, Arizona, Colorado, and Utah are advised to avoid any activities that may result in contact with wild rodents or rodent excreta or disruption of rodent burrows. The following specific recommendations for residents and travelers are based on current knowledge of transmission of other hantaviruses: 1) avoid activities that can result in contact with wild rodents, disruption of rodent burrows, or aerosolization of dried rodent excreta; 2) store food appropriately to avoid contamination with rodents and rodent excreta; and 3) dispose of food and trash properly to avoid attracting rodents.

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- 3. CDC. Update: outbreak of hantavirus infection—Southwestern United States, 1993. MMWR 1993;42:477–9.

Surveillance Summaries

Publication of CDC Surveillance Summaries

Since 1983, CDC has published the *CDC Surveillance Summaries* under separate cover as part of the *MMWR* series. Each report published in the *CDC Surveillance Summaries* focuses on public health surveillance; surveillance findings are reported for a broad range of risk factors and health conditions.

Summaries for each of the reports published in the most recent (June 4, 1993) issue of the *CDC Surveillance Summaries* (1) are provided below. All subscribers to *MMWR* receive the *CDC Surveillance Summaries*, as well as the *MMWR Recommendations and Reports*, as part of their subscriptions.

SURVEILLANCE FOR DIABETES MELLITUS — UNITED STATES, 1980–1989

Problem/Condition: In 1989, approximately 6.7 million persons reported that they had diabetes, and a similar number probably had this disabling chronic disease without being aware of it. Diabetes mellitus is the most important cause of lower extremity amputation and end-stage renal disease, the major cause of blindness among working-age adults, a major cause of disability, premature mortality, congenital malformations, perinatal mortality, and health-care costs, and an important risk factor for the development of many other acute and chronic conditions (e.g., diabetic ketoacidosis, ischemic heart disease, and stroke). Surveillance data describing diabetes and its complications are critical to increasing recognition of the public health burden of diabetes, formulating health-care policy, identifying high-risk groups, developing strategies to reduce the burden of this disease, and evaluating progress in disease prevention and control.

Reporting Period Covered: This report summarizes data from CDC's diabetes surveillance system, evaluating trends in diabetes and its complications by age, sex, and race for the years 1980–1989 (end year depending on data source).

Description of System: CDC has established an ongoing and evolving surveillance system to analyze and compile periodic, representative data on the disease burden of diabetes and its complications in the United States. Data sources currently include vital statistics, the National Health Interview Survey, the National Hospital Discharge Survey, and Medicare claims data for end-stage renal disease.

Results and Interpretation: These surveillance data indicate that the disease burden of diabetes and its complications is likely to grow as the population ages, that effective intervention strategies are needed to prevent diabetes and its complications, that prevention efforts need to be intensified among groups at highest risk, including blacks, and that important gaps exist in periodic and representative data for describing the burden of diabetes and its complications.

Actions Taken: CDC is currently exploring possible data sources to address the surveillance data gaps on blindness, adverse outcomes of pregnancy, and the public health burden of diabetes among minority groups.

Authors: Linda S. Geiss, M.A., William H. Herman, M.D., Merilyn G. Goldschmid, M.D., Frank DeStefano, M.D., M.P.H., Division of Diabetes Translation, National Center for Chronic Disease Prevention and Health Promotion, CDC. Mark S. Eberhardt, Ph.D., Office of Analysis and Epidemiology, National Center for Health Statistics, CDC. Earl S. Ford, M.D., M.P.H., Robert R. German, M.P.H., Jeffrey M. Newman, M.D., M.P.H., David R. Olson, Ph.D., Stephen J. Sepe,

Surveillance Summaries - Continued

M.P.H., John M. Stevenson, Ph.D., Frank Vinicor, M.D., M.P.H., Scott F. Wetterhall, M.D., M.P.H., Julie C. Will, Ph.D., Division of Diabetes Translation, National Center for Chronic Disease Prevention and Health Promotion, CDC.

LABORATORY-BASED SURVEILLANCE FOR MENINGOCOCCAL DISEASE IN SELECTED COUNTIES — UNITED STATES, 1989–1991

Problem/Condition: *Neisseria meningitidis* is a leading cause of bacterial meningitis and septicemia in the United States. Accurate surveillance for meningococcal disease is required to detect trends in patient characteristics, antibiotic resistance, and serogroup-specific incidence of disease.

Reporting Period Covered: January 1989 through December 1991.

Description of System: A case of meningococcal disease was defined by the isolation of *Neisseria meningitidis* from a normally sterile site, such as blood or cerebrospinal fluid, in a resident of a surveillance area. Cases were reported by contacts in each hospital laboratory in the surveillance areas. The surveillance areas consisted of three counties in the San Francisco metropolitan area, eight counties in the Atlanta metropolitan area, four counties in Tennessee, and the entire state of Oklahoma.

Results: Age- and race-adjusted projections of the U.S. population suggest that approximately 2600 cases of meningococcal disease occurred annually in the United States. The case-fatality rate was 12%. Incidence declined from 1.3 per 100,000 in 1989 to 0.9 per 100,000 in 1991. Seasonal variation occurred, with the highest attack rates in February and March and the lowest in September. The highest rates of disease were among infants, with 46% of cases in those ≤ 2 years of age. Males accounted for 55% of total cases, with an incidence among males of 1.2 per 100,000, compared with 1.0 per 100,000 among females (relative risk [RR]=1.3, 95% confidence interval [CI]=1.0–1.6). The incidence was significantly higher among blacks (1.5 per 100,000) than whites (1.1 per 100,000), with a relative risk of disease for blacks of 1.4 (95% CI=1.1–1.8). Sero-group B caused 46% of cases and serogroup C, 45%. Thirty-eight percent of isolates were reported to be resistant to sulfa; none were reported to be resistant to rifampin.

Interpretation: The decline in incidence of meningococcal disease from 1989 to 1991 cannot be explained by any change in public health control measures; this trend should be monitored by continued surveillance. The age, sex, and race distribution and seasonality of cases are consistent with previous reports. The proportion of *N. meningitidis* isolates resistant to sulfa continues to be substantial. A relatively small proportion of cases is potentially preventable by the use of the currently available polysaccharide vaccine, which induces protection against serogroups A, C, Y, and W135 and is effective only for persons >2 years of age.

Actions Taken: Current recommendations against the use of sulfa drugs for treatment or prophylaxis of meningococcal disease unless the organism is known to be sulfa sensitive should be continued. Since resistance to rifampin is rarely reported, it continues to be the drug of choice for prophylaxis. The development of vaccines effective for infants and vaccines inducing protection against serogroup B would be expected to have a substantial impact on disease.

Authors: Lisa A. Jackson, M.D., Jay D. Wenger, M.D., Meningitis and Special Pathogens Branch, Division of Bacterial and Mycotic Diseases, National Center for Infectious Diseases, CDC. The Meningococcal Disease Study Group.

Reference

1. CDC. CDC surveillance summaries (June 4). MMWR 1993;42(no. SS-2).

Notice to Readers

Change in Source of Information: Availability of Varicella Vaccine for Children with Acute Lymphocytic Leukemia

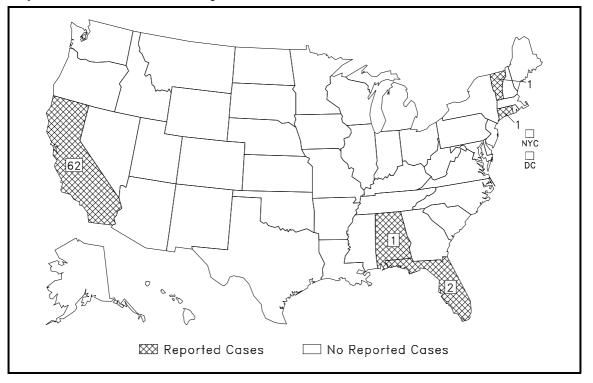
An investigational, live, attenuated varicella vaccine continues to be available free of charge through Merck Research Laboratories (West Point, Pennsylvania) to any physician requesting it for certain pediatric patients (aged 12 months–17 years) with acute lymphocytic leukemia (ALL) (1). However, the source of information about eligibility criteria and vaccine administration has changed (1) and is now available from the Varivax Coordinating Center, Bio-Pharm Clinical Services, Inc., 4 Valley Square, Blue Bell, PA 19422; telephone (215) 283-0897.

An Investigational New Drug application for the vaccine is on file with the Food and Drug Administration. Varicella vaccine is being provided to this group of patients for use through a study protocol to monitor and evaluate safety. Patients must meet specified criteria, including no clinical history of varicella and continuous remission for at least 12 months. The physician must provide information outlined in the protocol, and the protocol and consent form for the study must be approved by the institution's Investigational Review Board.

Reported by: National Immunization Program, CDC.

Reference

1. CDC. Availability of varicella vaccine for children with acute lymphocytic leukemia. MMWR 1992;41:326–7.



Reported cases of measles, by state - United States, weeks 21-25, 1993

The Morbidity and Mortality Weekly Report (MMWR) Series is prepared by the Centers for Disease Control and Prevention (CDC) and is available on a paid subscription basis from the Superintendent of Documents, U.S. Government Printing Office, Washington, DC 20402; telephone (202) 783-3238.

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Director, Centers for Disease Control and Prevention William L. Roper, M.D., M.P.H.
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