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# **Current Trends**

# Repeat Injuries in an Inner City Population – Philadelphia, 1987–1988

Findings from the Philadelphia Injury Prevention Program (PIPP) indicate that repeat injuries account for 10% of visits to emergency rooms (ERs) for injuries. Because understanding of the epidemiology of repeat injuries is limited, data from PIPP were used to examine this problem. PIPP is a cooperative effort by the Philadelphia Department of Public Health, the University of Pennsylvania School of Medicine, and CDC. PIPP maintains surveillance of virtually all emergency-room (ER)–treated injuries sustained by 68,103 inner city residents living in 17 census tracts in Philadelphia. This report focuses on repeat injuries among this predominantly (97%) black, indigent population (estimated 1987 median household income: \$11,810) (Donnelley Marketing Information Services, unpublished data).

An injury was defined as a visit to an ER in Philadelphia for care or evaluation of an injury. A repeat injury was defined as a discrete injury in the same person after an initial injury during a 1-year surveillance period (March 1, 1987, to February 29, 1988). Excluded from this definition were visits to an ER for follow-up care of an injury (e.g., suture removal, cast replacement, or wound redressings).

During the study period, the surveillance system identified 9567 injuries in 8600 persons (12.6% of the study population). Of these, 2161 (22.6%) resulted from falls, 1960 (20.5%) from interpersonal violence, 1500 (15.7%) from motor vehicle-related incidents, 965 (10.1%) from unintentional blunt trauma, and 895 (9.4%) from unintentional cuts. The remaining 2086 injuries (21.8%) resulted from other causes.

Of the 9567 injuries, 967 (10.1%) were repeat injuries to 802 (9.3%) persons. Of the repeat injuries, 268 (27.7%) resulted from interpersonal violence, 193 (20.0%) from falls, 111 (11.5%) from motor vehicle-related incidents, 93 (9.6%) from unintentional blunt trauma, and 82 (8.5%) from unintentional cuts. The remaining 220 (22.8%) resulted from other causes.

Of the 802 persons with repeat injuries, 521 (65.0%) were male. Two hundred eighty-two (35.2%) were 0–19 years of age; 499 (62.2%) were 20–64; and 21 (2.6%) were  $\geq$ 65. Six hundred eighty-three (85.2%) persons were injured twice; 85 (10.6%), three times; 25 (3.1%), four times; six (0.7%), five times; and three (0.4%), six times. Seven persons died from a repeat injury.

### Repeat Injuries - Continued

The cause of initial injury differed little between persons with repeat injuries and those with one injury (Figure 1). Of the 802 persons with repeat injuries, 224 (27.9%) had two or more injuries from a similar cause. Interpersonal violence was the most common cause of a repeat injury resulting from a similar cause. Of the 184 persons with repeat injuries and at least one injury from interpersonal violence, 88 (47.8%) had at least one additional injury from interpersonal violence and 15 had three or more injuries from interpersonal violence. Of the 172 persons whose first injury resulted from a fall, 67 (39.0%) fell again. Among persons  $\geq$ 65 years of age with repeat injuries, 61.5% with an initial fall injury suffered a subsequent fall injury.

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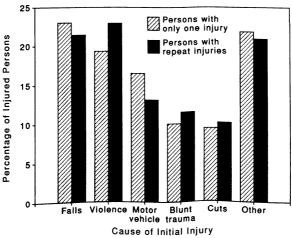
**Editorial Note:** Preliminary results from PIPP surveillance found that in 1 year 12.6% of the study population visited an ER for treatment of injuries caused primarily by falls, interpersonal violence, and motor vehicle incidents. The finding that 9.3% of persons who came to the ER for treatment of an injury returned within 1 year for treatment of another injury has important implications for injury prevention.

Falls and interpersonal violence were the main causes of both initial and repeat injuries. The PIPP data on repeat fall injuries are consistent with other studies that found that once an older person has fallen, the risk of falling again increases substantially (1-3). Accordingly, one aspect of a fall-prevention program for older persons should focus on persons with histories of previous falls.

The occurrence of repeat injuries from interpersonal violence is probably underestimated because such injuries may not have been specifically identified on the ER record as having resulted from interpersonal violence. In particular, many women and children diagnosed as having unintentional cuts, blunt trauma, or falls may have been injured intentionally.

Hospital emergency departments are an important interface between the healthcare system and victims of interpersonal violence. One cause of repeat violent injury is interpersonal violence associated with family and intimate relationships (e.g.,





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# Repeat Injuries - Continued

spouse and child abuse). The introduction of standard protocols in hospital emergency departments for the proper identification, treatment, and referral of victims of family violence has been advocated as a strategy for preventing repeat injury and other consequences associated with family violence (4). This strategy has proven useful in the identification of battered women among trauma patients in emergency departments (5). These strategies can lower mortality and morbidity associated with family violence but require further evaluation.

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# Epidemiologic Notes and Reports

# Group A Beta-Hemolytic Streptococcal Bacteremia - Colorado, 1989

From January through August 1989, group A beta-hemolytic *Streptococcus* (GABHS) was isolated from blood cultures obtained from 19 patients who had been admitted to a hospital in metropolitan Denver, Colorado. In comparison, this organism was cultured from blood from eight patients in 1988 and three in 1987 at this hospital (Figure 1). The Colorado Department of Health was notified of this increase, and in August, the department reviewed these patients' medical charts to describe GABHS bacteremia cases and to determine whether they represented community-acquired disease, nosocomial acquisition, or laboratory artifact.

Patients ranged in age from 3 weeks to 96 years (median: 67 years); 12 were male. All but one were residents of Colorado when hospitalized; 16 patients lived in the Denver metropolitan area. Twelve patients were admitted from private residences and four from nursing homes; three were transferred from acute- or extended-care facilities.

For 13 patients, a blood specimen was obtained within 6 hours of arrival at the hospital; 12 patients had presenting manifestations consistent with bacteremia or sepsis (primarily fever, chills, rigors, and a focus of infection). In only three patients was bacteremia first documented >48 hours after hospitalization.

Seven patients had no identified source of bacteremia; four had a possible cutaneous source of infection (cellulitis, impetigo, open sores, or an abscess); seven had pneumonia, and one had both cutaneous and respiratory infections. Two of the seven without identified sources of infection developed GABHS bacteremia following

### Streptococcal Bacteremia - Continued

major trauma in motor vehicle crashes. A third patient developed GABHS bacteremia following elective laser hemorrhoidectomy. Two patients developed acute respiratory distress syndrome; two developed acute renal failure; and nine (47%) died after developing GABHS bacteremia.

Eight group A *Streptococcus* blood culture isolates from this hospital were sent to CDC for typing. Three were M-type 1, T-type 1; three were M-type 3, T-type 3/13; and two were not typable.

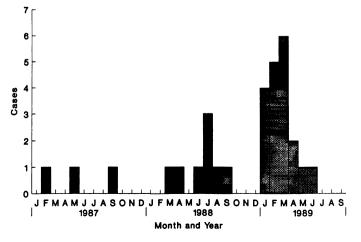
Retrospective surveys of this and 17 other metropolitan Denver hospital microbiology laboratories identified 73 cases of GABHS bacteremia between January and August—a rate of seven cases per 100,000 per year in the general population of the Denver metropolitan area.

Active surveillance for GABHS bacteremia was established in September 1989 in metropolitan Denver. Bacterial isolates are sent to CDC for analysis, and ongoing case investigation is examining potential explanations for the increasing incidence of group A streptococcal disease and possible risk factors, such as age, underlying illnesses, socioeconomic status, and race/ethnicity.

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**Editorial Note:** During the last half century, the incidence of severe infections with group A *Streptococcus* (including bacteremia, puerperal sepsis, and scarlet fever) and nonsuppurative sequelae (such as rheumatic fever) has decreased markedly in the United States. Potential explanations for this decline may relate to antibiotic therapy, improved living conditions, and decreased virulence of the infecting organism (1). Recently, however, group A streptococcal infections have re-emerged as a public health problem. In 1985 and 1986, clusters of patients with rheumatic fever were reported from several areas (2-6), and since 1987, a severe syndrome associated with group A streptococcal infection (streptococcal toxic-shock–like syndrome) has been recognized (7,8).





# Streptococcal Bacteremia - Continued

The findings in Denver are consistent with a trend toward increasing rates of GABHS bacteremia. While a direct comparison of the GABHS bacteremia incidence rate in Denver for 1989 and for previous years is not possible because of incomplete laboratory records, the 1989 rate is more than double the most recent population-based estimate of three cases per 100,000 persons per year from Charleston County, South Carolina (1985–1987) (CDC, unpublished data). In addition, during 1989, CDC received an increased number of reports of GABHS bacteremia from other areas in the United States and serotyped strains from several Scandinavian countries that have increased rates of GABHS bacteremia. These reports suggest widespread changes in the epidemiology of group A streptococcal disease.

The descriptive epidemiology of GABHS bacteremia in the Denver patients is similar to that in previous reports (9,10). Even though all age groups may be affected, disease occurs primarily in the elderly, including nursing home residents. Underlying medical conditions in affected persons may include chronic debilitating disease, immunosuppressive illness or medication, and intravenous-drug use. In addition, surgery, traumatic injury, or other disruption of the cutaneous barrier may predispose persons to invasive infection. Meningitis, endocarditis, osteomyelitis, septic arthritis, and genitourinary infections can occur in association with bacteremia. Despite appropriate therapy, illness can progress to shock, disseminated intravascular coagulation, and death; reported mortality rates range from 5% to 45% (9).

Streptococcal toxic-shock-like syndrome also has been reported recently from the Rocky Mountain area (7). This condition has been associated with pyrogenic exotoxin-producing group A streptococcal strains. Clinical features include fever, shock, localized erythema, renal failure, severe tissue injury (myositis and fasciitis), and adult respiratory distress syndrome; bacteremia may occur and illness is often fatal (7,8). The clinical manifestations in some of the Denver patients were compatible with this syndrome.

Factors contributing to the apparent recent increase in GABHS bacteremia are unclear. No single serotype, to suggest a common source, was found in the isolates from Denver. Serologic typing of group A streptococcal strains based on antigenic differences in the M-protein suggests that an increase in the proportion of virulent organisms may play a role. M-types 1 and 3, comprising 75% of the Denver isolates, may be more invasive and more likely to cause clusters of infection than most other streptococcal M-types (*11*). Analysis of serotyping data from isolates submitted to CDC since 1972 shows an increase in the proportion of these two M-types in the 1980s (*11*).

Efforts to prevent and control invasive group A streptococcal disease should be directed at surveillance to establish the incidence of this problem in different geographic areas; further epidemiologic studies of toxic-shock-like syndrome; and improved understanding of the roles of immune responses, exotoxin, and other virulence factors. Physicians should continue to diagnose and treat mild streptococcal infections to prevent their progression to severe invasive disease.

On July 14, 1989, CDC notified state and territorial health officials of a possible increase in severe group A streptococcal infections and requested that clusters of invasive group A streptococcal infection or illness in previously healthy persons be reported through state health departments to the Respiratory Diseases Branch, Division of Bacterial Diseases, Center for Infectious Diseases, CDC; telephone (404) 639-3021.

### Streptococcal Bacteremia - Continued

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(Continued on page 11)

### TABLE I. Summary – cases of specified notifiable diseases, United States

	1:	at Week Endi	ng	Cumulat	Cumulative, 1st Week Ending				
Disease	Jan. 6, 1990	Jan. 7, 1989	Median 1985-1989	Jan. 6, 1990	Jan. 7, 1989	Median 1985-1989			
Acquired Immunodeficiency Syndrome (AIDS) Aseptic meningitis Encephalitis: Primary (arthropod-borne	1,821 63	U* 63	266 63	1,821 63	462 63	266 63			
& unspec) Post-infectious Gonorrhea: Civilian	10 3	9 1	12 1	10 3	9 1	12 1			
Hepatitis: Type A	11,354 121 273	10,920 145 397	11,677 164 278	11,354 121 273	10,920 145 397	11,677 164 278			
Type B Non A, Non B	182 25	263 52	266 52	182 25	263 52	266 52			
Unspecified Legionellosis Leprosy	24 12	43 10 7	43 10	24 12	43 10 7	43 10			
Malaria Measles: Total <sup>†</sup>	10 88	8 40	8 9	10 88	, 40	8 9 8			
Indigenous Imported	80 8	40	8	80 8	40	-			
Meningococcal infections Mumps Pertussis	20 31 16	32 72 49	32 72 26	20 31 16	32 72 49	32 72 26			
Rubella (German measles) Syphilis (Primary & Secondary): Civilian	3 349	-3 461	20 3 461	3 349	46 <u>1</u>	3 461			
Military Toxic Shock syndrome Tuberculosis	3 233	4 333	2 4 141	3 233	4 333	2 4 141			
Tularemia Typhoid Fever Typhus fever, tick-borne (RMSF)	1 4 -	2 3 2	- 3 1	1 4	2 3 2	- 3 1			
Rabies, animal	27	47	47	27	47	47			

### TABLE II. Notifiable diseases of low frequency, United States

	T		
	Cum. 1990		Cum. 1990
Anthrax		Leptospirosis	
			1 -
Botulism: Foodborne	- I -	Plague	- I -
Infant		Poliomyelitis, Paralytic, <sup>3</sup>	- 1
Other	-	Psittacosis (N.C. 1, Ariz. 1)	2
Brucellosis (Calif. 2)	2	Rabies, human	· ·
Cholera	-	Tetanus	
Congenital rubella syndrome	-	Trichinosis (N.J. 1)	1
Congenital syphilis, ages < 1 year			1
Diphtheria	-		
	1	1	

\*Because AIDS cases are not received weekly from all reporting areas, comparison of weekly figures may be misleading. Four of the 88 reported cases for this week were imported from a foreign country or can be directly traceable to a known internationally imported case within two generations.

<sup>5</sup>No cases of suspected poliomyelitis have been reported in 1990; none of the 13 suspected cases in 1989 have been confirmed to date.

	r	Acontia	Encephalitis					enatitie	(Viral), by	type	<u> </u>	<u> </u>
Reporting Area	AIDS	Aseptic Menin- gitis	Primary	Post-in- fectious	Gona (Civi	rrhea ilian)	A .	в	NA,NB	Unspeci- fied	Legionel- losis	Leprosy
	Cum. 1990	Cum. 1990	Cum. 1990	Cum. 1990	Cum. 1990	Cum. 1989	Cum. 1990	Cum. 1990	Cum. 1990	Cum. 1990	Cum. 1990	Cum. 1990
UNITED STATES	1,821	63	10	3	11,354	10,920	273	182	25	24	12	-
NEW ENGLAND	50	2	1	-	436	473	4	17	1	1	-	-
Maine N.H.	- 16	- 1	-		4	4	-	-	:	-		
Vt.	-	-	-	-	3	-	-	-	-	-	-	-
Mass. R.I.	1	1	1	-	91 38	153 25	4	17	1	1		-
Conn.	33	-	-		300	291	-	-				
MID. ATLANTIC	985	-	-		487	1,323	9	10	-	-	3	-
Upstate N.Y.	30	-	-	-	-	- 650	-	-	-	-	-	-
N.Y. City N.J.	790 131	-	-	-	366	215	9	10	-	-	3	-
Pa.	34	-	-	-	121	458	-	-	•	-	-	-
E.N. CENTRAL	116	7	-	1	2,780	1,690	13	18	4	2	3	-
Ohio Ind.	24 22	3	-	1	1,019 375	466	5	5	3	1	3	-
111.	67	-	-	-	909	480	-	-	-	-	•	-
Mich. Wis.	1 2	4	-	-	466 11	522 222	8	13	1	1		-
		-	-	•	562	223	4	3	1			
W.N. CENTRAL Minn.	2	-	-		562 89	47	4	-	-			-
lowa	-	-	-	-	124	21	-	1	-	•	-	-
Mo. N. Dak.	2	•	-	-	347	149 2	-	-	-		-	
S. Dak.	-	-	-	:	2	4	1	-	1	-	-	-
Nebr. Kans.	-		-	:	-	-	3	2	-	-	-	-
S. ATLANTIC	270	17	3		4,074	3,275	44	47	3	1		
Del.	279	1	-		34	43	2	-	-	-		
Md.	24	5	1	-	182	200	18	9	-	1	-	-
D.C. Va.	43 121	1	-	-	185 58	183	1	1	-			
W. Va.	-	-	-		20	65	-	-	-	-	-	•
N.C. S.C.	54 25	4	1		403 534	636 699	9 4	13 13	2	:		
Ga.	8	-	1	-	897	489	-	-	-	-	-	-
Fla.	4	6	-	-	1,761	960	10	11	1	-	-	-
E.S. CENTRAL Ky.	38	6	1	-	618 64	1,131 82	7 2	18 4	1 1	1 1	4 1	-
Tenn.	10 25	-		-	117	300	1	10	-	-	2	
Ala. Miss.	3	5	1	-	238 199	471 278	4	4	-	-	1	-
	-	1	-	-			-	-	-	-		-
W.S. CENTRAL Ark.	40	2	-	-	778 199	971 93	16 3	2	-		1	-
La.	26	-	-	-	150	120	-	-	-		-	-
Okla. Tex.	9 5	2		-	88 341	142 616	13	2	-	-	1	-
MOUNTAIN	71	2	1	-	213	80	32	19	_	3		-
Mont.		-	-	-	213	3	1	1	-	-		
ldaho Wyo.	1	-	-		1	7 1	4	2	-	-	-	-
Colo.	22	-	-		2	-	-	-	-	-	-	:
N. Mex.	-	-	-	-	10	19	5	2	-	-	-	-
Ariz. Utah	32 8		1	-	133 13	6 7	18	7	-	-		-
Nev.	8	2	-		51	37	4	7	-	3	-	-
PACIFIC	240	27	4	2	1,406	1,754	144	48	15	16	1	-
Wash. Oreg.	75	-	-	:	- 57	102 62	- 5	- 3	-	-	-	-
Calif.	14 151	27	4	2	1,320	1,559	139	44	15	16	- 1	-
Alaska Hawaii	-	-	-	-	29	29	-	1	-	-	-	-
	-	-	-	-	-	2	-	-	-	•	-	-
Guam P.R.	- 1	-	-	-	-	4 22	-	-	-		-	-
V.I.	-	-	-	-	-	7	-	-	-	-	-	-
Amer. Samoa C.N.M.I.	-	-	-	-	-	1	:	:	-	-	-	-
	-	-	-	-	-	1	-	-	-	-	-	-

# TABLE III. Cases of specified notifiable diseases, United States, weeks ending January 6, 1990 and January 7, 1989 (1st Week)

N: Not notifiable

Reporting Area	Malaria Measles (Rubeola)					Menin- gococcal				Pertussi	-	Rubelia			
	Ivialaria	Indigenous		Impo	rted*	Total	Infections	Mumps		r er cusers					
	Cum. 1990	1990	Cum. 1990	1990	Cum. 1990	Cum. 1989	Cum. 1990	1990	Cum. 1990	1990	Cum. 1990	Cum. 1989	1990	Cum. 1990	Cum. 1989
UNITED STATES	10	80	80	8	8	40	20	31	31	16	16	49	3	3	3
NEW ENGLAND	2	•	-	-	-	•	2	•	-	-	-	6	•	•	•
Maine N.H.		:	:	:	:	:	1	:	:	:		2 3	:	:	2
Vt.	-	•	-	-	•	-	:	•	•	•	•	-	•	-	-
Mass. R.I.	2	:	:	:	-	:	1	:	:	-	:	1	:	:	-
Conn.	-	•	-	•	-	•	•	•	•	•	-	-	•	•	•
MID. ATLANTIC	•	11	11	4	4	•	2	1	1	2	2	12	-	•	-
Upstate N.Y. N.Y. City		:		:	-	-	-	:	:	:	:	-	:		:
N.J. Pa.	-	11	11	- 45	4	•	2	1	1	2	2	11	•	·	-
E.N. CENTRAL	-			43	-	•	3	3	3	1	1	1 6	•	•	-
Ohio		:	:	-		-	1	3	3			1		:	-
Ind.	•	·	•	•	-	-	:	-	•	•	•	:	•	•	-
III. Mich.	-	:		:	-	:	1	3	3	1	1	1	:		-
Wis.	-	•	•	•	-	•	-	-	•	-	-	4	-		-
W.N. CENTRAL Minn.	-	10	10	-	•	30	1	•	-	•	•	2		-	-
lowa	-	10	10		-			:		-	:	2	:	-	:
Mo. N. Dak.	•	•	-	-	-	30	•	-	-	-	-	-	-	-	-
S. Dak.		:	-		:	:	1	:	-	2		-	:	-	2
Nebr. Kans.	-	•	-	:	-	-	•	-	•	-	-	-	•	-	-
	4		•			•	•	-	•	-	•	-	-	-	-
S. ATLANTIC Del.	4	1	1	3	3	-	3	14	14	1	1	1		-	
Md.	2	1	1	2†	2	-		5	5	-	-	-	-	-	-
D.C. Va.	1			1†	1	-	-	2	2	- 1	1	1	:	-	:
W. Va.	•	•	-	-	-	-	:	-	-	•	-	-		-	-
N.C. S.C.	-		:		:	-	1	4	4	-	:	-	:	-	-
Ga.		-	-	-	-	5	-	-	-	•	-	-	-	-	-
Fla.	1	-	•	-	•	-	1	2	2	•	•	-	-	-	-
E.S. CENTRAL Ky.	1			:	:	:		-	-	3	3	1	-	-	-
Tenn.	-	•	•	-	-	-	-	-	-	-	-	- 1		-	-
Ala. Miss.	1		:	-	-	:		N	N	3	3	1 '	•	-	-
W.S. CENTRAL				_			-	1	1	•	-	-	-	-	
Ark.	-	-	-	-	-		-	1	1	-	:	-	2	:	-
La. Okla.	:	:	-	-	-	-	-	:	-	-	-	-	-	•	-
Tex.			-	-	-	-		-	-	-	:	-	:	-	-
MOUNTAIN	-		-		-	1	-	3	3	1	1	2	-		-
Mont. Idaho		•	-	•	•	-	-		-	-	•	-	-	-	-
Wyo.	-			:		-		2	2	-	-	-	-	-	-
Colo. N. Mex.	-	•	-	•	•	-	-		-	•	-	1	-	-	-
Ariz.	-			:	:	1		N 1	N 1	1	1	- 1	-	-	-
Utah Nev.	-	-	-	•	•	-	-	-	-	-	-	•	-	-	-
PACIFIC	-					-	-	-	•	-	•	-	-	•	•
Wash.	3	58	58	1	1	9	9	9	9	8	8	19	3	3	3
Oreg. Calif.	-	-	-	•	:		1	N	N	1	1		-	-	-
Alaska	3	58	58	1†	1	9	8	9	9	7	7	19	3	3	3
Hawaii	-	U		υ	-	•		Ű	-	Ū	-	-	U	-	-
Guam	-	U	•	U	•		-	U	-	υ	-	-	υ	-	-
P.R. V.I.	-	Ū	•	Ū	:	23		2 U	2	Ū	-	-	Ū	•	-
Amer. Samoa	-	U	•	U	•			U	:	U	:	:	υ	-	:
C.N.M.I.	•	Ū	-	U	•	-	•	U	•	υ	-	-	υ	-	-

# TABLE III. (Cont'd.) Cases of specified notifiable diseases, United States, weeks ending January 6, 1990 and January 7, 1989 (1st Week)

\*For measles only, imported cases includes both out-of-state and international importations. N: Not notifiable U: Unavailable <sup>†</sup>International <sup>‡</sup>Out-of-state

Reporting Area	Syphilis (Civilian) (Primary & Secondary)		Toxic- shock Syndrome	Tuber	culosis	Tula- remia	Typhoid Fever	Typhus Fever (Tick-borne) (RMSF)	Rabies, Animal
	Cum. 1990	Cum. 1989	Cum. 1990	Cum. 1990	Cum. 1989	Cum. 1990	Cum. 1990	Cum. 1990	Cum. 1990
UNITED STATES	349	461	3	233	333	1	4	•	27
NEW ENGLAND	20	12	1	-	2	-	-	-	•
Maine N.H.	:	-	-	2	:	:	:	-	
Vt.	-			-	-	•	-	-	-
Mass. R.I.	6	12	1	:	:		-	-	•
Conn.	14	-		-	2	:	-		
MID. ATLANTIC	40	82		87	54	1	1		9
Upstate N.Y.	-	-	-			•	•	-	-
N.Y. City N.J.	40	8 29	:	77	35 10	1	1		- 9
Pa.		45	-	10	9	-	-		
E.N. CENTRAL	33	15		27	12		-		-
Ohio	12	•	-	-	4	-	-		-
Ind. III.	1 16	2 8	:	22	1 6	:	:		
Mich.	3	3	•	-			-	-	
Wis.	1	2	-	5	1	-	-	-	-
W.N. CENTRAL	5	1	1	9	3	-	-	-	5
Minn.	1		-	4	1	•	-	-	4
lowa Mo.	4	1	-	1	1		-	-	-
N. Dak.	-	-	-	1	1	-	-	-	-
S. Dak.	-	-	:	1	-	-	-	-	•
Nebr. Kans.	-	-	1	2	-	-	-	-	- 1
S. ATLANTIC	127	208		18	40				6
Del.	5	208	-	1	40		-	-	-
Md.	9	23	-	7	-	•	-	-	-
D.C. Va.	28	18 7		-	4 11	-	-	-	2
W. Va.	- 28	· ·	-	2	1		-	-	2
N.C.	14	15	-		10	-	-	-	-
S.C. Ga.	28	4 38		8	14		-	-	2 2
Fla.	43	102	-	-	-	-	-	-	-
E.S. CENTRAL	28	46	1	8	13		-		2
Ky.	-	-		4	5	-	-	-	ĩ
Tenn.	17	34	- 1	4	7	•	-	-	
Ala. Miss.	11	12	-	4	1		-	-	1
W.S. CENTRAL	35	19		12			_		3
Ark.	4	-	-	9		-	-		1
La.	21	11	•	-	-	•	-		-
Okla. Tex.	10	8	-	3			-		2
MOUNTAIN	5	2		2					
Mont.	-	2	-	2	-		-	-	1
Idaho	1	-	-		-	-	-	-	-
Wyo. Colo.	-	-	-	•	-	-	-	•	-
N. Mex.	-	-	-	1	-		-	-	1
Ariz.	3	-	-	-	-	-	-	-	
Utah	1	2	-		-	-	-	•	•
Nev.			-	1	-	-	-	-	-
PACIFIC Wash.	56	76 6	-	70 3	209	•	3	-	1
Oreg.	1	3	-	2					-
Calif.	55	67	-	65	209	-	3	-	-
Alaska Hawaii	-		-	-		:	•	-	1
Guam			-	-	-	-	•	-	-
P.R.	-	-		-	:		•	-	-
V.I.	-	-	-	-	:				9
Amer. Samoa C.N.M.I.	-	-		-	-	-	•	•	•
G.N.W.L		-	•	-	•	-	-	•	-

# TABLE III. (Cont'd.) Cases of specified notifiable diseases, United States, weeks ending January 6, 1990 and January 7, 1989 (1st Week)

U: Unavailable

### All Causes, By Age (Years) All Causes, By Age (Years) P&I\*\* P&I\*\* **Reporting Area** Reporting Area All All 25-44 Total ≥65 45-64 1-24 45-64 <1 ≥65 25-44 1-24 <1 Total Ages Ages NEW ENGLAND S. ATLANTIC 1.324 Boston, Mass. Atlanta, Ga. Bridgeport, Conn. Baltimore, Md. Cambridge, Mass. з Charlotte, N.C. Fall River, Mass. Jacksonville, Fla. Hartford, Conn. Miami, Fla. Lowell, Mass. Norfolk, Va. Lvnn, Mass. Richmond, Va. New Bedford, Mass. Savannah, Ga. New Haven, Conn. St. Petersburg, Fla. 7 Providence, R.I. Tampa, Fla. Somerville, Mass. Washington, D.C. Springfield, Mass. Wilmington, Del. Waterbury, Conn. E.S. CENTRAL Worcester, Mass. Birmingham, Ala. MID. ATLANTIC 2.877 .939 Chattanooga, Tenn. Albany, N.Y. Knoxville, Ťenn. Allentown, Pa. Louisville, Ky. Buffalo, N.Y. Memphis, Tenn. Camden, N.J. Mobile, Ala. Elizabeth, N.J. Λ Montgomery, Ala. Erie, Pa.t Nashville, Tenn. Jersey City, N.J.§ W.S. CENTRAL 1,868 N.Y. Ćity, Ń.Y. 1.698 1.087 Newark, N.J. Austin, Tex. Baton Rouge, La. Paterson, N.J. ž Corpus Christi, Tex.§ Philadelphia, Pa. з ĩ Dallas, Tex. Pittsburgh, Pa.† El Paso, Tex. Reading, Pa. ā Fort Worth, Tex Rochester, N.Y Houston, Tex.§ Schenectady, N.Y. Little Rock, Ark. Scranton, Pa.† з з New Orleans, La. Syracuse, N.Y. San Antonio, Tex. Trenton, N.J. Shreveport, La. Utica, N.Y. Tulsa, Okla. ġ Yonkers, N.Y. MOUNTAIN E.N. CENTRAL 2,533 1,731 Albuquerque, N. Mex. Akron, Ohio Colo. Springs, Colo. Canton, Ohio Δ Denver, Colo. Chicago, III.§ Las Vegas, Nev. Cincinnati, Ohio Cleveland, Ohio Ogden, Utah Phoenix, Ariz. Columbus, Ohio Pueblo, Colo. Dayton, Ohio ·6 Salt Lake City, Utah Detroit, Mich. Tucson, Ariz. Evansville, Ind. Δ Fort Wayne, Ind. PACIFIC 1.930 1,298 Gary, Ind. Berkeley, Calif. Grand Rapids, Mich. Fresno, Calif. Indianapolis, Ind. Glendale, Calif. Madison, Wis. Honolulu, Hawaii Milwaukee, Wis. Δ Long Beach, Calif.§ з Peoria, III. Los Angeles Calif. Rockford, III. Oakland, Calif. South Bend, Ind. з Pasadena, Calif. Toledo, Ohio Portland, Oreg. Youngstown, Ohio Sacramento, Čalif. San Diego, Calif. W.N. CENTRAL San Francisco, Calif. Des Moines, Iowa San Jose, Calif. Duluth, Minn. Kansas City, Kans. Seattle, Wash. Kansas City, Mo. Spokane, Wash. Δ Tacoma, Wash. Lincoln, Nebr. Λ Minneapolis, Minn. 13,684<sup>††</sup> 9,215 2,622 1,130 TOTAL Omaha, Nebr. St. Louis, Mo. Ì. St. Paul, Minn. Wichita, Kans.

## TABLE IV. Deaths in 121 U.S. cities,\* week ending January 6, 1990 (1st 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.

†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.

t†Total includes unknown ages.

§Data not available. Figures are estimates based on average of past available 4 weeks.

# Streptococcal Bacteremia – Continued

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# Group A Beta-Hemolytic Streptococcal Pharyngitis Among U.S. Air Force Trainees – Texas, 1988–89

In December 1988 and January 1989, an outbreak of pharyngitis caused by group A beta-hemolytic *Streptococcus* (GABHS) occurred among military trainees at Lackland Air Force Base, Texas. From January through November 1988, the incidence of culture-positive GABHS pharyngitis was 2.1 cases per 1000 trainees per month. By comparison, in December and January, the rates were 9.5 per 1000 and 18.7 per 1000, respectively. The outbreak prompted the administration of penicillin prophylaxis to all trainees (>6000)—the first time in at least 15 years that a mass prophylaxis program was implemented at this Air Force base.

Between December 23 and January 10, GABHS was isolated from throat swabs of 186 trainees from 13 flights (one flight = approximately 50 trainees) in four of eight squadrons (one squadron = 16–20 flights). Each flight occupies a single open-bay sleeping area; each squadron occupies a single dormitory building. The outbreak was detected by a surveillance system for streptococcal pharyngitis: when streptococcal pharyngitis is detected in three or more trainees in a flight within a 7-day period, throat cultures are obtained from all members of that flight. If GABHS is isolated from >10% of throat cultures, all persons in the flight who are not allergic to penicillin are given penicillin prophylaxis.

In the last week of December and first 2 weeks of January, 11 flights from three squadrons exceeded the threshold for prophylaxis. Six additional flights with less than three trainees with streptococcal pharyngitis were screened; two of those flights exceeded the threshold. Throat cultures were positive for 17% (17/101) of a sample of male trainees in the first half of their 4-week training course, compared with 31% (101/330) of a sample of trainees in the second half (relative risk = 1.8; 95% confidence interval = 1.1-2.9). All women were in the second half of their training.

In the second week of January, benzathine penicillin G (1.2 million units IM) was administered to >6000 trainees already on the base. In addition, penicillin prophylaxis was initiated for all nonallergic, incoming personnel during their second week of training. During the third week of January, no flights exceeded the threshold. However, after two flights had three or more positive throat cultures in the fourth week of January, timing of prophylaxis was changed to the first week of training; subsequently, incidence rates of GABHS pharyngitis decreased markedly (Figure 1).

Routine prophylaxis was discontinued on April 30. In July, one flight exceeded the threshold for prophylaxis. No cases of acute rheumatic fever or other sequelae have been reported from Lackland Air Force Base or secondary training bases.

Reported by: AH Mumm, COL, MC; CA Smith, LT COL, BSC, Epidemiology Div, US Air Force School of Aerospace Medicine, Human Systems Div, Brooks Air Force Base, CE Gookins, MAJ,

### Streptococcal Pharyngitis - Continued

BSC, Environmental Health Office, Wilford Hall US Air Force Medical Center, Lackland Air Force Base, Texas. Respiratory Diseases Br, Div of Bacterial Diseases, Center for Infectious Diseases, CDC.

**Editorial Note:** During World Wars I and II, GABHS disease caused substantial morbidity among recruits (1), probably because of the introduction of large numbers of susceptible persons into crowded environments. A marked decrease in the incidence of GABHS disease and sequelae in military recruits occurred from 1965 to 1985 (1). This coincided with penicillin prophylaxis programs to prevent GABHS disease in military trainees and with a nationwide decrease in the incidence of rheumatic fever (2). By 1979, most military training centers had discontinued year-round prophylaxis of incoming recruits against GABHS disease (1).

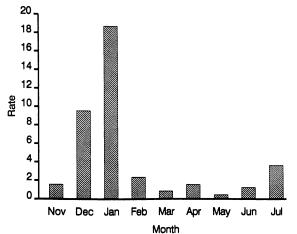
This investigation and other recent reports demonstrate that military training centers remain at risk for outbreaks of GABHS-related disease and underscore the importance of surveillance in these settings (3,4). Although outbreaks of GABHS infections can be seasonal and self-limited, the abrupt decrease in incidence after January suggests that surveillance and antibiotic prophylaxis were important in limiting this outbreak. During other outbreaks of GABHS pharyngitis in military recruits, mass prophylaxis has been used successfully (5,6).

Some military training centers have policies for use of penicillin prophylaxis when the incidence of GABHS infections exceeds a specified threshold. The Armed Forces Epidemiological Board has suggested that when the incidence of GABHS disease exceeds 10 cases per 1000 trainees per week, epidemics of acute rheumatic fever may occur (7). The appropriateness of such policies for other institutional settings requires evaluation. State health departments are requested to notify CDC's Respiratory Diseases Branch, Division of Bacterial Diseases, Center for Infectious Diseases, at (404) 639-3021 about outbreaks of GABHS infections and/or their sequelae in such settings.

### References

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FIGURE 1. Rate of group A beta-hemolytic streptococcal pharyngitis per 1000 trainees per month – Lackland Air Force Base, Texas, November 1988–July 1989



# Streptococcal Pharyngitis - Continued

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International Notes

# Epidemic Meningococcal Disease – Kenya and Tanzania: Recommendations for Travelers, 1990

Epidemic meningococcal disease has occurred in Nairobi, Kenya, and the Arusha area of northern Tanzania. From April through October 1989, approximately 4000 cases of meningitis due to *Neisseria meningitidis* serogroup A were reported in Nairobi, an attack rate of 267 per 100,000 during this period. Cases have also been reported in other regions of Kenya. Attack rates in Nairobi were equally distributed by sex; the highest rate occurred among children 0–4 years of age. Control measures in Nairobi included a vaccination program directed at persons in areas most affected by disease; the incidence appears to be decreasing.

In Tanzania, from July through November 1989, 1249 cases of meningococcal disease were reported in four northern districts, including the Arusha area – an attack rate of 172 per 100,000. The Tanzanian Ministry of Health has implemented control measures and has reported a decrease in meningococcal disease. No cases of meningococcal disease have been reported in U.S. travelers returning from Kenya or Tanzania.

Reported by: FE Onyango, MD, Dept of Pediatrics, Univ of Nairobi, NN Agata, MD, Kenyatta National Hospital, Nairobi; JS Oliech, MD, Kenya Ministry of Health. PR Hiza, MD, Chief Medical Officer, Ministry of Health, Tanzania. Div of Immunization, Div of Quarantine, Center for Prevention Svcs; Meningitis and Special Pathogens Br, Div of Bacterial Diseases, Center for Infectious Diseases, CDC.

**Editorial Note:** Epidemic meningococcal disease in Kenya and Tanzania is uncommon. These outbreaks may be caused by the same III-1 clonal group of *N. meningitidis* serogroup A responsible for recent major epidemics in Nepal (1983–1984), Saudi Arabia (1987), and Chad (1988) (*1*).

Because of the potential risk for exposure in Kenya and northern Tanzania, CDC recommends that U.S. travelers to these areas receive the meningococcal polysaccharide vaccine. The serogroup A meningococcal vaccine is 85%–95% effective for at least 1 year in persons  $\geq$ 4 years of age (2,3). Vaccine efficacy data for children <4 years of age are limited. However, vaccine immunogenicity increases progressively with age from 3 months to 4 years. Adverse reactions to the meningococcal vaccine are mild and infrequent, consisting primarily of localized erythema lasting 1–2 days (2). Protective immunity is achieved 10–14 days after vaccination. Quadrivalent

# Meningococcal Disease - Continued

polysaccharide vaccine containing serogroups A, C, Y, and W135 is available to physicians from local distributors or Connaught Laboratories at (800) 822-2463.

Although group A meningococcal disease is rare in the United States, carriage of group A meningococcus has been reported in travelers returning from countries with epidemics (4). Physicians are encouraged to report all cases of meningococcal disease to their local and state health departments and to forward meningococcal isolates to CDC through their state public health laboratories.

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# Epidemiologic Notes and Reports

# Update: Eosinophilia-Myalgia Syndrome Associated with Ingestion of L-Tryptophan – United States, as of January 9, 1990

As of January 9, 1046 eosinophilia-myalgia syndrome (EMS) cases have been reported to CDC from 49 states, the District of Columbia, and Puerto Rico (Figure 1) (1-3). Only Alaska has reported no cases (Figure 1). Seven deaths have been reported in patients who met the surveillance case definition and who used L-tryptophan (LT).

As of January 9, CDC has received completed report forms from 38 states and Puerto Rico with information about 429 cases fitting the case definition. Ages of these patients ranged from 11 years to 84 years (median: 48 years); 96% of patients were non-Hispanic white, 2% were Hispanic, and 1% were black. Three hundred sixty (84%) were female. Four hundred nineteen (98%) had histories of LT ingestion preceding onset of symptoms; dosage ranged from 26 mg to 15,000 mg per day (median: 1500 mg per day). Three hundred seventy-two (87%) reported onset of symptoms during or after July 1989 (Figure 2). Of the EMS patients reported thus far, 139 (32%) have required hospitalization.

Physicians report suspected cases to state and local health departments, and information is recorded on a standardized case-report form. Total numbers of EMS cases are telephoned weekly to CDC by health departments, and case-report forms mailed when completed; this results in a timely accumulation of total numbers but a lag in availability of detailed data.

Reported by: State and territorial health departments. Div of Environmental Hazards and Health Effects, Center for Environmental Health and Injury Control, CDC.

# Update: EMS - Continued

### References

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# FIGURE 1. Eosinophilia-myalgia syndrome cases, by area – United States, as of January 9, 1990

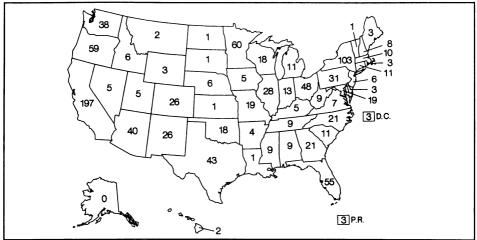
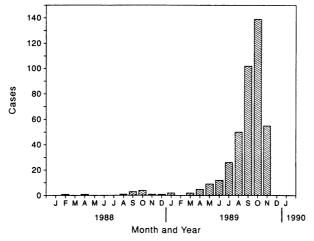


FIGURE 2. Eosinophilia-myalgia syndrome (EMS) cases, by month of onset\* – United States, January 1, 1988–January 9,  $1990^{+}$ 



\*Nine reports did not specify month of onset and are not included here. <sup>†</sup>Six persons had EMS onset before 1988; one case each occurred in August 1982, March 1983, September 1986, and February, July, and November 1987.

# Notices to Readers

# **Epidemiology in Action Course**

CDC and Emory University will cosponsor a course designed for practicing state and local health department professionals. This course, "Epidemiology in Action," will be held at CDC May 14–25, 1990. It emphasizes the practical application of epidemiology to public health problems and will consist of lectures, workshops, classroom exercises (including actual epidemiologic problems), discussions, and an on-site community survey. Applications are due March 1. For further information or an application form, contact Department PSB, Division of Public Health, Emory University, 1599 Clifton Road, N.E., Atlanta, GA 30329; telephone (404) 727-0199; FAX (404) 727-8744; TELEX (810) 751-8512.

# Nonpublication of MMWR on December 29, 1989

Because of the holidays, the *MMWR* was not published on December 29, 1989. Issues 51 and 52 of volume 38 were published together on January 5, 1990.

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The data in this report 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. The editor welcomes accounts of interesting cases, outbreaks, environmental hazards, or other public health problems of current interest to health officials. Such reports and any other matters pertaining to editorial or other textual considerations should be addressed to: Editor, *Morbidity and Mortality Weekly Report*, Centers for Disease Control, Atlanta, Georgia 30333; telephone (404) 332-4555.

Acting Director, Centers for Disease Control Walter R. Dowdle, Ph.D. Director, Epidemiology Program Office Stephen B. Thacker, M.D., M.Sc. Editor, *MMWR* Series Richard A. Goodman, M.D., M.P.H. Managing Editor Karen L. Foster, M.A.

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