



# MORBIDITY AND MORTALITY WEEKLY REPORT

- 21 Dengue Type 3 Infection
  - Race-Specific Differences in Influenza Vaccination Levels Among Medicare Beneficiaries — United States, 1993
- 33 Type B Botulism Associated with Roasted Eggplant in Oil Italy, 1993
- 36 Adult Blood Lead Epidemiology and Surveillance United States
- 37 Multistate Outbreak of Viral Gastroenteritis Associated with Consumption of Oysters — Apalachicola Bay, Florida, December 1994–January 1995
- 39 Notices to Readers

# International Notes

# Dengue Type 3 Infection — Nicaragua and Panama, October–November 1994

The geographic range and incidence of dengue virus activity in the Americas substantially increased from 1980 to 1994. During this period, all dengue activity in the Americas was associated with dengue serotypes 1, 2, and 4 (DEN-1, DEN-2, and DEN-4). On November 25, 1994, the Ministry of Health of Nicaragua announced the isolation of dengue type 3 (DEN-3) from two children hospitalized with minor hemorrhagic manifestations in Managua. Subsequently, DEN-3 virus was isolated from two persons with dengue fever in Panama. These cases represent the first isolation of DEN-3 from autochthonous cases in the Americas since 1977. This report describes these cases and dengue activity in Nicaragua and Panama and summarizes public health activities to control dengue fever in the Americas.

On October 22, 1994, a 5-year-old boy was hospitalized in Managua with symptoms of classic dengue fever (i.e., fever, vomiting, and rash). On October 23, a 10-year-old girl from Managua was hospitalized with similar symptoms. Both children had an uneventful clinical course and recovered. DEN-3 virus was isolated and reisolated (for confirmation) from blood samples from both patients by the Pedro Kourí Institute of Tropical Medicine in Havana. Serologic analyses confirmed that both were primary dengue infections.

DEN-3 virus also was isolated from blood samples of two patients with symptoms of dengue fever in Panama by the Gorgas Memorial Laboratory in Panama City. The first case occurred in the province of Chiriquí (sample collected on October 11) and the second, in the province of Panama (sample collected on November 14). Genetic typing of the initial DEN-3 isolate from Panama at CDC indicated the virus belongs to the Sri Lanka/India genotype, which caused major dengue hemorrhagic fever (DHF) epidemics in Sri Lanka and India during 1989–1992.

During 1994, a countrywide dengue epidemic occurred in Nicaragua (1994 estimated population: 4,300,000). A total of 20,469 dengue cases (4.8 per 1000 population) was reported: the attack rate was highest in the province of León (11.9 cases per 1000); the largest number of cases (7631; attack rate: 6.4 per 1000) was reported in Managua. The National Diagnostic and Reference Center documented probable dengue infection

Dengue — Continued

(i.e., presence of antiflavivirus immunoglobulin M) in patients from 15 (83%) of the 18 health-care regions. Of the 20,469 reported cases, 1247 (6.1%) had hemorrhagic manifestations; 900 (4.4%) patients were hospitalized. Reviews of medical records at two hospitals indicated that 35 (5.2%) of 676 hospitalized patients met the diagnostic criteria for dengue hemorrhagic fever (DHF). Six cases of suspected DHF were fatal.

In October and November, as the result of increased dengue transmission in Managua, the Ministry of Health initiated a multifaceted response to control *Aedes aegypti*, the mosquito vector of dengue. This response included ultralow-volume application of insecticides (deltamethrin or cypermethrin); indoor application of mosquito larvicide (temephos [Abate®\*]); and a national, community-based campaign to eliminate mosquito production sites. These activities were supported by efforts to educate the community. By the end of November, the weekly number of reported cases decreased substantially. In late November, an international team sponsored by the Pan American Health Organization (PAHO) traveled to Nicaragua to reinforce laboratory diagnostic capabilities, obtain epidemiologic information, evaluate the severity of disease produced by DEN-3, and assist national authorities in their efforts to control the outbreak.

An outbreak of dengue, which began in July 1994, also occurred in Panama; DEN-1 was the predominant virus serotype isolated. As of December 28, a total of 716 laboratory-diagnosed cases originating from eight of the 13 provinces (attack rate: 27.4 per 100,000 persons) had been reported.

Reported by: M Palacio, JJ Amador, F Acevedo, J de los Reyes, A Ramírez, A González, G Huelva, Ministry of Health, Managua; R Jiménez, Hospital La Mascota, F Ruiz, Hospital Manolo Morales, Managua; R Cuadra, Hospital Escuela, León, Nicaragua. MG Guzmán, G Kourí, M Soler, M Alvarez, R Rodriguez, Pedro Kourí Institute of Tropical Medicine, E Martínez, Hospital William Soler, Havana. E Quiroz, V Bayard, C Campos, MO Vasquez, Ministry of Health, Panama. Div of Communicable Disease Prevention and Control, Pan American Health Organization, Washington, DC, and Nicaragua. Dengue Br, Div of Vector-Borne Infectious Diseases, National Center for Infectious Diseases, CDC.

Editorial Note: Dengue fever is an acute, mosquito-transmitted viral disease characterized by fever, headache, arthralgia, myalgia, rash, nausea, and vomiting. Infections are caused by any of the four virus serotypes. Although most dengue infections result in relatively mild illness, some can produce DHF. Based on the World Health Organization case definition (1), a case of DHF must meet the following criteria: 1) fever, 2) minor or major hemorrhagic manifestations, 3) thrombocytopenia (≤100,000/mm³), and 4) objective evidence of increased capillary permeability (e.g., hemoconcentration [hematocrit increased by ≥20%], pleural effusions [evidenced by chest radiography or other imaging method], or hypoproteinemia). A case of dengue shock syndrome (DSS) must meet all the criteria for DHF plus hypotension or narrow pulse pressure (≤20 mm Hg); the fatality rate for patients with DSS can be as high as 44% (2).

In 1994, outbreaks of dengue were reported from Brazil, Costa Rica, Dominican Republic, Haiti, Mexico, Puerto Rico, and Venezuela. In Nicaragua, an outbreak in 1992 was localized in León province and associated with DEN-2 and DEN-4 viruses; an outbreak in June and July 1993 was focused in León and adjacent Chinandega province, but no virus was isolated. A limited outbreak of DEN-2 (14 laboratory-confirmed

<sup>\*</sup>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.

Dengue — Continued

cases) occurred in Panama City in 1993 and was the first recorded outbreak of dengue in Panama since 1942.

DEN-3 was first isolated in the Americas (in Puerto Rico) in 1963 (3) and subsequently caused epidemics in Jamaica and the eastern Caribbean during that year (4). Although DEN-3 has been isolated from international travelers returning to the Americas from other geographic regions (5), this serotype was last isolated in the region (in Puerto Rico) in 1977 (6). The identification of autochthonous transmission of DEN-3 in two countries in the Americas (Nicaragua and Panama) in 1994 has important implications for public health because most residents of urban areas in the American tropics are susceptible to this serotype. In particular, all persons in the American tropics who are aged <16 years (approximately one third of the population of Latin America) and all persons living in Ae. aegypti-infested areas who were not infected with DEN-3 during 1963–1978 are at risk for infection with this virus. The appearance of DEN-3 also may increase the risk for DHF resulting from secondary, heterotypic infection (7) or from the introduction of a particularly virulent genotype of DEN-3 (8).

The international response to the outbreak in Nicaragua was specified by a contingency plan developed by health agencies in 1993 to address the potential for reintroduction of DEN-3 into the Americas. This plan highlighted the importance of effective, laboratory-based surveillance programs for dengue and DHF in the Americas. Because of increased DEN-3 activity in other regions and repeated detection of DEN-3 in travelers returning to the Americas, PAHO alerted member countries of these events during April–May 1994 and recommended actions to follow after the identification of an autochthonous case of DEN-3. These actions include prompt investigation to define the magnitude and distribution of the DEN-3 virus in the country and, if the distribution is determined to be limited and circumscribed, efforts to eradicate mosquitoes in the affected area.

PAHO recently published a document for the prevention and control of dengue in the Americas (9), which includes a detailed emergency plan for the control of epidemic dengue and DHF. During 1992–1994, these guidelines were presented by PAHO to national representatives of *Ae. aegypti*-infested countries in the Americas. In addition, during 1994, PAHO teams reviewed national dengue-control programs in selected countries and assisted national authorities in preparing or updating contingency plans for outbreaks.

Health-care providers should consider dengue in the differential diagnosis of all patients who have symptoms compatible with dengue and who reside in or have visited any tropical areas. When dengue is suspected, the patient's blood pressure, hematocrit, and platelet count should be monitored for evidence of hypotension, hemoconcentration, and thrombocytopenia. Acetaminophen products are recommended for management of fever because of the anticoagulant properties of acetylsalicylic acid (i.e., aspirin). Acute- and convalescent-phase serum samples should be obtained for viral isolation and serodiagnosis.

Suspected dengue cases should be reported to the state or territorial health department; the report should include a clinical summary, dates of onset of illness and blood collection, and other epidemiologic information (e.g., a detailed travel history with dates and location of travel). Serum samples should be sent for confirmation through state health department laboratories to CDC's Dengue Branch, Division of Vector-

## Dengue — Continued

Borne Infectious Diseases, National Center for Infectious Diseases, 2 Calle Casia, San Juan, PR 00921-3200; telephone (809) 766-5181; fax (809) 766-6596.

#### References

- 1. World Health Organization. Dengue hemorrhagic fever: diagnosis, treatment, and control. Geneva: World Health Organization, 1986.
- 2. Tassniyom S, Vasanawathana S, Chirawatkul A, Rojanasuphot S. Failure of high-dose methyl-prednisolone in established dengue shock syndrome: a placebo-controlled, double-blind study. Pediatrics 1993;92:111–5.
- 3. Russell PK, Buescher EL, McCown JM, Ordoñez J. Recovery of dengue viruses from patients during epidemics in Puerto Rico and East Pakistan. Am J Trop Med Hyg 1966;15:573–9.
- 4. Ehrenkranz NJ, Ventura AK, Cuadrado RR, Pond WL, Porter JE. Pandemic dengue in Caribbean countries and the southern United States: past, present, and potential problems. N Engl J Med 1971;285:1460–9.
- 5. Rigau-Pérez JG, Gubler DJ, Vorndam AV, Clark GG. Dengue surveillance—United States, 1986–1992. In: CDC surveillance summaries (July). MMWR 1994;43(no. SS-2):7–19.
- 6. Gubler DJ. Dengue and dengue hemorrhagic fever in the Americas. P R Health Sci J 1987;6: 107–11.
- 7. Halstead SB. Pathogenesis of dengue: challenges to molecular biology. Science 1988;239: 476–81.
- 8. Lanciotti RS, Lewis JG, Gubler DJ, Trent DW. Molecular evolution and epidemiology of DEN-3 virus. J Gen Virol 1994;75:65–75.
- 9. Pan American Health Organization. Dengue and dengue hemorrhagic fever in the Americas: guidelines for prevention and control of dengue and dengue hemorrhagic fever in the Americas. Washington, DC: Pan American Health Organization, 1994; scientific publication no. 548.

# Health Objectives for the Nation

# Race-Specific Differences in Influenza Vaccination Levels Among Medicare Beneficiaries — United States, 1993

One national health objective for the year 2000 is to provide annual influenza vaccination to 60% of all noninstitutionalized, high-risk populations in the United States, (objective 20.11) (1). Since May 1, 1993, Medicare has reimbursed providers for the cost of influenza vaccine; reimbursement for the administration of the vaccine also has been provided for beneficiaries with part B coverage, which allows them to receive the vaccine without a copayment and without having to meet the annual deductible amount for part B reimbursement. Approximately 96% of all persons aged ≥65 years in the United States have Medicare part B coverage (Health Care Financing Administration [HCFA], unpublished data, 1994). To characterize patterns of vaccine use by Medicare beneficiaries, HCFA and CDC estimated influenza vaccine use by Medicare beneficiaries during September–December 1993. Because of disparities in vaccine use by race, this analysis focused on race-specific differences between blacks and whites. This report presents the findings of that analysis.

Claims submitted for services provided during September 1–December 31, 1993, and paid by Medicare were used to identify persons who received influenza vaccine. The percentage of beneficiaries who received Medicare-paid vaccinations was calculated using the HCFA 1993 denominator file for beneficiaries aged ≥65 years for the United States and for each state and county, by sex, 10-year age group, and race

(white and black [data for racial groups other than whites and blacks are grouped together in the Medicare claims data system and were not analyzed separately]) (Tables 1 and 2). Medicare claims are not submitted by managed-care plans; therefore, beneficiaries who are members of such plans (approximately 6% of the Medicare population) were excluded from the analysis. Because 1993 was the first year influenza vaccination was reimbursed by Medicare, approximately 10%–20% of Medicare beneficiaries may have been vaccinated in 1993 and not had claims filed with Medicare (CDC, unpublished data, 1994).

During 1993, a total of 9,831,884 (35%) beneficiaries received Medicare-reimbursed influenza vaccinations. However, the vaccination rate for blacks (17%) was less than half that for whites (37%) (Table 1). Among whites, the vaccination rate for women aged ≥85 years (30%) was lower than that for women aged 65–84 years by approximately eight percentage points and lower than that for men aged ≥85 years by

TABLE 1. Number and rate of influenza vaccinations paid for by Medicare part B, by recipient age, sex, and race — United States, 1993

Race*/Sex/ Age group (yrs)	No. non-HMO <sup>†</sup> part B enrollees	No. Medicare-paid influenza vaccinations	Medicare-paid vaccination rate
WHITE			
Men			
65–74	5,483,480	1,883,175	34%
75–84	3,404,559	1,367,802	40%
≥85	844,735	291,706	35%
Total	9,732,774	3,542,683	36%
Women			
65–74	6,882,485	2,556,524	37%
75–84	5,561,678	2,136,587	38%
≥85	2,256,534	684,299	30%
Total	14,700,697	5,377,410	37%
Overall		• •	
65–74	12,365,965	4,439,699	36%
75–84	8,966,237	3,504,389	39%
≥85	3,101,269	976,005	31%
Total	24,433,471	8,920,093	37%
BLACK			
Men			
65–74	456,776	64,127	14%
75–84	251,261	43,612	17%
≥85	68,566	10,990	16%
Total	776,603	118,729	15%
Women			
65–74	655,873	118,960	18%
75–84	467,096	92,067	20%
≥85	187,636	31,590	17%
Total	1,310,605	242,617	19%
Overall			
65–74	1,112,649	183,087	16%
75–84	718,357	135,679	19%
≥85	256,202	42,580	17%
Total	2,087,208	361,346	17%

<sup>\*</sup>Data for racial groups other than whites and blacks are grouped together in the Medicare claims data system and were not analyzed separately.

<sup>&</sup>lt;sup>†</sup>Health maintenance organization.

TABLE 2. Rates of influenza vaccination paid for by Medicare, by state and race\* of recipient — United States, 1993

		Race				Race		
State	White	Black	Total <sup>†</sup>	State	White	Black	Total <sup>†</sup>	
Alabama	39	19	35	Montana	48	35	48	
Alaska	19	18	16	Nebraska	46	22	45	
Arizona§	44	22	42	Nevada	23	15	23	
Arkansas	46	24	44	New Hampshire	35	28	34	
California§	28	15	26	New Jersey§	27	16	26	
Colorado	47	21	46	New Mexico	28	14	27	
Connecticut	35	22	35	New York	34	13	31	
Delaware	32	17	30	North Carolina	41	18	37	
District of				North Dakota	41	28	41	
Columbia	32	14	20	Ohio	36	21	35	
Florida	41	18	40	Oklahoma	39	18	38	
Georgia	38	16	33	Oregon	46	25	45	
Hawaii	34	24	36	Pennsylvania	40	22	38	
ldaho	47	31	46	Rhode Island	41	27	40	
Illinois	31	11	29	South Carolina§	36	18	32	
Indiana	42	19	41	South Dakota	41	32	40	
lowa	49	27	49	Tennessee	46	21	43	
Kansas	46	21	45	Texas	34	15	32	
Kentucky	35	21	34	Utah	34	20	34	
Louisiana	29	14	26	Vermont	34	33	33	
Maine	42	31	42	Virginia	45	24	41	
Maryland	37	17	34	Washington	42	24	42	
Massachusetts	18	9	17	West Virginia	29	19	29	
Michigan	33	18	32	Wisconsin	45	27	45	
Minnesota	43	26	43	Wyoming	29	24	29	
Mississippi	27	15	24					
Missouri	36	16	34	Total	37	17	35	

<sup>\*</sup>Data for racial groups other than whites and blacks are grouped together in the Medicare claims data system and were not analyzed separately.

approximately five percentage points (p<0.01 for both comparisons). Among blacks, vaccination rates varied 1%–3% between different age-sex groups (Table 1).

Coverage rates for Medicare-reimbursed influenza vaccination ranged from 16% (Alaska) to 49% (Iowa) (Table 2). Vaccination levels were  $\geq$ 40% in 20 (40%) of the 50 states and in the District of Columbia. In 33 (66%) states and in the District of Columbia, vaccination rates for blacks were below 60% of the rates for whites. Vaccination rates for blacks were at least 60% of the rates for whites in 17 states\*; in these states, the total black population aged  $\geq$ 65 years with Medicare part B coverage was 65,515 (3% of the national black population that has Medicare part B coverage).

<sup>†</sup>Includes persons in all racial groups and persons of unknown race.

<sup>§</sup> After the mailing of the original estimates from the Medicare claims data to the 63 federal vaccination grant programs, California and South Carolina reported 473,062 and 23,322 influenza vaccinations administered to persons aged ≥65 years, respectively, which were not billed to Medicare in 1993. Arizona estimated that an additional 40,000 persons received vaccine that was not billed to Medicare, and New Jersey estimated 80,000–100,000 doses were not billed. This additional information is not reflected in the table.

<sup>\*</sup>Alaska, Connecticut, Hawaii, Idaho, Kentucky, Maine, Minnesota, Montana, Nevada, New Hampshire, North Dakota, Rhode Island, South Dakota, Vermont, West Virginia, Wisconsin, and Wyoming.

Reported by: AM McBean, MD, Univ of Minnesota School of Public Health, Minneapolis. JD Babish, MPH, Office of Research and Demonstrations, Health Care Financing Administration. Adult Vaccine Preventable Disease Br, National Immunization Program, CDC.

Editorial Note: The findings in this report are consistent with previous surveys that have documented lower influenza vaccination coverage among blacks than whites. For example, based on the 1991 National Health Interview Survey, among all persons aged ≥65 years, 41% had been vaccinated; however, within this age group, blacks were less likely than whites to have been vaccinated (27% and 43%, respectively) (2). Findings from the Medicare Current Beneficiary Survey (September–December 1992) indicated that, for noninstitutionalized beneficiaries, the vaccination rate during winter 1991–92 was 48% overall but 29% among blacks and 50% among whites (Office of the Actuary, HCFA, unpublished data, 1994). These variations may reflect differences in factors such as socioeconomic status, access to medical care, and prevalence of specific risks.

The finding in this report that  $\ge 40\%$  of beneficiaries in 20 states and the District of Columbia had received vaccine indicates that, in these areas, substantial progress has been made toward achieving the national health objective for the year 2000 (1,3). Because the wide variations in state-specific vaccination levels (Table 2) also have been documented for Medicare-reimbursed pneumococcal vaccination claims (4), analysis of these variations may assist in planning programs for increasing vaccine coverage.

Because not all providers submitted claims to Medicare for reimbursement, the rates for Medicare-reimbursed influenza vaccination claims in this report are lower than those based on other national surveys. However, failure to submit claims to Medicare in 1993 for influenza vaccination services is not known to have differentially affected claims submitted for vaccinations administered to black beneficiaries compared with white beneficiaries. In the future, reporting may be enhanced through communication with public and private providers; the use of simplified billing procedures; and helping public-sector providers, visiting nurse groups, and others obtain Medicare provider numbers (5).

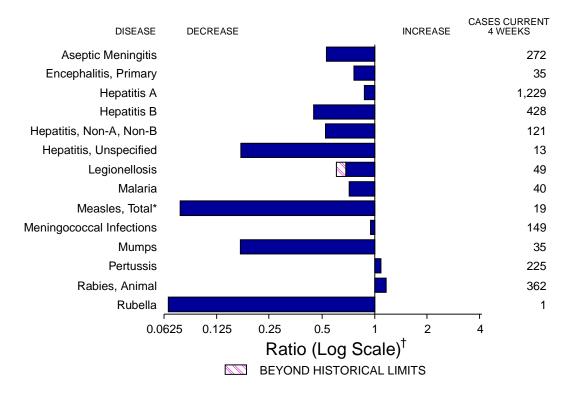
HCFA is collaborating with a coalition representing approximately 160 community organizations to identify strategies to improve coverage in 1995. In addition, as part of HCFA's Consumer Information Strategy (6), demographic- and county-specific vaccination rates for 1993 (7) were provided to health-care providers; consumer-based organizations; local, state, and other federal agencies; and Medicare beneficiaries. This information should assist in increasing beneficiary use of influenza vaccine and addresses consumer and provider concerns about the risks for influenza and the effectiveness and safety of influenza vaccine (8–10). The county-specific vaccination rates also may assist programs receiving federal childhood vaccination grants to improve influenza vaccination coverage.

#### References

- 1. Public Health Service. Healthy people 2000: national health promotion and disease prevention objectives. Washington, DC: US Department of Health and Human Services, Public Health Service, 1991:122–3; DHHS publication no. (PHS)91-50213.
- 2. Heath KA, Strikas RA, Stevenson J, Williams WW. Influenza and pneumococcal vaccination among older adults: results of the 1991 National Health Interview Survey [Abstract]. In: Program and abstracts of the CDC Epidemic Intelligence Service 43rd annual conference. Atlanta: US Department of Health and Human Services, Public Health Service, CDC, 1994:33.

(Continued on page 33)

FIGURE I. Notifiable disease reports, comparison of 4-week totals ending January 14, 1995, with historical data — United States



<sup>\*</sup>The large apparent decrease in the number of reported cases of measles (total) reflects dramatic fluctuations in the historical baseline.

TABLE I. Summary — cases of specified notifiable diseases, United States, cumulative, week ending January 14, 1995 (2nd Week)

	Cum. 1995		Cum. 1995
Anthrax Aseptic Meningitis Brucellosis Cholera Congenital rubella syndrome Diphtheria Encephalitis, primary Encephalitis, post-infectious	79 5 - - - 6 1	Plague Poliomyelitis, Paralytic <sup>†</sup> Psittacosis Rabies, human Rocky Mountain Spotted Fever Syphilis, congenital, age < 1 year <sup>§</sup> Tetanus Toxic shock syndrome	2 4
Haemophilus influenzae* Hansen Disease Hepatitis, unspecified Leptospirosis	34 - 2 2	Trichinosis Tularemia Typhoid fever	- - 10

<sup>&</sup>lt;sup>†</sup>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 the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

<sup>\*</sup>Of 34 cases of known age, 8 (24%) were reported among children less than 5 years of age.

†Updated quarterly from reports to the Division of Sexually Transmitted Diseases and HIV Prevention, National Center for Prevention Services. First quarter data not yet available.

<sup>-:</sup> no reported cases

TABLE II. Cases of selected notifiable diseases, United States, weeks ending January 14, 1995, and January 15, 1994 (2nd Week)

			Hepatitis (Viral), by type								
Reporting Area	AIDS*	Gonor	rhea	,	1				,NB	Legion	ellosis
neporting Area	Cum. 1995	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994
UNITED STATES	-	8,911	14,305	370	677	97	340	32	154	17	57
NEW ENGLAND	-	154	309	5	13	2	13	-	7	-	1
Maine N.H.	-	1 1	2	-	-	-	-	-	-	-	-
Vt.	-	-	-	-	-	-	-	-	-	-	-
Mass. R.I.	-	135 17	127 19	1 2	7 6	1 1	8 2	-	2 5	-	1
Conn.	-	-	161	2	-	-	3	-	-	-	-
MID. ATLANTIC Upstate N.Y.	-	276	1,116	1	57 4	2 2	53 1	6 6	14 2	-	2
N.Y. City	-	-	948	-	31	-	19	-	-	-	-
N.J. Pa.	-	26 250	24 144	1	14 8	-	21 12	-	8 4	-	1 1
E.N. CENTRAL	-	2,840	3,380	100	80	15	72	8	21	7	25
Ohio Ind.	-	1,155 60	1,330 319	78 2	12 16	2 2	7 19	1	- 1	7	6 9
III.	-	662	806	-	36	-	17	_	4	-	4
Mich. Wis.	-	921 42	622 303	20	7 9	11	18 11	7	16	-	5 1
W.N. CENTRAL	-	479	697	6	35	2	20	1	-	2	5
Minn. Iowa	-	110 75	112 34	1 3	1 1	2	2	- 1	-	2	3
Mo.	-	164	386	-	23	-	18	-	-	-	-
N. Dak. S. Dak.	-	1	1 1	-	-	-	-	-	-	-	-
Nebr.	-	120	5 150	2	8 2	-	-	-	-	-	1
Kans. S. ATLANTIC	_	129 3,743	158 3,941	12	∠ 17	- 21	38	- 7	14	3	1 2
Del.	-	81	51	-	-	1	1	-	-	-	-
Md. D.C.	_	567 149	506 362	7	6 3	3 5	10 2	2	2	-	-
Va. W. Va.	-	195	672	- 1	- 1	-	-	- 1	- 1	-	-
N.C.	-	53 540	17 1,165	1	1	2 7	20	3	6	1 2	1
S.C. Ga.	-	380 1,050	500	-	3 1	1	2	-	4	-	1
Fla.	-	728	668	3	2	2	3	1	1	-	-
E.S. CENTRAL	-	300	1,394	6	103	7	62	-	58	1	17
Ky. Tenn.	_	15 -	190 411	5 -	14 6	3 4	11 45	_	3 55	-	2
Ala. Miss.	-	- 285	384 409	- 1	5 78	-	6	-	-	- 1	- 15
W.S. CENTRAL	_	494	1,449	10	4	5	10	1	4	-	1
Ark.	-	-	220	-	-	-	-	-	-	-	-
La. Okla.	-	494 -	711 5	7	4	5	10	1	4	-	1
Tex.	-	-	513	3	-	-	-	-	-	-	-
MOUNTAIN Mont.	-	198 -	307 15	89 2	90	13 1	12	2 1	18 -	-	4 1
ldaho	-	2	1	3	3	1	-	-	5	-	-
Wyo. Colo.	-	3 104	1 160	40	9	5	2	1	2 6	-	1
N. Mex. Ariz.	-	11 77	38 18	39 1	33 44	6	7 1	-	1	-	1
Utah	-	1	8	4	1	-	1	-	2	-	-
Nev.	-	-	66	-	-	-	1	-	2	-	1
PACIFIC Wash.	-	427 -	1,712 139	141 -	278 18	30	60 3	7 -	18 1	4	-
Oreg. Calif.	-	392	44 1,471	17 120	9 243	1 29	1 51	- 7	- 15	2	-
Alaska	-	22	25	1	5	-	-	-	-	-	-
Hawaii	-	13	33	3	3	-	5	-	2	2	-
Guam P.R.	-	- 18	3 23	-	-	-	-	-	-	-	-
V.I. Amer. Samoa	-	-	23 2 1	-	- 1	-	-	-	-	-	-
C.N.M.I.	-	-	4	-	-	-	-	-	-	-	-

N: Not notifiable U: Unavailable -: no reported cases C.N.M.I.: Commonwealth of Northern Mariana Islands \*Updated monthly to the Division of HIV/AIDS, National Center for Infectious Diseases.

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending January 14, 1995, and January 15, 1994 (2nd Week)

							Measle	es (Rube	eola)		Meningococcal Infections		1	
Reporting Area	Ly	me	Mal	aria	Indig	enous	Impo	orted*	To	tal			Mu	mps
	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	1995	Cum. 1995	1995	Cum. 1995	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994
UNITED STATES	32	125	10	16	-	4	-	_	4	1	57	153	11	34
NEW ENGLAND	3	6	3	3	-	2	-	-	2	-	4	9	-	1
Maine	-	-	-	-	-	-	-	-	-	-	-	1	-	-
N.H. Vt.	-	1	-	-	-	-	-	-	-	-	-	1 -	-	1 -
Mass.	3	-	1	Ī	-	-	-	-	-	-	4	3	-	-
R.I. Conn.	-	3	2	3	-	2	-	-	2	-	-	4	-	-
MID. ATLANTIC	20	111	_	4	_	_	_	_	_	1	2	8	_	2
Upstate N.Y.	-	83	-	-	-	-	-	-	-	-	2	1	-	-
N.Y. City	-	8	-	1	-	-	-	-	-	-	-	-	-	-
N.J. Pa.	20	16 4	-	2 1	Ū	-	Ū	-	-	1	-	4 3	-	2
E.N. CENTRAL	3	_	1	3	_	_	_	_	_	_	12	33	6	10
Ohio	3	_	-	-	_	_	_	_	_	_	4	2	2	-
Ind.	-	-	-	-	U	-	U	-	-	-	1	8	-	_
III. Mich.	-	-	1	2 1	-	-	-	-	-	-	6 1	11 5	4	7 3
Wis.	-	-		-	-	-	-	-	-	-	-	7	-	-
W.N. CENTRAL	2	1	_	1	_	_	_	_	_	-	3	7	2	2
Minn.	-	-	-	-	-	-	-	-	-	-	-	-	-	-
lowa Mo.	-	-	-	1	-	-	-	-	-	-	3	7	1 1	2
N. Dak.			-			-	-	-	-	-	-	-	-	-
S. Dak.	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Nebr. Kans.	2	1	-	-	-	-	-	-	-	-	-	-	-	-
		2	1	1	-	-	-	-	-				-	
S. ATLANTIC Del.	3	2	-	-	-	-	-	-	-	-	13	14	-	2
Md.	-	-	-	-	-	-	-	-	-	-	-	2	-	-
D.C. Va.	-	-	-	-	-	-	-	-	-	-	1	1	-	-
W. Va.	-	_	_	-		-	-	-	-	-	-	_	_	-
N.C.	2	-	1	1	-	-	-	-	-	-	3	3	-	1
S.C. Ga.	1	-	-	-	-	-	-	-	-	-	1	4	-	1
Fla.			-			-	-	-	-	-	8	4	-	-
E.S. CENTRAL	_	4	_	1	_	_	_	_	_	_	1	51	_	8
Ky.	-	4	-	-	-	-	-	-	-	-	1	3	-	-
Tenn.	-	-	-	-	- U	-	- U	-	-	-	-	2 9	-	-
Ala. Miss.	-	_	_	1	-	-	-	-	-	-	-	37	_	8
W.S. CENTRAL	_	_	_	-	_	_	_	_	_	_	3	1	_	_
Ark.	-	-	-	-	-	-	-	-	-	-	-		-	-
La.	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Okla. Tex.	-	-	-	-	-	-	-	-	-	-	1 2	1	-	-
MOUNTAIN		_	1	_	_	2	_	_	2	_	10	10	_	_
	_	_	-	-	_	-	_	_	-	_	-	1	-	_
Mont. Idaho	-	-	-	-	-	-	-	-	-	-	1	1	-	-
Wyo. Colo.	-	-	1	-	-	-	-	-	-	-	-	1	-	-
N. Mex.	_	_	-	-	_	2	_	_	2	_	4	i	N	N
Ariz.	-	-	-	-	-	-	-	-	-	-	5	4	-	-
Utah Nev.	-	-	-	-	Ū	-	Ū	-	-	-	-	2	-	-
PACIFIC	1	1	4	3	J			-			9	20	3	9
Wash.	-	-	-	-	-	-	-	-	-	-	2	1	-	1
Oreg.	-	-	-	-	-	-	-	-	-	-	1	1	N	N
Calif. Alaska	1	1	3 1	2	-	-	-	-	-	-	5 -	18	3	6 2
Hawaii	-	-	-	1	-	-	-	-	-	-	1	-	-	-
Guam	_	_	_	-	U	_	U	_	_	_	_	_	_	_
P.R.	-	-	-	-	-	-	-	-	-	-	-	-	-	-
V.I. Amer. Samoa	-	-	-	-	U	-	U	-	-	-	-	-	-	-
Amer Samoa	-	-	-	1	U U	-	U U	-	-	9	-	-	-	-

 $<sup>{\</sup>rm *For}\ imported\ measles,\ cases\ include\ only\ those\ resulting\ from\ importation\ from\ other\ countries.$ 

N: Not notifiable

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending January 14, 1995, and January 15, 1994 (2nd Week)

Reporting Area		Pertussis			Rubella		Sypt (Prima Secon	ary &	Tuberc	ulosis	Rab Anii	
. •	1995	Cum. 1995	Cum. 1994	1995	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994
UNITED STATES	58	66	163	-	-	1	305	730	283	754	127	160
NEW ENGLAND	5	5	5	-	-	-	5	9	2	4	43	47
Maine N.H.	4	4	3	-	-	-	-	-	-	-	3	3
Vt.	-	-	2	-		-	-	-	-	-	3	2
Mass.	1	1	-	-	-	-	3	3	-	-	25	24
R.I. Conn.	-	-	-	-	-	-	2	6	2	4	12	1 17
MID. ATLANTIC	-	-	37		-	1	1	46	22	21	37	53
Upstate N.Y.	-	_	-	-	-	1	-	40	-	1	23	33
N.Y. City	-	-	-	-	-	-	-	42	22	19	-	-
N.J. Pa.	Ū	-	2 35	Ū	-	-	1	1 3	-	1	12 2	11 9
	7				-	-						
E.N. CENTRAL Ohio	, 5	9 5	45 15	-	-	-	52 17	71 9	34 9	41 8	1 1	1 -
Ind.	ŭ	-	-	U	-	-	3	11	1	1	-	-
III. Mich.	2	4	16 3	-	-	-	27 5	30 9	23	32	-	-
Wis.	-	-	11	-		-	5 -	12	1	-	-	1
W.N. CENTRAL	3	4	6	_	_	_	16	52	7	1	3	5
Minn.	-	-	-	-	-	-	3	3	-		-	-
lowa	-	-	-	-	-	-	3	2	2	-	2	2
Mo. N. Dak.	-	1 -	1	-	-	-	10	47 -	1 -	-	-	-
S. Dak.	-	-	-	-	-	-	-	-	-	-	-	-
Nebr. Kans.	3	3	- 5	-	-	-	-	-	4	- 1	1	3
S. ATLANTIC	21	21	22	_	_	_	121	167	39	30	30	33
Del.			-	-	-	-	-	-	-	1	2	-
Md. D.C.	-	-	5	-	-	-	13	3 7	25	9	9	7
Va.	-	-	-	-	-	-	4 7	15	1 -	4	6	7
W. Va.			1	-	-	-	-	-	3	-	1	2
N.C. S.C.	21	21	12 4	-	-	-	41 21	75 20	3 6	16	7 2	1 2
Ga.	-	-	-		-	_	20	30	1	-	3	14
Fla.	-	-	-	-	-	-	15	17	-	-	-	-
E.S. CENTRAL	-	-	10	-	-	-	72	166	4	279	8	3
Ky. Tenn.	-	-	1	-	-	-	-	9 45	1 -	4	8	-
Ala.	Ū		1	Ū	-	-	-	32		16	-	3
Miss.	-	-	8	-	-	-	72	80	3	259	-	-
W.S. CENTRAL	-	-	8	-	-	-	35	149	1	-	1	2
Ark. La.	-	-	-	-	-	-	35	18 74	-	-	1	1
Okla.	-		8		-	-	-	-	1	-	-	1
Tex.	-	-	-	-	-	-	-	57	-	-	-	-
MOUNTAIN	22	22	2	-	-	-	3	13	5	19	4	2
Mont. Idaho	- 7	- 7	-	-	-	-	-	-	-	-	3	-
Wyo.	-	-	-	-	-	-	-	-	-	-	-	-
Colo.	-	-	1	-	-	-	2	7	-	-	-	-
N. Mex. Ariz.	- 15	15	1	-	-	-	- 1	2	- 5	- 17	1	2
Utah	-	-	-	-	-	-	-	1	-	-	-	-
Nev.	U	-	-	U	-	-	-	3	-	2	-	-
PACIFIC	-	5	28	-	-	-	-	57	169	359	-	14
Wash. Oreg.	-	-	3	-	-	-	-	1	4	4 4	-	-
Calif.	-	5	21	-	-	-	-	56	158	345	_	10
Alaska	-	-	-	-	-	-	-	-	-	3	-	4
Hawaii	-	-	1	-	-	-	-	-	7	3	-	-
Guam P.R.	U	-	-	U	-	-	3	13	-	-	1	-
V.I.	U	-	-	U	-	-	- -	13	-	-	-	-
Amer. Samoa	U	-	-	U	-	-	-	-	-	-	-	-
C.N.M.I.	Ü	-	-	U	-	-	-	-	-	6	-	

U: Unavailable -: no reported cases

TABLE III. Deaths in 121 U.S. cities,\* week ending January 14, 1995 (2nd Week)

	A	All Cau	ses, By	/ Age (Y	ears)		P&l <sup>†</sup>		,	All Cau	ıses, By	/ Age (Y	ears)		P&I <sup>†</sup>
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	580	402		47 16	11	14	45	S. ATLANTIC	1,636	1,051	325	177	52	29	117
Boston, Mass. Bridgeport, Conn.	182 61	117 42	35 9	16 8	5 1	9 1	8	Atlanta, Ga. Baltimore, Md.	238 454	134 291	51 90	42 53	7 15	4 5	8 53
Cambridge, Mass.	18	12		1	-	-	-	Charlotte, N.C.	69	48	10	9 9	1	1	6
Fall River, Mass. Hartford, Conn.	17 U	10 U	3 U	4 U	Ū	Ū	1 U	Jacksonville, Fla. Miami, Fla.	174 73	118 36	41 18	16	2 2	3 1	11 -
Lowell, Mass.	19	14		1	-	-	2	Norfolk, Va.	61	37	12	7	3	2 U	6
Lynn, Mass. New Bedford, Mass	12 s. 23	8 15	1 4	2 4	1	-	2	Richmond, Va. Savannah, Ga.	U 50	U 37	U 7	U 3	U 1	2	U 4
New Haven, Conn.	46	32	12	1	1	-	4	St. Petersburg, Fla.	94	70	15	5	2	2	5
Providence, R.I. Somerville, Mass.	49 3	40 3		3	1	-	11 -	Tampa, Fla. Washington, D.C.	240 176	176 101	42 38	12 20	10 8	9	17 7
Springfield, Mass.	53	36	14	1	1	1	6	Wilmington, Del.	7	3	1	1	1	-	-
Waterbury, Conn.	40 57	32 41		3 3	- 1	3	6 5	E.S. CENTRAL	885	579	190	73	21	22	62
Worcester, Mass.			-		-			Birmingham, Ala.	164	100	36	15	6	7	4
MID. ATLANTIC Albany, N.Y.	3,267 33	2,193 22	574 3	366 5	82 2	52 1	184 4	Chattanooga, Tenn. Knoxville, Tenn.	. 86 87	58 53	19 21	8 8	1 2	3	6 10
Allentown, Pa.	36	29	4	3	-	-	-	Lexington, Ky.	75	48	18	5	2	2	7
Buffalo, N.Y. Camden, N.J.	110 40	94 29	15 6	3	1	1 1	11 4	Memphis, Tenn. Mobile, Ala.	142 129	95 94	28 25	13 7	3 1	3 2	18 9
Elizabeth, N.J.	44	31	8	4	-	i	2	Montgomery, Ala.	36	23	8	5	-	-	2
Erie, Pa.§ Jersey City, N.J.	51 44	38 31	9 7	4 4	2	-	3 2	Nashville, Tenn.	166	108	35	12	6	5	6
New York City, N.Y.		1,166	350	234	41	28	82	W.S. CENTRAL	1,690	1,095	312	185	48	50	122
Newark, N.J.	80 39	39 25	23 6	13 5	4 3	1	7 3	Austin, Tex. Baton Rouge, La.	92 88	56 61	21 13	14 10	1 1	3	4 1
Paterson, N.J. Philadelphia, Pa.	510	346		58	16	8	33	Corpus Christi, Tex.	82	50	18	10	2	2	4
Pittsburgh, Pa.§	77	52		10	5	2	4	Dallas, Tex. El Paso, Tex.	263 101	167 76	58 14	26 9	6 2	6	11 13
Reading, Pa. Rochester, N.Y.	18 153	14 110	2 24	2 10	4	5	4 8	Ft. Worth, Tex.	130	84	25	15	5	1	10
Schenectady, N.Y.	29	26	3	-	-	-	1	Houston, Tex. Little Rock, Ark.	341 96	206 58	61 20	54 10	10 4	10 4	42 10
Scranton, Pa.§ Syracuse, N.Y.	30 102	27 77	1 16	2 5	3	1	1 10	New Orleans, La.	70	38	8	9	6	9	-
Trenton, N.J.	39	28	4	3	1	3	5	San Antonio, Tex.	269 35	183 26	48 5	17 3	8 1	13	17 3
Utica, N.Y. Yonkers, N.Y.	13 U	9 U	3 U	1 U	Ū	Ū	Ū	Shreveport, La. Tulsa, Okla.	123	90	21	8	2	2	3 7
E.N. CENTRAL	2,432	1,573	452	238	99	68	121	MOUNTAIN	986	662	187	87	26	23 3	76
Akron, Ohio	69	58	7	4	-	-	-	Albuquerque, N.M.		92	24	9	5		4
Canton, Ohio	45 438	33 185	8 92	3 84	63	1 14	6 11	Colo. Springs, Colo Denver, Colo.	. 46 152	27 99	14 25	4 21	1	1 6	6 14
Chicago, III. Cincinnati, Ohio	106	79	15	10	-	2	12	Las Vegas, Nev.	177	126	38	11	2	_	16
Cleveland, Ohio	184	114	40	24	3	3	2	Ogden, Utah Phoenix, Ariz.	18 225	13 130	51	1 29	2 12	2 2 U	2 23
Columbus, Ohio Dayton, Ohio	249 129	163 105	57 16	14 7	5	10 1	21 14	Pueblo, Colo.	U	U	U	U	U	ū	U
Detroit, Mich.	298	182		34	8	9	7	Salt Lake City, Utah Tucson, Ariz.	149	61 114	11 24	7 5	4	3 6	7 4
Evansville, Ind. Fort Wayne, Ind.	37 3	32 3	4	1	-	-	2	·					10		-
Gary, Ind.	33	17	9	3	3	1	-	PACIFIC Berkeley, Calif.	1,290 25	869 15	236 6	107 1	19	36 2	122 1
Grand Rapids, Micl Indianapolis, Ind.	h. 76 232	54 158		3 23	2 5	4 7	7 8	Fresno, Calif.	144	85	32	13	5	9	11
Madison, Wis.	60	44	13	3	-	-	2	Glendale, Calif. Honolulu, Hawaii	U 87	U 49	U 27	U 9	U 1	U 1	U 9
Milwaukee, Wis. Peoria, III.	170 27	125 12		9 4	2 1	5 6	12	Long Beach, Calif.	83	56	17	5	1	4	10
Rockford, III.	47	40		-	i	-	8	Los Angeles, Calif. Pasadena, Calif.	U 37	U 28	U 4	U 4	U	U 1	U 6
South Bend, Ind.	51	37	9	2	1	2	2	Portland, Oreg.	227	168	31	18	4	6	15
Toledo, Ohio Youngstown, Ohio	111 67	79 53	18 10	7 3	4 1	3	5 2	Sacramento, Čalif.	U	U	U	U	U	U	U U
W.N. CENTRAL	860	630	131	48	21	9	50	San Diego, Calif. San Francisco, Calif		99	U 26	U 18	U 2	3	13
Des Moines, Iowa	82	62	15	-	3	2	6	San Jose, Calif.	194	139	38	10	3	4	26
Duluth, Minn.	40	33	6 5	1	- 1	-	3	Santa Cruz, Calif. Seattle, Wash.	28 161	24 111	4 26	20	2	2	7 8
Kansas City, Kans. Kansas City, Mo.	26 100	18 57	5 11	2 6	3	2	1 4	Spokane, Wash.	52	37	12	1	-	2	6
Lincoln, Nébr.	35	29	5	1	-	-	1	Tacoma, Wash.	82	58	13	8	1	2	10
Minneapolis, Minn. Omaha, Nebr.	. 220 104	166 72		12 7	7 1	1 2	14 4	TOTAL	13,626 <sup>¶</sup>	9,054	2,513	1,328	379	303	899
St. Louis, Mo.	112	80	18	9	4	1	7								
St. Paul, Minn. Wichita, Kans.	86 55	73 40		4 6	1 1	1	8 2								
vvicinta, Nano.	55	40	,	U		'	2								

<sup>\*</sup>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.

<sup>-:</sup> no reported cases.

- 3. Williams WW, Hickson MA, Kane MA, Kendal AP, Spika JS, Hinman AR. Immunization policies and vaccine coverage among adults: the risk for missed opportunities. Ann Intern Med 1988;108:616–25.
- 4. McBean AM, Babish JD, Prihoda R. The utilization of pneumococcal polysaccharide vaccine among elderly Medicare beneficiaries, 1985 through 1988. Arch Intern Med 1991;151:2009–16.
- 5. CDC. Implementation of the Medicare influenza vaccination benefit—United States, 1993. MMWR 1994;43:771–3.
- 6. Vladeck BC. From the Health Care Financing Administration: the consumer information strategy. JAMA 1994;272:196.
- 7. Health Care Financing Administration. 1993 Influenza immunizations paid for by Medicare: state and county rates. Baltimore: US Department of Health and Human Services, Health Care Financing Administration, 1994.
- 8. CDC. Adult immunization: knowledge, attitudes, and practices—DeKalb and Fulton counties, Georgia, 1988. MMWR 1988;37:657–61.
- 9. Fiebach NH, Viscoli CM. Patient acceptance of influenza vaccination. Am J Med 1991;91:393–400
- 10. Williams WW. Hawaii pneumococcal disease initiative: surveys of consumer and physician knowledge. In: 26th National Immunization Conference proceedings. Atlanta: US Department of Health and Human Services, Public Health Service, CDC, National Center for Prevention Services, Division of Immunization, 1992;117–22.

# International Notes

# Type B Botulism Associated with Roasted Eggplant in Oil — Italy, 1993

In August and October 1993, public health officials in Italy were notified of seven cases of type B botulism from two apparently unrelated outbreaks in different communites. Investigations were initiated by the Regional Health Observatory of Campania and the Italian National Institute of Health. This report summarizes the outbreak investigations, which indicated that illness was associated with eating commercially prepared roasted eggplant in oil.

# **Outbreak 1**

On August 14, two waitresses working in a sandwich bar in Santa Maria di Castellabate were admitted to a local hospital with dysphagia, diplopia, and constipation; a clinical diagnosis of botulism was made. On August 12, the waitresses had prepared and eaten ham, cheese, and eggplant sandwiches. A third waitress also ate the sandwiches and developed dyspepsia for which vomiting was induced; she did not have neurologic symptoms. The owner of the bar, who had tasted a small piece of eggplant from the same jar later on August 12, remained asymptomatic. The cook had initially opened the jar of commercially prepared sliced roasted eggplant in oil and had tasted its contents on August 11 and developed diarrhea. Both the cook and the owner reported that the eggplant tasted spoiled.

Botulism was presumptively diagnosed in the two hospitalized patients; both were treated with trivalent botulism antitoxin and gradually improved. No food samples were available for testing. No botulism toxin was detected in the serum of the two hospitalized patients. However, cultures of their stools subsequently yielded type B *Clostridium botulinum*.

Type B Botulism — Continued

#### **Outbreak 2**

During October 5–6, four of nine members of an extended family who had dined together on October 2 were hospitalized in Naples with suspected botulism. The meal consisted of green olives, prosciutto, bean salad, green salad, mozzarella cheese, sausages, and commercially prepared roasted eggplant in oil. Based on an investigation and analysis of food histories, the eggplant was implicated as the probable source (relative risk=undefined; p<0.01). All of the patients were treated with trivalent botulism antitoxin and gradually improved. Investigation indicated that on September 27, another family member had opened and dipped a fork into the implicated jar of eggplant; although he did not eat any eggplant, he used the fork for other food items. On September 28, he had developed vomiting, dysphagia, and double vision but was not hospitalized; his symptoms resolved spontaneously. On October 8, he was asymptomatic but was hospitalized and treated with trivalent botulism antitoxin after botulism was diagnosed in other family members.

One of the hospitalized patients developed respiratory muscle weakness and required mechanical ventilation. A serum specimen from one patient was negative for botulism toxin. Cultures of stool specimens from three patients yielded proteolytic type B *C. botulinum*. No eggplant was available for testing.

### Follow-Up

The commercially prepared eggplant suspected of causing both outbreaks was produced by one company and sold only in Italy. The company reported preparing the eggplant in the following manner: eggplant slices were washed and soaked overnight in a solution of water, vinegar, and salt; roasted in an oven; and subsequently placed in glass jars. Garlic, peppers, oregano, and citric acid were added. The mixtures then were covered with sunflower oil and sealed with screw-on lids; after being filled, the jars were boiled in water for 30 minutes. The pH of the product was not consistently monitored. A total of 119 jars of eggplant from the same lot that caused the outbreaks was tested; neither *C. botulinum* spores nor botulism toxin were detected. The pH of the product varied from 3.9 to 5.1; the pH was >4.6 in 24 (20%) jars tested.

Public health officials issued a national warning and recalled unused jars of eggplant. No additional cases of botulism associated with this product were reported.

Reported by: P D'Argenio, MD, F Palumbo, MD, R Ortolani, MD, R Pizzuti, MD, M Russo, DBiol, Regional Health Observatory of Campania, R Carducci, MD, Cardarelli Hospital, M Soscia, MD, Contugno Hospital, Naples; P Aureli, DBiol, L Fenicia, DBiol, G Franciosa, DBiol, National Institute of Health, Rome; A Parella, MD, Public Health Dept, Eboli; V Scala, MD, Public Health Dept, Vallo della Lucania, Italy. Foodborne and Diarrheal Diseases Br, Div of Bacterial and Mycotic Diseases, National Center for Infectious Diseases; Global EIS Program, International Br, Div of Field Epidemiology, Epidemiology Program Office, CDC.

**Editorial Note:** Foodborne botulism is a paralytic illness caused by the ingestion of botulism toxin, a neurotoxin produced by the ubiquitous spore-forming bacterium *C. botulinum*. Although botulism toxin in food can be destroyed by heating to boiling (212 F [100 C]) for 10 minutes, the spores are heat resistant and can survive prolonged boiling. To destroy *C. botulinum* spores, food must be heated under pressure to temperatures substantially >212 F. Certain environmental conditions, such as absence of oxygen (anaerobic conditions), pH >4.6, warm temperatures (generally >39 F [>4 C]), high moisture content (water activity), and lack of competing bacterial flora promote production of botulism toxin in foods contaminated with *C. botulinum* spores. The

Type B Botulism — Continued

process used to produce the eggplant epidemiologically implicated in the outbreaks in Italy probably failed to remove *C. botulinum* spores and may have provided such conditions.

Covering foods in oil may provide the anaerobic conditions required for the production of botulism toxin. Outbreaks of botulism in the United States and Canada have been caused by covering vegetables with oil or grease. For example, in 1983, onions covered in grease and left overnight on a grill caused a large outbreak of botulism in Illinois (1), and commercially processed garlic in oil caused outbreaks in 1985 and 1989 (2,3). As a result of these outbreaks, the Food and Drug Administration recommended the addition of antimicrobial growth inhibitors or acidifying agents to such products (4).

In Italy, approximately 50 cases of botulism are reported annually (National Institute of Statistics, unpublished data, 1994), compared with approximately 20 cases annually in the United States (CDC, unpublished data, 1994). However, because the clinical and laboratory diagnoses of botulism can be difficult, these counts of incident cases probably underestimate the actual occurrence. In the United States, surveillance for botulism is linked to the release of botulism antitoxin for treatment of suspected cases. CDC maintains supplies of botulism antitoxin at quarantine stations nationwide for rapid release at the request of state health officials who report suspected cases. This centralized system for controlling antitoxin supplies results in reporting of botulism cases and a well-maintained, reliable source of antitoxin; no such system exists in Europe.

In Italy, as in the United States, outbreaks of botulism associated with commercial products are uncommon; most result from eating improperly preserved home-canned foods (National Institute of Statistics, unpublished data, 1994). However, two previous outbreaks in Italy have been linked to commercial products-mushrooms in oil and pickled olives (5). The outbreaks described in this report probably resulted from a commercial process that was inadequate to prevent contamination of the final product with C. botulinum spores. Another potential explanation is that the jars of eggplant may have been contaminated after they were opened; however, this is less likely because both outbreaks were caused by the same commercial product. Once contaminated with spores, the pH, oil covering, and lack of refrigeration probably provided conditions conducive to the production of botulism toxin. Strict control of the commercial processes used to manufacture such products and the addition of antimicrobial growth inhibitors or acidifying agents could assist in preventing such outbreaks. Persons who prepare roasted vegetables in oil at home should be aware that this practice may be hazardous, especially if such foods are allowed to remain above refrigerator temperature (generally >39 F [>4 C]).

# References

- 1. MacDonald KL, Spengler RF, Hatheway CL, Hargrett NT, Cohen ML. Type A botulism from sauteed onions: clinical and epidemiologic observations. JAMA 1985;253:1275–8.
- 2. St. Louis ME, Shaun HS, Peck MB. Botulism from chopped garlic: delayed recognition of a major outbreak. Ann Intern Med 1988;108:363–8.
- 3. Morse DL, Pickard LK, Guzewich JJ, Devine BD, Shayegani M. Garlic-in-oil associated botulism: episode leads to product modification. Am J Public Health 1990;80:1372–3.
- 4. Food and Drug Administration. Press release no. P89-20. Washington, DC: US Department of Health and Human Services, Public Health Service, Food and Drug Administration, April 17, 1989.

Type B Botulism — Continued

5. Fenicia L, Ferrini AM, Aureli P, Padovan MT. Epidemic of botulism caused by black olives [Italian]. Industrie Alimentari 1992;31:307–8.

# Current Trends

# Adult Blood Lead Epidemiology and Surveillance — United States, Third Quarter 1994

CDC's National Institute for Occupational Safety and Health (NIOSH) Adult Blood Lead Epidemiology and Surveillance (ABLES) program monitors elevated blood lead levels (BLLs) in adults in the United States. Blood lead data from laboratory reports are transmitted to state-based lead surveillance programs and are compiled by NIOSH for quarterly reporting (1).

The total number of elevated blood lead reports during the first three quarters of 1994 increased 18% over third quarter totals from 1993 (Table 1). This increase is consistent with the yearly increase in total reports from 1992 to 1993 ( $\it 2$ ). However, the number of 1994 third quarter reports of higher BLLs (50–59  $\mu g/dL$  and  $\geq$ 60  $\mu g/dL$ ) decreased from 1993 to 1994.

Reports of elevated BLLs represent initial reports of newly identified cases or reflect periodic monitoring of known cases or ongoing exposures. The decrease in reports of higher BLLs during 1993–1994 may represent an increase in detection and prevention efforts by industry. Opportunities for prevention include identification of workers with elevated BLLs, reduction of worker exposures, implementation of workplace controls, and/or timely medical removal. The decrease also may have resulted from underreporting of elevated BLLs.

BLL surveillance allows states to respond rapidly to reports of individual elevated BLLs (or workplace-specific clusters) and to conduct appropriate follow-up and prevention activities. Standardization of BLL reporting by states is progressing and helps to identify the magnitude of the problem and trends among U.S. workers. A year

TABLE 1. Reports of elevated blood lead levels (BLLs) among adults — 22 states,\* third quarter 1994

Reported BLL	Third qu	arter 1994	Cumulative reports,	Cumulative reports,			
(μ <b>g/dL</b> )	No. reports	No. persons <sup>†</sup>	third quarter 1994	third quarter 1993§			
25–39	4,637	2,569	13,267	11,261			
40-49	1,316	641	4,058	3,163			
50-59	233	135	769	788			
≥60	86	51	312	411			
Total	6,272	3,396	18,406	15,623			

<sup>\*</sup>Reported by Alabama, Arizona, California, Connecticut, Illinois, Iowa, Maryland, Massachusetts, Michigan, New Hampshire, New Jersey, New York, North Carolina, Oklahoma, Oregon, Pennsylvania, South Carolina, Texas, Utah, Vermont, Washington, and Wisconsin.

<sup>&</sup>lt;sup>†</sup>Individual reports are categorized according to the highest reported BLL for a person during the given quarter. Pennsylvania reports only numbers of reports on a quarterly basis; quarterly summaries of numbers of persons do not include Pennsylvania data.

<sup>&</sup>lt;sup>§</sup>Data for first quarter 1993 reported from 17 states (Alabama, Connecticut, Illinois, Iowa, Maryland, Massachusetts, Michigan, New Hampshire, New Jersey, New York, Oregon, Pennsylvania, South Carolina, Texas, Utah, Vermont, and Wisconsin). Data for second and third quarters 1993 also include reports from Arizona, California, and Washington.

#### Blood Lead Epidemiology — Continued

2000 national health objective is to eliminate exposures that result in workers having blood lead concentrations >25  $\mu$ g/dL (baseline: 4804 workers with BLLs >25  $\mu$ g/dL in seven states in 1988) (objective 10.8) (3). Substantial progress is needed to meet this objective.

Reported by: JP Lofgren, MD, Alabama Dept of Public Health. C Fowler, MS, Arizona Dept of Health Svcs. FJ Mycroft, PhD, Occupational Health Br, California State Dept of Health Svcs. BC Jung, MPH, Connecticut Dept of Public Health and Addiction Svcs. M Lehnherr, Occupational Disease Registry, Div of Epidemiologic Studies, Illinois Dept of Public Health. R Gergely, Iowa Dept of Public Health. E Keyvan-Larijani, MD, Lead Poisoning Prevention Program, Maryland Dept of the Environment. R Rabin, MSPH, Div of Occupational Hygiene, Massachusetts Dept of Labor and Industries. A Carr, MBA, Bur of Child and Family Svcs, Michigan Dept of Public Health. D Solet, PhD, Div of Public Health Svcs, New Hampshire State Dept of Health and Human Svcs. B Gerwel, MD, Occupational Disease Prevention Project, New Jersey State Dept of Health. R Stone, PhD, New York State Dept of Health. S Randolph, MSN, North Carolina Dept of Environment, Health, and Natural Resources. E Rhoades, MD, Oklahoma State Dept of Health. M Barnett, MS, State Health Div, Oregon Dept of Human Resources. J Gostin, MS, Occupational Health Program, Div of Environmental Health, Pennsylvania Dept of Health. R Marino, MD, Div of Health Hazard Evaluations, South Carolina Dept of Health and Environmental Control. D Perrotta, PhD, Bur of Epidemiology, Texas Dept of Health. D Beaudoin, MD, Bur of Epidemiology, Utah Dept of Health. L Toof, Div of Epidemiology and Health Promotion, Vermont Dept of Health. J Kaufman, MD, Washington State Dept of Labor and Industries. H Anderson, MD, D Higgins, Wisconsin Dept of Health and Social Svcs. Div of Surveillance, Hazard Evaluations, and Field Studies, National Institute for Occupational Safety and Health, CDC.

#### References

- 1. CDC. Surveillance of elevated blood lead levels among adults—United States, 1992. MMWR 1992;41:285–8.
- 2. CDC. Adult blood lead epidemiology and surveillance—United States, 1992–1994. MMWR 1994;43:483–5.
- 3. Public Health Service. Healthy people 2000: national health promotion and disease prevention objectives—full report, with commentary. Washington, DC: US Department of Health and Human Services, Public Health Service, 1991; DHHS publication no. (PHS)91-50212.

# Epidemiologic Notes and Reports

# Multistate Outbreak of Viral Gastroenteritis Associated with Consumption of Oysters — Apalachicola Bay, Florida, December 1994–January 1995

On January 3, 1995, the Florida Department of Health and Rehabilitative Services (HRS) was notified of an outbreak of acute gastroenteritis associated with eating oysters. The subsequent investigation by HRS has identified 34 separate clusters of cases, many of which were associated with oysters harvested during December 29–31 from 13 Mile Area and Cat Point in Apalachicola Bay. Oysters were shipped to other states, but additional clusters of illness associated with these oysters have been reported only in Georgia. Most of these oysters were served steamed or roasted. This report summarizes the preliminary findings of the ongoing investigation of this outbreak.

On January 4, Apalachicola Bay was closed to harvesting even though levels of fecal coliforms in the water and in the oyster meat were within acceptable limits. The preliminary investigation identified no gross breaches of sanitation; however, during

Viral Gastroenteritis — Continued

the holiday season, the bay was used heavily by recreational boaters and commercial fishermen. Clusters of cases identified since the bay was closed prompted concern regarding the continued marketing of these oysters as unshelled and as shucked product both in Florida and other states.

Following the detection of cases associated with oysters from Apalachicola Bay, enhanced surveillance detected three additional clusters of cases in Florida and two in Texas initially linked to oysters harvested in Galveston Bay. As a result, on January 13, Galveston Bay was closed to harvesting.

Norwalk-like viruses have been detected by electronmicroscopy in stool specimens from seven of 11 persons who ate oysters from Apalachicola Bay.

Reported by: C Aristeguieta, MD, Dept of Family Medicine, Univ of Miami; I Koenders, Districts 1 and 2 Health Office, Tallahassee; D Windham, Districts 3 and 13 Health Office, Ocala; K Ward, MSEH, Districts 4 and 12 Health Office, Daytona Beach; E Gregos, Districts 5 and 6 Health Office, Tampa; L Gorospe, E Ngo-Seidel, MD, Nassau County Public Health Unit, Fernandina Beach; J Walker, MD, District 4 Health Office, Jacksonville; WG Hlady, MD, R Hammond, PhD, RS Hopkins, MD, State Epidemiologist, Florida Dept of Health and Rehabilitative Svcs. DM Simpson, MD, State Epidemiologist, Texas Dept of Health. Viral Gastroenteritis Section, Respiratory and Enteric Viruses Br, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases; Div of Field Epidemiology, Epidemiology Program Office, CDC.

**Editorial Note:** Outbreaks of oyster-associated gastroenteritis affect substantially more persons than those identified in the few documented sentinel clusters (1–3). An important feature of these outbreaks is the inherent delays in removing contaminated oysters from the market. Although oyster tags permit traceback to the general harvest areas, they are not sufficiently detailed to allow recall of oysters from a specific site, and they can be lost when oysters are shucked. In this outbreak, the continued occurrence of cases 1 week after the bay was closed and the product was recalled suggests that the contaminated product was still available to consumers. Cooking (i.e., steaming and roasting) did not always render the oysters noninfectious. In addition, enhanced surveillance in Florida prompted by the investigation led to the closing of an oyster bed in Texas. The observation that both the quality of water in the Florida beds and the meat in the implicated oysters met national standards underscores the inherent limitations of the existing methods and the urgent need for improved indicators of viral contamination. In the absence of such indicators, it is difficult to determine when a bed can be safely reopened.

The findings in this investigation indicate the outbreak resulted from consumption of oysters contaminated with Norwalk-like virus. In a previous oyster-associated Norwalk virus outbreak, identification of the identical sequence of the virus genome in specimens from patients in five states established a clear link between those cases and the oysters from one harvest site (1,2). For the outbreaks described in this report, molecular analysis of fecal specimens will be required to determine the number of linked outbreaks and help assess the usefulness of the specific control measures (4,5). However, the preliminary findings suggest that Apalachicola Bay oysters may have become contaminated by sewage dumped overboard by recreational and commercial boaters. Long-term solutions to eliminate fecal contamination of oyster beds will require either that boaters not be permitted to dump sewage overboard or that beds used for harvesting be limited to those in pristine waters. Improved methods to detect virus in these oysters are needed to understand the extent of the contamination and to strengthen prevention efforts and enforcement.

Viral Gastroenteritis — Continued

#### References

- 1. Kohn MA, Farley TA, Ando T, et al. A large outbreak of Norwalk virus gastroenteritis associated with eating raw oysters: implications for maintaining safe oyster beds. JAMA 1995 (in press).
- Dowell SF, Groves C, Kirkland KB, et al. A multistate outbreak of oyster-associated gastroenteritis: implications for interstate tracing of contaminated shellfish. J Infect Dis 1994 (in press).
- 3. CDC. Viral gastroenteritis associated with consumption of raw oysters—Florida, 1993. MMWR 1994:43:446–9.
- 4. Ando T, Monroe SS, Gentsch JR, Jin Q, Lewis DC, Glass RI. Detection and differentiation of antigenically distinct small round-structured viruses (Norwalk-like viruses) by reverse transcription-PCR and Southern hybridization. J Clin Microbiol 1995;33:64–71.
- 5. Lew JF, LeBaron CW, Glass RI, et al. Recommendations for collection of laboratory specimens associated with outbreaks of gastroenteritis. MMWR 1990;39(no. RR-14).

# Notice to Readers

# **Hospital Epidemiology Course**

CDC, the Society for Healthcare Epidemiology of America (SHEA), and the American Hospital Association will cosponsor a hospital epidemiology training course May 6–9, 1995, in San Francisco. The course—designed for infectious disease fellows, hospital epidemiologists, and infection-control practitioners—provides hands-on exercises to improve skills in detection, investigation, and control of epidemiologic problems encountered in the hospital setting and lectures and seminars on fundamental aspects of hospital epidemiology.

Additional information is available from the SHEA Meetings Department, Suite 200, 875 Kings Highway, Woodbury, NJ 08095-3172; telephone (609) 845-1720; fax (609) 853-0411.

### Notice to Readers

# **Publication of Work-Related Lung Disease Surveillance Report**

CDC's National Institute for Occupational Safety and Health (NIOSH) has released the Work-Related Lung Disease (WoRLD) Surveillance Report, 1994 (1).\* This report summarizes surveillance data for occupational respiratory diseases (e.g., asbestosis, coal workers' pneumoconiosis, silicosis, byssinosis, hypersensitivity pneumonitis, and occupational asthma) and contains case-based and rate-based surveillance data, age-adjusted death rates, geographic distribution of occupational respiratory diseases, and proportionate mortality ratios.

#### Reference

NIOSH. Work-related lung disease (WoRLD) surveillance report, 1994. Morgantown, West Virginia: US Department of Health and Human Services, Public Health Service, CDC, 1994; DHHS publication no. (NIOSH)94-120.

<sup>\*</sup>Single copies of this report are available without charge from the Surveillance Section, Epidemiological Investigations Branch, Division of Respiratory Disease Studies, NIOSH, CDC, 1095 Willowdale Road, Morgantown, WV 26505-2845; telephone (304) 285-6115; fax (304) 285-6111.

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.

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.

All material in the MMWR Series is in the public domain and may be used and reprinted without special permission; citation as to source, however, is appreciated.

Director, Centers for Disease Control and Prevention
David Satcher, M.D., Ph.D.

Deputy Director, Centers for Disease Control
and Prevention
Claire V. Broome, M.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, MMWR (weekly)
Karen L. Foster, M.A.
Writers-Editors, MMWR (weekly)
David C. Johnson
Patricia A. McGee
Darlene D. Rumph-Person
Caran R. Wilbanks

☆U.S. Government Printing Office: 1995-633-175/05049 Region IV