



MORBIDITY AND MORTALITY WEEKLY REPORT

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Health Objectives for the Nation

Rates of Cesarean Delivery — United States, 1991

Cesarean deliveries have accounted for nearly 1 million of the approximately 4 million annual deliveries in the United States since 1986 (Table 1). The cesarean rate in the United States is the third highest among 21 reporting countries, exceeded only by Brazil and Puerto Rico (1). This report presents data on cesarean deliveries from CDC's National Hospital Discharge Survey (NHDS) for 1991 and compares these data with previous years.

Data on discharges from short-stay, nonfederal hospitals have been collected annually since 1965 in the NHDS, conducted by CDC's National Center for Health Statistics. For 1991, medical and demographic information were abstracted from a sample of 274,000 inpatients discharged from 484 participating hospitals. The 1991 cesareans and vaginal births after a prior cesarean (VBAC) presented in this report are based on weighted national estimates from the NHDS sample of approximately 31,000 (11%) women discharged after delivery. The estimated numbers of live births by type of delivery were calculated by applying cesarean rates from the NHDS to live births from national vital registration data. Therefore, estimates of the number of cesareans in this report will not agree with previously published data based solely on the NHDS (2). Stated differences in this analysis are significant at the 95% confidence level, based on the two-tailed t-test with a critical value of 1.96.

In 1991, there were 23.5 cesareans per 100 deliveries, the same rate as in 1990 and similar to rates during 1986–1989 (Table 1). The primary cesarean rate (i.e., number of first cesareans per 100 deliveries to women who had no previous cesareans) for 1986–1991 also was stable, ranging from 16.8 to 17.5. In 1991, the cesarean rate in the South was 27.6, significantly (p<0.05) higher than the rates for the West (19.8), Midwest (21.8), and Northeast (22.6). Rates were higher for mothers aged ≥30 years than for younger women; in proprietary hospitals than in nonprofit or government hospitals; in hospitals with fewer than 300 beds than in larger hospitals; and for deliveries for which Blue Cross/Blue Shield* and other private insurance is the expected source of

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

TABLE 1. Estimated rates of cesarean delivery and number of live births,* by type of delivery — United States, selected years, 1965–1991

					Cesarean	delivery [†]				
	Cesarea	Cesarean rate		No.	Re	peat		VBAC¶		Other
Year	Primary**	Total ^{††}	No. live births ^{§§}	primary	No.	(%)§	Total	No.	Rate ^{¶¶}	vaginals†
1991	17.1	23.5	4111***	628	338	35.0	966	108	24.2	3037
1990	16.8	23.5	4158	626	351	35.9	977	90	20.4	3091
1989	17.1	23.8	4041	620	342	35.6	962	78	18.5	3001
1988	17.5	24.7	3910	615	351	36.3	966	50	12.6	2894
1987	17.4	24.4	3809	601	328	35.3	929	36	9.8	2844
1986	17.4	24.1	3757	595	310	34.3	905	29	8.5	2823
1985	16.3	22.7	3761	559	295	34.6	854	21	6.6	2886
1980	12.1	16.5	3612	418	178	29.9	596	6†††	3.4†††	3010
1975	7.8	10.4	3144	238	89	27.1	327	2†††	2.0†††	2815
1970	4.2	5.5	3731	153	52	25.2	205	1†††	2.2 ^{†††}	3525
1965	$NA^{\S\S}$	4.5	3760	NA	NA	NA	169	NA	NA	NA

^{*}In thousands.

[†] Estimated by applying cesarean rates derived from the National Hospital Discharge Survey (NHDS) to the number of live births from national vital registration data.

[§] Proportion of all cesareans that are repeat cesareans; standard error does not exceed 1.8% for any year.

Vaginal birth following a previous cesarean delivery. Estimated by applying cesarean rates derived from the NHDS to the number of live births from national vital registration data.

^{**}Number of first cesareans per 100 deliveries to women who did not have a previous cesarean; standard error does not exceed 1.1% for any year.

^{††} Number of cesarean deliveries per 100 total deliveries; standard error does not exceed 1.5% for any year.

^{§§} Source: National vital registration data.

Mumber of women with a VBAC per 100 deliveries of women with a previous cesarean delivery; standard error does not exceed 1.0% for any year.

^{***}Provisional data.

^{†††} Figure does not meet standards of reliability of precision because the weighted numerator is fewer than 10,000 deliveries.

^{§§§} Not available.

Cesarean Delivery — Continued

payment than for other sources of payment (Table 2). The same pattern characterized primary cesarean deliveries.

Since the early 1970s, the number and percentage of births to older women increased; however, if the age distribution of mothers in 1991 had remained the same as in 1986, the overall cesarean rate in 1991 would have been 23.3, essentially the same as the 23.5 observed.

Based on the NHDS, of the approximately 4,111,000 live births in 1991, an estimated 966,000 (23.5%) were by cesarean delivery. Of these, an estimated 338,000 (35.0%) births were repeat cesareans, and 628,000 (65.0%) were primary cesareans. Since 1986, approximately 600,000 primary cesareans have been performed annually. In 1986, 8.5% of women who had a previous cesarean delivered vaginally, compared with 24.2% in 1991. Of all cesareans in 1991, 35.0% were associated with a previous cesarean, 30.4% with dystocia (i.e., failure of labor to progress), 11.7% with breech

TABLE 2. Estimated total and primary cesarean rates,* by region, age of mother, hospital size and ownership, and expected source of payment — United States, 1991

	Estimated to	otal cesarean	Estimated pri	Estimated primary cesarean				
Category	Rate	(SE†)	Rate	(SE)				
Region								
Northeast	22.6	(0.5)	15.6	(0.5)				
Midwest	21.8	(0.5)	15.3	(0.4)				
South	27.6	(0.3)	20.5	(0.3)				
West	19.8	(0.5)	15.1	(0.5)				
Age (yrs) of mother								
<20	18.2	(1.5)	16.8	(0.6)				
20–24	21.0	(0.5)	15.9	(0.4)				
25–29	24.3	(0.5)	17.2	(0.4)				
30–34	26.7	(0.6)	17.6	(0.5)				
≥35	28.4	(0.9)	19.8	(8.0)				
Hospital size (no. beds)								
<100	24.6	(0.5)	17.9	(0.4)				
100–299	24.1	(0.3)	17.6	(0.3)				
300–499	22.4	(0.4)	16.4	(0.3)				
≥500	22.4	(0.5)	16.1	(0.5)				
Hospital ownership								
Nonprofit	23.3	(0.2)	16.7	(0.2)				
State and local government	20.7	(0.5)	15.6	(0.5)				
Proprietary	28.8	(0.6)	22.1	(0.6)				
Expected source of payment								
Blue Cross/Blue Shield§	27.6	(0.6)	20.1	(0.6)				
Other private insurance	25.3	(0.3)	18.3	(0.3)				
Medicaid	21.4	(0.3)	15.7	(0.3)				
Other government sources	21.3	(0.7)	15.8	(0.7)				
Self	20.7	(0.8)	15.7	(0.7)				
Other	17.8	(0.9)	13.0	(0.8)				
Total	23.5	(0.2)	17.1	(0.2)				

^{*}Total=number of cesarean deliveries per 100 total deliveries; primary=number of first cesareans per 100 deliveries to women who did not have a previous cesarean.

Standard error.

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Cesarean Delivery — Continued

presentation, 9.2% with fetal distress, and 13.7% with all other specified complications.

The average hospital stay for all deliveries in 1991 was 2.8 days. In comparison, the hospital stay for a primary cesarean delivery was 4.5 days, and for a repeat cesarean, 4.2 days—nearly twice the duration for VBAC deliveries (2.2 days) or for vaginal deliveries that were not VBACs (2.3 days). In 1986, the average hospital stay for all deliveries was 3.2 days, for primary cesareans 5.2 days, for repeat cesareans 4.7 days, and for VBAC and non-VBAC vaginal deliveries 2.7 and 2.6 days, respectively.

Reported by: Office of Vital and Health Statistics Systems, National Center for Health Statistics, CDC.

Editorial Note: The cesarean rate in the United States steadily increased from 1965 through 1986; however, the findings in this report indicate that rates have been stable since 1986 (3). Because there is little evidence that maternal and child health status has improved during this time and because cesareans are associated with an increased risk for complications of childbirth, a national health objective for the year 2000 (4) is to reduce the overall cesarean rate to 15 or fewer per 100 deliveries and the primary cesarean rate to 12 or fewer per 100 deliveries (objective 14.8).

Postpartum complications—including urinary tract and wound infections—may account in part for the longer hospital stays for cesarean deliveries than for vaginal births (5). Moreover, the prolonged hospital stays for cesarean deliveries substantially increase health-care costs. For example, in 1991, the average costs for cesarean and vaginal deliveries were \$7826 and \$4720, respectively. The additional cost for each cesarean delivery includes \$611 for physician fees and \$2495 for hospital charges (6). If the cesarean rate in 1991 had been 15 (the year 2000 objective) instead of 23.5, the number of cesarean births would have decreased by 349,000 (617,000 versus 966,000), resulting in a savings of more than \$1 billion in physician fees and hospital charges.

Despite the steady increase in VBAC rates since 1986, several factors may impede progress toward the year 2000 national health objectives for cesarean delivery. For example, VBAC rates substantially reflect the number of women offered trial of labor, which has been increasingly encouraged since 1982 (7). Of women who are offered a trial of labor, 50%–70% could deliver vaginally (7)—a level already achieved by many hospitals (8). Trial of labor was routinely offered in 46% of hospitals surveyed in 1984 (the most recent year for which national data are available) (9) when the VBAC rate (according to NHDS data) was 5.7%. The year 2000 objective specifies a VBAC rate of 35%, based on all women who had a prior cesarean, regardless of whether a trial of labor was attempted. To reach the overall cesarean rate goal, however, increases in the VBAC rate will need to be combined with a substantial reduction in the primary rate.

One hospital succeeded in reducing the rate of cesarean delivery by applying objective criteria for the four most common indications for cesarean delivery, by requiring a second opinion, and by instituting a peer-review process (10). Other recommendations for decreasing cesarean delivery rates include eliminating incentives for physicians and hospitals by equalizing reimbursement for vaginal and cesarean deliveries; public dissemination of physician- and hospital-specific cesarean delivery rates to increase public awareness of differences in practices; and addressing malpractice concerns, which may be an important factor in maintaining the high rates of cesarean delivery (4).

Cesarean Delivery — Continued

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International Notes

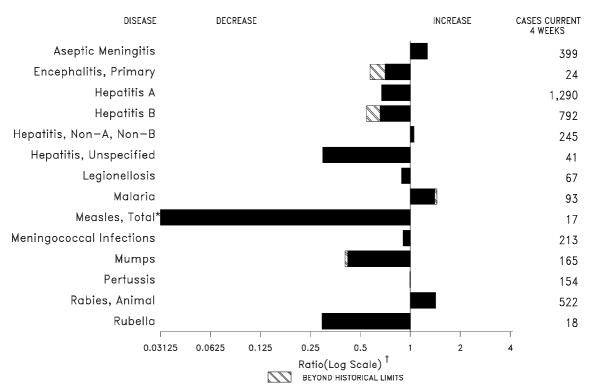
Malaria Among U.S. Embassy Personnel — Kampala, Uganda, 1992

The treatment and prevention of malaria in Africa has become a challenging and complex problem because of increasing drug resistance. Although the risk of acquiring malaria for U.S. citizens and their dependents stationed overseas generally has been low, this risk varies substantially and unpredictably. During May 1992, the Office of Medical Services, Department of State (OMS/DOS), and CDC were notified of an increased number of malaria cases among official U.S. personnel stationed in Kampala, Uganda. A review of the health records from the Embassy Health Unit (EHU) in Kampala indicated that 27 cases of malaria were diagnosed in official personnel from March through June 1992 compared with two cases during the same period in 1991. EHU, OMS/DOS, and CDC conducted an investigation to confirm all reported malaria cases and identify potential risk factors for malaria among U.S. Embassy personnel. This report summarizes the results of the investigation.

Malaria blood smears from 25 of the 27 reported case-patients were available for review by OMS/DOS and CDC. A case of malaria was confirmed if the slide was positive for *Plasmodium* sp. Of the 25 persons, 17 were slide-confirmed as having malaria.

A questionnaire was distributed to all persons served by the EHU to obtain information about residence, activities, use of malaria chemoprophylaxis, and use of personal protection measures (i.e., using bednets and insect repellents, having window and

FIGURE I. Notifiable disease reports, comparison of 4-week totals ending April 17, 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 fifteen is 0.02159.)

TABLE I. Summary — cases of specified notifiable diseases, United States, cumulative, week ending April 17, 1993 (15th Week)

	Cum. 1993		Cum. 1993
AIDS* Anthrax Botulism: Foodborne Infant Other Brucellosis Cholera Congenital rubella syndrome Diphtheria Encephalitis, post-infectious Gonorrhea Haemophilus influenzae (invasive disease)†	37,227 2 12 1 21 8 3 51 105,239 379	Measles: imported indigenous Plague Poliomyelitis, Paralytic [§] Psittacosis Rabies, human Syphilis, primary & secondary Syphilis, congenital, age < 1 year Tetanus Toxic shock syndrome Trichinosis Tuberculosis	13 73 1 - 16 - 7,646 - 5 72 7
Hansen Disease Leptospirosis Lyme Disease	379 39 11 777	Tularemia Typhoid fever Typhus fever, tickborne (RMSF)	4,377 15 81 23

[†]Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where thehatched area begins is based on the mean and two standard deviations of these 4-week totals.

^{*}Updated monthly; last update April 17, 1993.

†Of 349 cases of known age, 126 (36%) 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; all were vaccine associated.

TABLE II. Cases of selected notifiable diseases, United States, weeks ending April 17, 1993, and April 11, 1992 (15th Week)

	April 17, 1993, and April 11, 1992 (15th Week) Aseptic Encephalitis Hepatitis (Viral), by type													
_		Aseptic	Enceph				He	oatitis (\	/iral), by	type	Legionel-	Lyme		
Reporting Area	AIDS*	Menin- gitis	Primary	Post-in- fectious	Gond		Α	В	NA,NB	Unspeci- fied	Ĭosis	Dišease		
	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	37,227	1,777	149	51	105,239	142,690	5,844	3,014	1,181	161	306	777		
NEW ENGLAND	1,651	42	4	1	2,248	3,032	174	124	6	5	11	74		
Maine N.H.	51 50	6 4	1 -	-	27 13	32 39	8 4	3 13	-	-	2	- 7		
Vt. Mass.	8 819	5 23	3	- 1	9 874	7 1,150	3 100	2 95	1 2	- 5	8	24		
R.I.	80	4	-	-	109	244	39	11	3	-	1	19		
Conn. MID. ATLANTIC	643 6,434	140	- 5	4	1,216 11,194	1,560 15,439	20 267	331	84	3	63	24 566		
Upstate N.Y.	1,414	74	-	1	2,194	2,409	104	113	43	1	16	397		
N.Y. City N.J.	2,774 1,570	5 -	-	-	3,355 2,019	6,327 2,382	10 96	1 97	- 27	-	9	47		
Pa.	676	61	5	3	3,626	4,321	57	120	14	2	38	122		
E.N. CENTRAL Ohio	2,709 497	270 84	48 16	12 2	21,820 6,645	26,238 8,103	644 109	331 76	229 24	2	82 45	7 7		
Ind.	433	44	3	5	2,245	2,574	334	55	4	-	11	-		
III. Mich.	858 839	55 79	8 18	5	7,073 4,523	8,132 6,335	135 63	52 145	6 186	1 1	1 19	-		
Wis.	82	8	3	-	1,334	1,094	3	3	9	-	6	-		
W.N. CENTRAL Minn.	1,941 322	97 20	6 3	-	5,268 320	7,781 993	868 127	238 18	50 1	2 1	13 -	20 2		
Iowa	120	26 22	-	-	569	494	10 570	9 191	2 34	1	1 3	1 3		
Mo. N. Dak.	1,188	2	2	-	3,012 10	4,202 29	20	191	- 34	-	- -	- -		
S. Dak. Nebr.	18 88	4 1	1	-	58 141	59 454	9 96	- 5	- 7	-	- 7	-		
Kans.	205	22	-	-	1,158	1,550	36	15	6	-	2	14		
S. ATLANTIC Del.	7,778 158	457 2	27 1	22	29,817 374	46,226 489	352 2	485 43	172 52	22	57 6	67 47		
Md.	591	41	7	-	4,889	4,690	60	83	5	3	14	7		
D.C. Va.	354 566	13 55	- 7	3	1,798 2,716	2,433 5,633	2 50	10 44	12	10	7 2	1 5		
W. Va. N.C.	19 254	5 38	6 5	-	185 6,276	274 6,046	- 14	9 51	9 18	-	- 5	2 3		
S.C.	590	2	-	-	2,627	3,179	4	10	-	-	1	-		
Ga. Fla.	1,345 3,901	29 272	1 -	19	4,128 6,824	15,656 7,826	35 185	26 209	20 56	9	12 10	2		
E.S. CENTRAL	989	98	7	3	12,532	13,545	81	328	270	-	18	3		
Ky. Tenn.	79 393	45 22	2 4	3	1,340 3,715	1,452 4,515	46 16	31 264	4 262	-	6 10	2		
Ala. Miss.	350 167	25 6	1	-	4,626 2,851	4,427 3,151	17 2	31 2	2 2	-	2	1		
W.S. CENTRAL	4,497	100	10	-	12,863	13,353	409	345	53	36	7	9		
Ark. La.	181 595	9	-	-	1,717	2,522 1,824	16 18	16 35	2 17	-	2	1		
Okla.	421	-	3	-	3,222 953	1,444	27	60	17	5	5	5		
Tex.	3,300	88	7	-	6,971	7,563	348	234	17	31	-	3		
MOUNTAIN Mont.	2,252 10	101 -	8 -	3 1	3,023 13	3,296 21	1,225 43	192 4	87 -	35 -	28 3	3 -		
ldaho Wyo.	33 28	2	-	-	37 23	37 14	72 7	14 7	- 21	1	1 3	2		
Colo.	729	27	3	-	1,002	1,370	319	21	12	17	1	-		
N. Mex. Ariz.	186 799	13 39	2 2	2	304 1,029	266 989	94 376	92 27	30 6	1 7	1 6	-		
Utah Nev.	161 306	4 16	1	-	84 531	59 540	296 18	8 19	14 4	9	3 10	1		
PACIFIC	8,976	472	34	6	6,474	13,780	1,824	640	230	56	27	28		
Wash. Oreg.	139 459	-	-	-	1,020 457	1,224 413	196 34	52 16	49 4	5	2	-		
Calif.	8,360	446	31	6	4,714	11,767	1,336	562	174	50	23	28		
Alaska Hawaii	7 11	4 22	2 1	-	133 150	227 149	232 26	4 6	1 2	1	2	-		
Guam	-	-	-	-	14	30	1	_1	-	1	-	-		
P.R. V.I.	953 33	14	-	-	134 22	15 33	13	56 1	12 -	-	-	-		
Amer. Samoa C.N.M.I.	1	2	<u>-</u> -	-	7 18	10 11	6	-	-	-	<u>-</u> -	-		
O.1 V.1V1.1.	- 1		-		10	- 11		-		-				

N: Not notifiable

U: Unavailable

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

^{*}Updated monthly; last update April 17, 1993.

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending April 17, 1993, and April 11, 1992 (15th Week)

		1	Measle	s (Rube	eola)		Menin-	Menin-							
Reporting Area	Malaria	Indig	enous	Impo	orted*	Total	gococcal Infections	Mu	mps	ı	Pertussi	s		Rubella	ì
	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	5 252	1	73	-	13	542	794	24	486	38	700	334	9	46	42
NEW ENGLAND Maine	23	1	41	-	4	8	49 3	-	4	9	193 5	34 2	-	1 1	4
N.H. Vt.	2	1	-	-	1	1	7	-	-	4	119	13	-	-	-
Mass.	10	-	26 7	-	2	5	4 26	-	1	3 2	33 27	16	-	-	-
R.I. Conn.	1 9	-	8	-	1	2	1 8	-	2 1	-	2 7	3	-	-	4
MID. ATLANTIC Upstate N.Y.	29 15	-	5 1	-	1	94 26	96 45	5	47 13	17 5	128 48	60 20	-	7 1	6 4
N.Y. City	2	-	-	-	-	26	3	-	-	-	-	5	-	-	-
N.J. Pa.	7 5	-	4	-	1	39 3	11 37	- 5	6 28	12	20 60	16 19	-	5 1	2
E.N. CENTRAL Ohio	17 5	-	-	-	-	17 3	118 36	5 2	81 38	4 1	102 73	29 5	-	-	6
Ind.	3	-	-	-	-	9	20	-	-	2	11	9	-	-	-
III. Mich.	7 2	-	-	-	-	4	38 23	3	20 23	1	4 12	5 1	-	-	6
Wis. W.N. CENTRAL	3	-	-	-	- 1	1	1 44	- 1	- 15	- 1	2 27	9 26	-	- 1	2
Minn.	-	-	-	-	-	3	2	-	-	-	-	9	-	-	-
lowa Mo.	1 1	-	-	-	-	-	6 19	1 -	4	1 -	1 11	1	-	1	-
N. Dak. S. Dak.	1	-	-	-	-	-	2	-	4	-	1 1	4 1	-	-	-
Nebr. Kans.	-	-	-	-	- 1	-	2 13	-	1	-	4 9	2	-	-	2
S. ATLANTIC	87	-	12	-	2	61	160	3	122	4	53	37	3	5	2
Del. Md.	1 6	-	-	-	1	1 4	6 18	-	3 23	3	23	11	1	1 1	-
D.C. Va.	5 6	-	-	-	- 1	- 6	4 13	- 1	13	-	- 5	4	-	-	-
W. Va. N.C.	2 50	-	-	-	-	19	5 28	-	3 57	-	1	2	-	-	-
S.C. Ga.	2	-	-	-	-	-	13 42	1	12	-	5	7	-	-	-
Fla.	15	-	12	-	-	31	31	1	11	1	8	7	2	3	2
E.S. CENTRAL Ky.	4	-	-	-	-	243 227	52 9	2	17	1	27 3	2	-	-	-
Tenn. Ala.	1 2	-	-	-	-	-	14 16	1 1	8 6	- 1	16 8	1 1	-	-	-
Miss.	1	-	-	-	-	16	13	-	3	-	-	-	-	-	-
W.S. CENTRAL Ark.	7 1	-	1	-	-	62	66 6	3	75 3	-	15 1	13 7	-	8	-
La. Okla.	3	-	1	-	-	-	16 6	-	5 2	-	4 10	6	-	- 1	-
Tex.	3	-	-	-	-	62	38	3	65	-	-	-	-	7	-
MOUNTAIN Mont.	7 1	-	3	-	-	2	69 5	1	38	-	51	46	-	2	-
Idaho Wyo.	-	-	-	-	-	- 1	3 2	- 1	3 2	-	10 1	11	-	1	-
Colo.	4	-	2	-	-	1	9	-	4	-	20	19	-	-	-
N. Mex. Ariz.	2	-	1	-	-	-	2 41	N -	N 20	-	13 3	10	-	-	-
Utah Nev.	-	-	-	-	-	-	3 4	-	3 6	-	4	5 1	-	1	-
PACIFIC Wash.	75 5	-	11	-	5	52 7	140 18	4	87 6	2	104 7	87 24	6	22	22
Oreg. Calif.	2 67	-	- 5	-	-	1 35	14 99	N 4	N 71	2	, - 92	7 54	- 4	1 14	- 22
Alaska	- 1	-	- 6	-	- 5	9	4	-	4	-	1 4	- 2	- 2	1	-
Hawaii Guam	1	- U	-	- U	5	4	5 1	- U	6 4	- U	4	-	2 U	6	-
P.R. V.I.	-	Ŭ	72		-	35	5	Ŭ	2	Ŭ U	-	8	Ŭ	-	-
Amer. Samoa	-	U	1	U	-	-	-	U	-	U	2	-	Ú	-	-
C.N.M.I.	-	-	-	=	-	-	-	-	9	-	-	1	-	-	

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

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending April 17, 1993, and April 11, 1992 (15th Week)

Reporting Area		hilis 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	7,646	10,024	72	4,577	5,359	15	81	23	1,914
NEW ENGLAND	118	204	8	78	70	-	8	2	364
Maine N.H.	2 4	- 15	1 2	7 1	3	-	-	-	- 15
Vt. Mass.	-	1	4	1	-	-	-	-	7
R.I.	63 3	89 12	1	32 16	46 -	-	6	2	114 -
Conn.	46	87	-	21	21	-	2	-	228
MID. ATLANTIC Upstate N.Y.	678 68	1,407 106	16 9	1,001 77	1,221 155	-	10 4	2	590 406
N.Y. City N.J.	448 107	752 202	-	607 153	688 196	-	2 2	2	106
Pa.	55	347	7	164	182	-	2	-	78
E.N. CENTRAL	1,215	1,370	24	564	547	2	10	-	13
Ohio Ind.	349 114	205 60	12 1	81 54	90 51	1	4 1	-	2
III. Mich.	441 193	577 289	1 10	298 112	271 114	- 1	3 2	-	-
Wis.	118	239	-	19	21	-	-	-	11
W.N. CENTRAL	494	360	5	108	113	2	-	-	100
Minn. Iowa	14 28	28 9	2 2	26 5	33 7	-	-	-	18 16
Mo. N. Dak.	380	262 1	-	51 1	41 3	1	-	-	1 20
S. Dak.	-	-	-	6	8	-	-	-	10
Nebr. Kans.	7 65	14 46	- 1	5 14	5 16	- 1	-	-	1 34
S. ATLANTIC	2,089	2,842	7	755	1,072	-	12	5	523
Del. Md.	38 114	66 226	-	10 116	16 74	-	3	-	46 147
D.C.	142	146	-	51	43	-	-	-	4
Va. W. Va.	182 1	235 3	-	133 24	100 19	-	1	-	92 28
N.C. S.C.	496 343	677 350	3	114 91	148 111	-	-	4	10 42
Ga.	376	632	-	216	223	-	1	1	134
Fla.	397	507	4	-	338	-	7	-	20
E.S. CENTRAL Ky.	957 76	1,406 44	2 1	316 87	307 107	3	1 -	3 2	26 4
Tenn. Ala.	251 246	322 669	1	62 122	- 118	2 1	- 1	-	- 22
Miss.	384	371	-	45	82	-	-	1	-
W.S. CENTRAL Ark.	1,780 269	1,582 225	1	386 46	454 37	5 3	1	11	142 8
La.	707	716	-	-	26	-	1	-	-
Okla. Tex.	111 693	73 568	1 -	34 306	34 357	1 1	-	11 -	26 108
MOUNTAIN	65	139	2	143 5	145	-	3	-	21
Mont. Idaho	-	2 1	-	5 2	- 8	-	-	-	3
Wyo.	1	1	-	1	-	-		-	2
Colo. N. Mex.	21 12	22 16	1	8 18	17 20	-	2	-	2
Ariz. Utah	30 1	60	- 1	67 9	59 19	-	1	-	14
Nev.	-	37	-	33	22	-	-	-	-
PACIFIC	250	714	7	1,226	1,430	3	36	-	135
Wash. Oreg.	15 44	36 15	-	70 21	77 28	1	2	-	-
Calif. Alaska	185 2	657 2	7	1,046 8	1,237 22	2	32	-	121 14
Hawaii	4	4	-	81	66	-	2	-	-
Guam	- 157	1	-	18	24	-	-	-	- 14
P.R. V.I.	157 15	63 16	-	44 2	40 2	-	-	-	16 -
Amer. Samoa C.N.M.I.	-	2	-	1 7	- 10	-	-	-	-
O.1 W.1V1.1.	-	۷			10	-		-	

U: Unavailable

TABLE III. Deaths in 121 U.S. cities,* week ending April 17, 1993 (15th Week)

	April 17, 1993 (15th Week) All Causes, By Age (Years) All Causes, By Age (Years) Post														
	F	All Cau	ises, By	y Age (\	ears)		P&I [†]		All Causes, By Age (Years)						P&I [†]
Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	Total
NEW ENGLAND Boston, Mass. Bridgeport, Conn. Cambridge, Mass. Fall River, Mass. Hartford, Conn. Lowell, Mass. Lynn, Mass. New Bedford, Mas New Haven, Conn. Providence, R.I. Somerville, Mass. Springfield, Mass. Waterbury, Conn.		416 103 19 13 22 36 24 10 25 34 28 35 31	31 4 2 3 8 7 2 5 3	42 17 4 - 6 2 - 2 4 - 3	18 5 2 - 1 3 - - 2 2 1 -	8 4 1 1 1 - - - 1	57 24 3 1 1 1 4 2 1 5	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.	1,443 196 226 104 135 150 55 86 53 69 155 187 27	908 118 146 76 84 80 32 51 35 53 117 93 23	273 37 36 18 27 42 12 20 9 9 25 34 4	180 34 29 7 17 19 5 11 7 3 10 38	38 3 8 2 2 6 - 2 1 1 3 10	43 4 7 1 5 3 6 2 1 3 -	102 9 26 9 5 6 5 5 4 16 8
Waterbury, Corni. Worcester, Mass. MID. ATLANTIC Albany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Elizabeth, N.J. Erie, Pa.§ Jersey City, N.J. New York City, N.Y 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.	51 2,503 65 21 U 63 27 58 39	1,684 48 17 U 43 47 22 799 42 13 261 66 12 107 17 23 71 33 31 8	4 484 13 4 U 111 7 100 100 259 21 6 77 17 14 3 5 10 10	243 2 - U 4 2 1 7 140 4 4 44 5 2 6 2 1 3 7 7	54 1 - U 3 28 6 - 10 2 1 1 1 2	38 1 U 2 20 7 - 5 1	36 134 4 - U 1 1 4 - 45 5 - 366 7 2 14 1 1 5 7 2	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.	127 92 145 67 62 127 1,493 75 33	593 103 65 97 60 96 43 46 83 923 53 21 42 122 44 72 213 69 44 117 34	164 32 11 17 19 30 17 11 27 280 10 8 7 48 11 17 74 14 12 42 8	66 17 2 7 10 7 5 4 14 161 7 2 6 10 8 8 3 5 18 22 7	19 6 2 3 2 3 1 2 73 3 1 2 10 8 3 14 1 22 6 3	22 5 1 3 1 9 2 1 5 6 2 1 1 3 2 6 7 2 3 6 1 7 2 6 6 1 7 2 6 6 1 7 6 1 7 6 1 7 7 8 6 7 8 7 8 7 8 7 8 7 8 8 7 8 7 8 8 7 8 7	71 9 3 17 13 13 10 1 5 97 3 1 2 2 7 10 33 14
Yonkers, N.Y. E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind. Gary, Ind. Grand Rapids, Mic Indianapolis, Ind. Madison, Wis. Milwaukee, Wis. Peoria, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio W.N. CENTRAL Des Moines, Iowa Duluth, Minn. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn Omaha, Nebr. St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	32 2,387 91 358 428 428 176 183 262 46 72 21 h. 56 159 55 176 40 56 56 116 72 824 78 78 37 122 36	27 1,556 63 29 178 101 131 113 16 153 35 55 8 42 121 128 27 47 92 57 593 54 25 88 89 139 139 149 159 159 159 159 159 159 159 159 159 15	3 439 15 93 39 314 22 52 8 12 7 7 12 11 127 20 4 26 19 21 5	2 216 9 13 24 11 6 5 37 1 1 10 5 8 1 1 6 2 6 2 18 3 3 6 6 2 18 18 18 18 18 18 18 18 18 18 18 18 18	120 2 -3 63 7 7 6 -11 1 1 2 12 -1 -1 -1 -1 -1 -1 -1 -1 -1 -1 -1 -1 -1	56 2 111 5 3 3 5 5 9 9 1 1 2 2 6 6 - 1 1 2 2 2 2 3 5 - 4 4 4 5 5 1 1 1 1	148 19 13 3 9 10 11 65 4 21 2 18 5 4 68 7 11 1 8 2 17 2 9 4 5	Tulsa, Okla. 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. Los Angeles, Calif. Pasadena, Calif. Portland, Oreg. Sacramento, Calif. San Diego, Calif. San Francisco, Calif. San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash. TOTAL	o. 50 113 161 33 198 26 1 96 151 2,058 37 79 25 69 87 562 34 125 147	92 642 85 400 22 112 20 62 116 1,397 21 45 21 47 61 373 30 90 108 108 108 133 21 92 54 100 8,712	29 171 22 6 15 39 5 48 3 18 15 343 5 19 1 12 89 1 19 24 22 39 37 3 26 7 26 2,371	7 88 13 2 10 17 4 21 1 9 11 224 2 6 2 7 13 60 2 10 14 25 33 13 4 22 10 10 11 11 11 11 11 11 11 11 11 11 11	32 5 1 1 4 4 46 5 1 18 2 1 5 5 3 2 2 2 2 4 419	2 21 1 1 2 1 2 6 6 2 1 5 3 4 - 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	9 91 6 1 21 15 2 26 2 10 8 153 4 2 1 7 11 30 7 8 17 6 8 11 904

^{*}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.

Because of changes in reporting methods in these 3 Pennsylvania cities, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks.

Total includes unknown ages.

U: Unavailable.

Malaria — Continued

door screens, and wearing long sleeves and pants in the evening). Of the 157 persons eligible for the survey, 128 (82%) responded.

Risk for malaria was not associated with sex or location of residence in Kampala. Although the risk for malaria was higher among children aged ≤15 years (6/32 [19%]) than among persons >15 years (11/94 [12%]), this difference was not significant (relative risk [RR]=1.6; 95% confidence interval [CI]=0.6–4.0). Eighty-two percent of the cases occurred among persons who had been living in Kampala for 1–5 years, compared with those living there <1 year. Travel outside of the Kampala area to more rural settings was not associated with increased risk for malaria.

Four malaria chemoprophylaxis regimens were used by persons who participated in the survey: mefloquine, chloroquine and proguanil, chloroquine alone, and proguanil alone. In addition, 23 (18%) persons who responded were not using any malaria chemoprophylaxis. The risk for malaria was significantly lower among persons using either mefloquine or chloroquine and proguanil (8/88 [9%]) than among persons using the other regimens or no prophylaxis (9/37 [24%]) (RR=0.4; 95% CI=0.2–0.9). Twelve persons not using prophylaxis reported side effects or fear of possible side effects as a reason.

The risk for malaria was lower among persons who reported using bednets all or most of the time (2/27 [7%]) than among persons who sometimes or rarely used bednets (15/99 [15%]) (RR=0.5; 95% Cl=0.1–2.0). The risk for malaria was also lower among persons who consistently used insect repellent in the evening (0/16), compared with those who rarely used repellent (17/110 [15%]) (RR=0; upper 95% confidence limit=1.2). Risk for malaria was not associated with failure to have window or door screens or wear long sleeves or pants in the evening.

As a result of this investigation, EHU staff reviewed with all personnel the need to use and comply with the recommended malaria chemoprophylaxis regimens. EHU staff also emphasized the need to use personal protection measures and made plans to obtain insecticide-impregnated bednets and to provide window and door screens for all personnel.

Reported by: U.S. Embassy Health Unit, Kampala, Uganda; Office of Medical Svcs, Dept of State, Washington, D.C. Malaria Br, Div of Parasitic Diseases, National Center for Infectious Diseases, CDC.

Editorial Note: In Uganda, the increase in malaria among U.S. personnel was attributed to poor adherence to both recommended malaria chemoprophylaxis regimens and use of personal protection measures during a period of increased malaria transmission and intensified chloroquine resistance in sub-Saharan Africa. The findings in this report underscore the need to provide initial and continued counseling regarding malaria prevention for persons living abroad in malaria-endemic areas—preventive measures that are also important for short-term travelers to such areas.

Mefloquine is an effective prophylaxis regimen in Africa and in most other areas with chloroquine-resistant *P. falciparum*; however, in some areas (e.g., Thailand), resistance to mefloquine may limit its effectiveness. In Africa, the efficacy of mefloquine, compared with chloroquine alone, in preventing infection with *P. falciparum* is 92% (1). Mefloquine is safe and well tolerated when given at 250 mg per week over a 2-year period. The risk for serious adverse reactions possibly associated with mefloquine prophylaxis (e.g., psychosis and convulsions) is low (i.e., 1.3–1.9 episodes per 100,000 users [2]), while the risk for less severe adverse reactions (e.g., dizziness,

Malaria — Continued

gastrointestinal complaints, and sleep disturbances) is similar to that for other antimalarial chemoprophylactics (1).

Doxycycline has similar prophylactic efficacy to mefloquine, but the need for daily dosing may reduce compliance with and effectiveness of this regimen (3,4). Chloroquine alone is not effective as prophylaxis in areas of intense chloroquine resistance (e.g., Southeast Asia and Africa). In Africa, for persons who cannot take mefloquine or doxycycline, chloroquine and proguanil is an alternative, although less effective, regimen. Chloroquine should be used for malaria prevention in areas only where chloroquine-resistant *P. falciparum* has not been reported.

Country-specific recommendations for preventing malaria and information on the dosage and precautions for malaria chemoprophylaxis regimens are available from *Health Information for International Travel, 1992* (i.e., "yellow book") (5) or 24 hours a day by telephone or fax, (404) 332-4555.

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Notice to Readers

FDA Approval of Use of a New *Haemophilus* b Conjugate Vaccine and a Combined Diphtheria-Tetanus-Pertussis and *Haemophilus* b Conjugate Vaccine for Infants and Children

Haemophilus influenzae type b (Hib) conjugate vaccines have been recommended for use in infants since 1990, and their routine use in infant vaccination has contributed to the substantial decline in the incidence of Hib disease in the United States (1–3). Vaccines against diphtheria, tetanus, and pertussis during infancy and childhood have been administered routinely in the United States since the late 1940s and has been associated with a greater than 90% reduction in morbidity and mortality associated with infection by these organisms. Because of the increasing number of vaccines now routinely recommended for infants, a high priority is the development of combined vaccines that allow simultaneous administration with fewer separate injections.

The Food and Drug Administration (FDA) recently licensed two new products for vaccinating children against these diseases: 1) the *Haemophilus* b conjugate vaccine (tetanus toxoid conjugate, ActHIB TM),* for vaccination against Hib disease only and 2) a combined diphtheria and tetanus toxoids and whole-cell pertussis vaccine (DTP)

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

Notice to Readers — Continued

and Hib conjugate vaccine (TETRAMUNETM), a combination of vaccines formulated for use in vaccinating children against diphtheria, tetanus, pertussis, and Hib disease.

ActHIBTM

On March 30, 1993, the FDA approved a new *Haemophilus* b conjugate vaccine, polyribosylribitol phosphate-tetanus toxoid conjugate (PRP-T), manufactured by Pasteur Merieux Serum et Vaccins and distributed as ActHIBTM by Connaught Laboratories, Inc. (Swiftwater, Pennsylvania). This vaccine has been licensed for use in infants in a three-dose primary vaccination series administered at ages 2, 4, and 6 months. Previously unvaccinated infants 7-11 months of age should receive two doses 2 months apart. Previously unvaccinated children 12-14 months of age should receive one dose. A booster dose administered at 15 months of age is recommended for all children. Previously unvaccinated children 15-59 months of age should receive a single dose and do not require a booster. More than 90% of infants receiving a primary vaccination series of ActHIB™ (consecutive doses at 2, 4, and 6 months of age) develop a geometric mean titer of anti-Haemophilus b polysaccharide antibody >1 µg/mL (4). This response is similar to that of infants who receive recommended series of previously licensed *Haemophilus* b conjugate vaccines for which efficacy has been demonstrated in prospective trials. Two U.S. efficacy trials of PRP-T were terminated early because of the concomitant licensure of other Haemophilus b conjugate vaccines for use in infants (4). In these studies, no cases of invasive Hib disease were detected in approximately 6000 infants vaccinated with PRP-T. These and other studies suggest that the efficacy of PRP-T vaccine will be similar to that of the other licensed Hib vaccines.

TETRAMUNETM

On March 30, 1993, the FDA approved a combined diphtheria and tetanus toxoids and whole-cell pertussis vaccine (DTP) and *Haemophilus* b conjugate vaccine. TETRA-MUNE $^{\text{TM}}$, available from Lederle-Praxis Biologicals (Pearl River, New York), combines two previously licensed products, DTP (TRIIMMUNOL $^{\text{®}}$, manufactured by Lederle Laboratories [Pearl River, New York]) and *Haemophilus* b conjugate vaccine (HibTITER $^{\text{®}}$, manufactured by Praxis Biologics, Inc. [Rochester, New York]).

This vaccine has been licensed for use in children aged 2 months–5 years for protection against diphtheria, tetanus, pertussis, and Hib disease when indications for vaccination with DTP vaccine and *Haemophilus* b conjugate vaccine coincide. Based on demonstration of comparable or higher antibody responses to each of the components of the two vaccines, TETRAMUNETM is expected to provide protection against Hib, as well as diphtheria, tetanus, and pertussis, equivalent to that of already licensed formulations of other DTP and *Haemophilus* b vaccines.

The Advisory Committee for Immunization Practices (ACIP) recommends that all infants receive a primary series of one of the licensed *Haemophilus* b conjugate vaccines beginning at 2 months of age and a booster dose at age 12–15 months (5). The ACIP also recommends that all infants receive a four-dose primary series of diphtheria and tetanus toxoids and pertussis vaccine at 2, 4, 6, and 15–18 months of age, and a booster dose at 4–6 years (6–8). A complete statement regarding recommendations for use of ActHIBTM and TETRAMUNETM is being developed.

Notice to Readers — Continued

Reported by: Office of Vaccines Research and Review, Center for Biologics Evaluation and Research, Food and Drug Administration. Div of Immunization, National Center for Prevention Svcs; Meningitis and Special Pathogens Br, Div of Bacterial and Mycotic Diseases, National Center for Infectious Diseases, CDC.

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The data in the weekly *MMWR* are provisional, based on weekly reports to CDC by state health departments. The reporting week concludes at close of business on Friday; compiled data on a national basis are officially released to the public on the succeeding Friday. Inquiries about the *MMWR* Series, including material to be considered for publication, should be directed to: Editor, *MMWR* Series, Mailstop C-08, Centers for Disease Control and Prevention, Atlanta, GA 30333; telephone (404) 332-4555.

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