CENTER FOR DISEASE CONTROL



MORBIDITY AND MORTALITY WEEKLY REPORT

Epidemiologic Notes and Reports

Plague – Arizona, California, New Mexico

Three bacteriologically confirmed bubonic plague cases have thus far been reported to CDC for 1978. One case was acquired in Arizona, 1 in California, and 1 in New Mexico (Table 1). None of the cases had secondary pneumonic involvement; there were no deaths.

IABLE	1. Reported	contirmed	cases of	nlamia	United	States	1978
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Case	Aye	Sex	Onset	County	State
1	22	F	March	Coconino	Arizona
2	57	F	May	Siskiyou	California
3	14	м	June	Bernalillo	New Mexico

The case history of the third patient is of particular interest because it is fairly characteristic of the majority of plague cases acquired in New Mexico. On the evening of June 11, 1978, a 14-year-old boy had onset of fever, chills, headache, dizziness, and left inquinal pain and swelling. On June 12, nausea, vomiting, and abdominal pain developed, and he was seen by his physician. The initial evaluation did not result in a specific diagnosis, and the patient was sent home with instructions to return the following day if unimproved. The patient returned on June 13 and was admitted to the hospital. Admission physical examination revealed a blood pressure of 95/50, pulse of 95, respiratory rate of 22, and temperature of 39.4 C (102.9 F). An extremely painful, left inguinal lymph node, 1½ X 2 cm in diameter, was detected; it was non-fluctuant and associated with overlying soft tissue edema without erythema. Several scratches on the trunk and extremities, not suggestive of insect bites, were attributed to working with bales of hay. The remainder of the examination was unremarkable. Clinical laboratory data revealed a white blood cell count of 11,600/mm³; the differential was 6 metamyelocytes, 51 bands, 39 segmented neutrophils, 3 lymphocytes, and 1 eosinophile. There was a mild elevation of SGOT and LDH, but the bilirubin and alkaline phosphatase were normal. A chest X ray was also normal.

Based on the history, physical examination, and preliminary laboratory studies, a tentative diagnosis of bubonic plague was made, and the patient was placed in strict isolation. Streptomycin and tetracycline therapy was begun after 2 blood cultures and a bubo aspirate were obtained. Both blood cultures and the bubo aspirate—tentatively positive for *Yersinia pestis* by direct fluorescent antibody testing on the day of admission—subsequently yielded *Y. pestis* on culture.

The patient's hospital course was unremarkable; inguinal pain remained the predominant symptom. He was removed from isolation after receiving specific antimicrobial

July 28, 1978 / Vol. 27 / No. 30

Epidemiologic Notes and Reports

- 259 Plague Arizona, California, New Mexico
- 260 Nosocomial Respiratory Syncytial Virus Infections in an Intensive Care Nursery – California
- 267 Human Rabies Texas ACIP Recommendation
- 268 Yellow Fever Vaccine

Plague - Continued

therapy for 48 hours. His temperature gradually returned to normal by June 18, and he was discharged on June 19.

No additional illness was detected in the patient's family, in a friend with whom he worked, or in that friend's family. The patient and his family resided on a ranch in Bernalillo County. They owned, in addition to horses, 2 cats and 4 dogs, a number of which were allowed to roam freely. All pet cats and dogs wore flea collars, but none of the animals had been dusted with insecticides this year. No illness was reported in any of the domestic animals.

The patient specifically denied contact with wild animals. Although he hunts, he had not done so this season. In the week before onset of illness, he had spent most of his time in or around his home. On June 9, the patient, his father, and the boy's friend had transferred hay from a storage barn to feeder stables, where horses are kept. On June 10, they replenished the storage barn with a fresh supply of hay. Although the patient denied receiving flea bites, he noted generalized pruritis after working with the hay.

Environmental investigation revealed that the ranch was located in piñon-juniper habitat. Numerous sources of man-made and naturally occurring rodent harborage were found. Although several abandoned rodent burrows were found on the premises, one, located at an unused stable adjacent to the storage barn, had numerous fleas at its entrance. Of the rodent and flea specimens submitted for culture, only pools of fleas which were obtained from the abandoned burrow at the stable have thus far been positive for Y. *pestis* on culture. No fleas were recovered from the dogs or cats.

Reported by W Milburn, MD, Albuquerque; F Malone, C Montman, Albuquerque Environmental Health Dept; J Mann, MD, State Epidemiologist, A Pressman, MD, G Graves, P Matzner, New Mexico Health and Environmental Dept; A Kelter, MD, State Epidemiologist, Arizona State Dept of Health Services; SB Werner, MD, California Dept of Health; Plague Br, Vector-borne Diseases Div, Bur of Laboratories, Field Services Div, Bacterial Diseases Div, Bur of Epidemiology, CDC.

Editorial Note: A review of potential exposures for plague cases from 1949-1976 in New Mexico reveals that often no history of exposure (e.g., eviscerating or skinning wild animals or traveling to plague-endemic areas to work) was given for the week before onset of illness. In many, the immediate vicinity of the home would have to be considered the most likely source of infection. Man-made sources of rodent harborage and food may be important in attracting wild rodents to their homes and is an area presently being investigated. The role of domestic pets in transporting infective wild rodent fleas to the home is an area requiring further investigation.

Nosocomial Respiratory Syncytial Virus Infections in an Intensive Care Nursery – California

An outbreak of upper respiratory infection and pneumonia involving 9 infants and caused by respiratory syncytial virus (RSV) occurred in a 16-bed intensive care nursery (ICN) of a hospital and medical center in San Francisco, California, from February ²⁵ through March 19, 1978 (Table 2).

On February 25, an 18-week-old premature infant with hyaline membrane disease and bronchopulmonary dysplasia developed fever with respiratory distress and had a convulsion. A nasopharyngeal viral culture taken then was subsequently positive for RSV. Two days later 2 other premature infants (aged 13 and 35 weeks) developed sneezing, cough, rhonchi, and rales. The 13-week-old was in isolation for a previously documented cytor megalovirus infection. Nasopharyngeal viral cultures from both of these infants were reported positive for RSV on March 2.

RSV Infection - Continued

TABLE 2. Clinical data on 9 infants with nosocomial RSV infection, California, Feb. 25-March 19, 1978

Patient	Age (weeks)	Sex	Date of first symptom	FA results	Culture results	Interval between +FA and +culture (days)	Fever	Upper respiratory symptoms	Pneumonia	Severity of illness
1	18	M	2/25	+	+	N/A*	ves	no	yes	severe
2	13	F	2/27	+	+	N/A*	yes	no	yes	severe
3	35	м	2/24	N/D1	+	N/A	ves	ves	ves	moderate
4	3	M	2/28	+	+	8	yes	yes	yes	moderate
5	11	F	unknown	+	-	N/A	no	no	no	unknowr
6	6	F	3/2	+	+	5	no	yes	no	mild
7	6	М	3/7	+	+	2	по	yes	no	mild
8	6 days	M	3/19	_	+	N/A	no	ves	no	mild
9**	2	F	3/10	+	+	5	no	yes	no	mild

N/A = not applicable, Patients 1 and 2 had positive RSV cultures reported before FA screening was begun.

[†]N/D = not done

•• This patient was not in the Intensive Care Nursery.

At that time the following procedures were instituted: (1) RSV fluorescent antibody (FA) screening and viral cultures were performed on nasopharyngeal swabs from all ICN patients; (2) all positive patients were isolated in a separate room; (3) strict handwashing, gowning, and gloving procedures were required before contact with all ICN patients (masking was not required); (4) certain nursing staff were assigned exclusively to infected infants; and (5) all nursing staff with upper respiratory symptoms were considered infected with RSV. If well enough to work, they were allowed to care only for alreadyinfected infants.

On the basis of direct FA screening, 2 additional infants were found positive for RSV on March 2 and were isolated. One (patient 4, Table 2) had a collapsed right upper lobe; a culture was positive for RSV. The other (patient 5) had no respiratory symptoms; 2 of ³ FA studies were borderline-positive for RSV, and none of 8 viral cultures was positive.

FA screening was repeated March 6 on the remaining 11 patients in the unit, but no new cases were identified. The following day 2 patients, both aged 6 weeks, developed mild upper respiratory symptoms. Repeat FA testing was positive on both; cultures taken at this time subsequently grew RSV.

On March 19, a 6-day-old infant who had been in the ICN since birth developed nasal congestion. FA studies were negative, and he was discharged from the hospital 2 days later. Viral cultures taken before discharge were later positive for RSV.

One additional infant developed RSV infection in association with this outbreak. The child, born on February 28, remained in the newborn nursery, a room adjoining the ICN, for 6 days because of neonatal hyperbilirubinemia. She was discharged on March 7 but was readmitted to another hospital ward on March 15 because of rhinorrhea and cough of ⁵ days' duration. FA studies and viral culture were both positive for RSV at the time of readmission. The child had had no direct contact with ICN babies during her first hospitalization. The nursing staffs of the intensive care and newborn nurseries are separate, but patients in both units are cared for by the same house staff members.

Routine viral screening of nursing and house staff members was not performed. However, from March 7-March 21 FA studies and viral cultures were performed on 2 pediatric house officers, 11 ICN nurses, 1 nursery X-ray technician, and 1 phlebotomist, who regularly bled patients in the ICN. All 15 reported upper respiratory illnesses with onset occurring from 1 to 13 days (mean 4.9 days) before viral testing. The RSV FA test was strongly positive in 1 nurse and weakly positive in 4 additional nurses and the phleboto-

RSV Infection -- Continued

mist. None of the adults was positive by culture, perhaps owing to the delay in obtaining cultures after onset of symptoms.

Reported by R Ballard, MD, WL Drew, MD, PhD, L Mintz, MD, R Roth, MD, S Sniderman, MD, Mount Zion Hospital and Medical Center, San Francisco; Respiratory Section, Respiratory and Special Pathogens Br, Viral Diseases Div, Bur of Epidemiology, CDC.

Editorial Note: Nosocomial RSV infections have been described in pediatric wards (1) and in newborn (2) and premature infant (3) nurseries. In these settings, transmission appears to occur primarily via the hands and clothing of staff members rather than by direct patient-to-patient contact or aerosol spread (1). Preventive measures such as handwashing, gowning and gloving, and isolation of infected children appear to be effective in reducing nosocomial spread (4).

Identification of RSV-infected patients by viral culture necessarily entails a delay of several days until culture results are known. The speed and sensitivity of viral isolation can be enhanced by direct bedside inoculation of tissue culture cells (5). This technique was utilized in the present outbreak and resulted in viral isolation within 3 to 8 days of culture (mean 4.9 days). The use of direct FA staining of nasopharyngeal smears, on the other hand, permitted identification of infected infants within hours of receipt of the specimen. This technique identified 7 of 8 culture-positive infants (87.5%). Direct fluorescent antibody staining was extremely useful in the rapid detection of RSV infection in infants and permitted prompt institution of specific infection-control measures.

Continued on page 267

	29th Wi	EEK ENDING		CUMULATIVE, FIRST 29 WEEKS				
DISEASE	July 22, 1978	July 23, 1977*	MEDIAN 1973-1977**	July 22, 1978	July 23, 1977*	MEDIAN 1973-1977*		
Aseptic meningitis	124	146	88	1,523	1,542	1,335		
Brucellosis	2	7	6	84	113	113		
Chickenpox	1,017	899	938	119,386	157,715	142,640		
Diphtheria	-	-	2	48	54	119		
Encephalitis: Primary (arthropod-borne & unspec.)	22	28	22	346	3 78	466		
Post infectious	6	2	5	113	116	169		
Hepatitis, Viral: Type B	258	371	240	8,154	9.171	6,221		
Type A	510	566	646	15,820	17,337	19,540		
Type unspecified	146	187	j 440	4,861	4,954			
Malaria	19	14	14	356	267	209		
Measles (rubeola)	278	499	289	21.532	51-159	23,240		
Meningococcal infections: Total	41	23	26	1.512	1.122	938		
Civilian	40	23	26	1,493	1,115	914		
Military	1 1	-	-	19	7	21		
Mumps	138	181	530	12,359	15.092	42,245		
Pertussis	39	44		997	533			
Rubella (German measles)	135	410	122	14,353	17.992	14.274		
Tetanus	1	2	3	41	35	35		
Tuberculosis	673	596	644	16,548	16,772	17,682		
Tularemia	13	5	5	57	76	79		
Typhoid fever	10	7	9	238	188	206		
Typhus fever, tick-borne (Rky. Mt. spotted)	51	72	48	501	6 0 9	411		
Venereal diseases:								
Gonorrhea: Civilian	22,490	22,265	21,423	527,339	530,099	530,099		
Military	472	541	541	13,693	15,135	15.759		
Syphilis, primary & secondary: Civilian	393	391	448	11,373	11,418	13,531		
Military	6	1	7	160	167	196		
Rabies in animals	30	61	61	1,658	1,673	1,639		

TABLE II. Notifiable diseases of low frequency, United States

	CUM, 1978		CUM. 1978
Anthrax	4	Polionyelitis: Total	58
Botulism	51	Paralytic	
Congenital rubella syndrome	20	Psittacosis (Calif. 1)	
Leprosy (Tex. 3, Calif. 1)	79	Rables in man	
Leptospirosis (Tex. 2)	33	Trichinosis (Mo. 1)	
Plague	2	Typhus fever, flea-borne (endemic, murine) (Tex. 1)	

* Delayed reports received for calendar year 1977 are used to update last year's weekly and cumulative totals.

"Medians for gonorrhea and syphilis are based on data for 1975-1977.

†Delayed report: Leptospirosis: Va. +1 (1977)

	ASEPTIC	BRU-	CHICKEN-			E	NCEPHALI	TIS	HEPATI	TIS (VIRAL), BY TYPE		
REPORTING AREA	MENIN- GITIS	CEL- Losis	POX	DIPHT	HERIA	Pri	mary	Post-in- fectious	в	A	Unspecified	MAL	ARIA
	1978	1978	1978	1978	CUM. 1978	1978	1977*	1978	1978	1978	1978	1978	CUM. 1978
NITED STATES	124	2	1,017	-	48	22	28	6	258	510	146	19	356
WEW ENGLAND	1	_	60	_	_		z	_		10	2		
	î	-	8	_	_	-	-	-	4	2	2	_	14
V.H.†	_	2	1	-	-	-		-	-	2	-	-	ź
Mass.	-		-	-	-	-	· · · ·	-	-	2	-	-	-
H.I.	_	-	27	-	-	-	2	-	3	3	2	-	3
Conn.	-	-	17	-	-		-	-	1	ī	-	-	17
			•				-	-	-	1	_	-	
MID. ATLANTIC	23	-	356		1	2	3	1	47	43	16	2	69
LY. City	9	-	251	-	-	2	-	1	5	15	4	1	10
	4	Ξ	96	_	L	_	1	-	3	2	2	-	29
Pa.	2	-	NN 9	-	-	-	2	-	27	18	7	1	15 15
E M. neu	~		,				2	-	12	•	3	-	15
E.N. CENTRAL	7		364	-	-	3	7	-	21	85	16	_	17
nd.t	2	-	66	-	-	-	1	-	11	25	-	-	3
1.	1	-	-	-	-	2	5	-	2	12	8		3
Mich.	4	1	144 40	_	-	-	-	-	2	22	-	-	4
Wis.	_	-	40	_	_	1	1	-	6	15	8	Ξ	6
WN OFFI		_	.14	-	-	-	-	-	-	11	-	-	1
W.N. CENTRAL	7	1	9	-	1	1	2	-	14	26	3	-	17
lowa	-	-	-	-	-	-	-	-	7	12	-	-	4
Mot	-	-	4	-	-	-	-	-	1	2	-	-	-
N. Dak.	6	_	3	_	1	1	-	-	5	8	1	-	6
o. Dak	-	_	1	-	-	-	1	2	-	_	-	_	1
Nebr. Kans.	1	-	_	_	-	-	1	_	1	2	1		3
Nans.		1	1	-	_		-	-	-	2	î	_	3
S. ATLANTIC													
will,	26	1	77	-	-	3	5	2	59	82	15	7	71
Md.	2	-	1	- 21	-	-		-	-	-	1	-	1
D.C.	1	-	-	_	-	_	2	-	2	4	3	-	15
Va. W. Va.	5	-	12	_		_	_	-	10	5	1	1	17
N.C.	-	L.	15	-	-	-	-	-	3	2	_	-	i
S.C.	11	-	NN	-	-	3	1	_	2	3	1	3	6
Ga.	1	-	5	-	-	-	2	-	3	3	-	-	4
Fla.	- 6	-		-	-	-	-	-	6	19		-	6
E	0	-	44	-	-	-	-	2	33	46	9	3	21
E.S. CENTRAL	11	-	10	_	-	6	_	2	27	38	1	-	3
Tenn	_	-	6	-	-	ĩ	-	-	2	-	-	_	1
Ala	5	-	NN	-	-	3	-	1	17	26	-	-	ī
Miss.	4	-	3	-	-	1	-	1	4	4	ĩ	-	ī
-	2	-	1	-	-	1	-	-	4	8	-	-	-
W.S. CENTRAL	27	-	37	_	1	5	1	_	20	58	31	1.1	19
Ark.	2		2	-	1	-	-	_	3	5	4	1	14
Okla	10	-	พท	-	-	-	-	_	ž	10			3
Tex.t	5	-	-	-	-	1	-	-	9	5	4	-	_
	LO	-	35	-	_	4	1	-	5	38	19	1	16
MOUNTAIN													= -
	-	-	75 10	_	3	_	_	_	14	70	16	1	4
ldaho Wyo,	_	-	10	_		_	_	-	1	15	-	-	_
Colo	_		_	-	_	_	-	_	_	19	-	-	-
N. Mau	-	-	30	-	2	-	-	-	3	10	3	-	1
Ariz.	-	-	-	-	-	-	-	-	2	9	-	-	1
Utah	-	-	NN	-	-	-	-	-	7	27	12	-	1
Nev.	-	-	34 1	-	-	-	-	-	-	23	-	-	
ACIFIC	_		1	-	I	-	-	-	I	د	1	-	1
	22	-	29	_	42	z	8	1	52	98	46	9	142
Uren		-	11	-	39	-	2		3	14	1	-	6
Galif +	4		-	-	-	-	-	-	8	18	3	-	3
Alaska	17		-	-	-	2	6	1	41	61	39	8	115
Hawaii	1	-	10	-	3	-	-	-	-	2	2	1	3
	-	-	8	-	-	-	-	-	-	3	1	-	15
Guam t													
+ n ,	NA	NA	NA	NA	-	NA	-	-	NA	NA	NA	NA	
V.I.	- 1	_	2	-	-	-	-	-	-	3	1	_	- 4
WN: Not notifiable	NA	NA	NA	NA	-	NA	_	-	NA	NA	NA	NA	- 1

TABLE III. Cases of specified notifiable diseases, United States, weeks ending July 22, 1978, and July 23, 1977 (29th week)

NW: Not notifiable. *Delayed reports received for 1977 are not shown below but are used to update last year's weekly and cumulative totals. The following delayed reports will be reflected in next week's cumulative totals: Asep. meng: N.H. -1, La. -1; Chickenpox: N.H. +3, Ind. +34, Calif. +1, Mont. -1, Mont. -1, La. -5, Mont +1; Hep. B: N.J. +2, Mo. -2, La. -1, Tex. +1; Hep. A: N.H. -1, N.J. +3, Ohio -1, La. -5, Mont +1; Hep. unsp.: Mo. -2, Tex. -1,

		EASLES (RU	8501 41	MENING	OCOCCAL IN	FECTIONS		AUMPS	PERTUSSIS	011	ELLA	TETANU
REPORTING AREA		· · · · · ·			TOTAL				PEHIUSSIS	RUE		-
-	1978	CUM. 1978	CUM. 1977*	1978	CUM. 1978	CUM. 1977*	1978	CUM. 1978	1978	1978	CUM. 1978	CUM. 1978
UNITED STATES	278	21,532	51,159	41	1,512	1,122	1 3 8	12,359	39	135	14,353	41
NEW ENGLAND	10	1,951	2,461	4	72	49	4	706	-	9	700	1
Maine N.H.	2	1,309	164	1	67	3		483		1	146	-
Vt.	-	25	290	-	2	4		11	_	1	27	ī
Mass.t	8	247	618	1	18	17	2	82	-	5	203	-
R.I.	_	7	61 818	2	16	1	2	31	-		40	-
Conn.	-	318	919	2	23	21	-	94	-	2	186	-
MID. ATLANTIC	47	2,013	8,111	7	254	147	3	529	7	23	2,805	2
Upstate N.Y.	19	1,317	3,696	3	82	32	-	184	3	2	488	1
N.Y. City N.J. †	24	295	193	1	62 64	40 32	3	127	4	9	105	
Pa.	-	332	3,560	2	66	43	-	92	-	3	635	1
E.N. CENTRAL	0 3	0 374	14 300	-	136	1.20	6.7	4	,	50		
Ohiot	83	9,274 466	10,399	2	49	120	52 16	4,902 771	1	50 12	6,664 1,327	2
Ind. 1	-	165	4,276	-	26	8	-	271	-	-	523	i
UI.	8	581	1,515	-	6	31	9	1,619	-	2	411	-
Mich. Wis.	62 13	6.652 1.410	908 2,363	-	44 11	32 12	7 20	1,320 921	-	20 16	2,923	-
						12	20	721	-	10	1.4400	-
W.N. CENTRAL	5	375	9,376	2	53	52	10	1.880	4	10	620	5
Minn. Iowa	2	34 51	2,596	2	12	19	2	17	4	3	127	1
Mo. 1	-	11	1,033	-	23	15	8	1,148	_	í	49	
N. Dak.	-	185	22		3	1	-	11	-	1	80	-
S. Dak.	-	- 5	66		2	4	-	6	-	-	110	-
Nebr. Kans.	3	85	192 1,208	2	8	1 5	-	18 560	-	3	34 129	
S. ATLANTIC	64	4,667	4,383	14	389	259	19	663	5	19	966	6
Del. Md.	5	5	22 371	2	12	17	3	48	-	-	34	
D.C.	-	42	14	-	1	17	3	60 1		<u> </u>	- 6 1	1
Va.	S	2,787	2,595	1	48	19	4	119	-	1	230	-
W. Va. N.C.	3	1,009	206	1	8 78	9 58	1	153	-	3	322	
S.C.	2	111	60 146	3	24	26	-	56 15	2	10	178 26	1
Ga.	2	17	760	1	45	37	1	62	3	1	3	-
Fla.	43	503	209	6	154	76	7	149	-	3	166	3
E.S. CENTRAL	13	1,349	1,921	4	121	127	25	1.050	-	3	479	1
Ky.		115	1,156	-	23	26	-	179	-	-	122	1
Tenn. Ala	11	937 89	657 77	1	30 37	30 47	2 20	433 373	-	3	189	
Miss.	2	208	31	ĩ	31	24	3	65	-	-	147	
W.S. CENTRAL	7	933	2.017	5	233	197	10	1,587	12	4	878	13
Ark. La. †	2	16 316	29 74	1	21 93	9 76	- 2	577 60	3	-	57	1
Okla.	-	13	54	-	16	10	- -	4	-	_	482	2
Tex.	5	588	1,860	2	103	102	8	946	9	4	328	9
MOUNTAIN	3	235	2,474	1	32	29	10	366	4	1	186	1
Mont.	-	102	1,152	-	1	2	-	136	2	-	17	-
Idaho	-	1	161	-	3	4	~	20	-	-	2	-
Wya. Cola.	1	29	15 496	_	2	1	-	74	-	-	-	-
N. Mex. t	-		253	_	7	7	_	15	-	_	43	1
Ariz.	ı	44	296	-	11	10		10	1	1	99	-
Utah Nev	1	44 15	8 93	-	4	3	10	107	-	-	23	1
					-							
PACIFIC Wash.	4 E 3 6	735	1C,017 524	2	222	142	5	676	6	16	1,055	10
Oreg.	- 0	140	347	ĩ	20	17	-	164 76	1	1	93 91	1
Calif.	9	456	9,052	i	154	80	5	406	5	15	863	10
Alaska		-	60		5	25	-	6	-	-	2	-
Hawaii	1	5	34	-	4	2	-	24	-	-	6	-
Guam t	NA	24	4	-	-	_	NA	31	NA	NA	1	1
P. R.	4	199	827	-	2	1	25	1,034		-	15	5
V.1.	NA	6	14	-	1	-	NA	1	NA	NA	1	-

TABLE III (Cont.'d). Cases of specified notifiable diseases, United States, weeks ending 40 • • a 1..... 22 1077 /204

NA: Not available. *Delayed reports received for 1977 are not shown below but are used to update last year's weekly and cumulative totals. †The following delayed reports will be reflected in next week's cumulative totals: Measles: Mass. -2, Ind. +1; Men. inf.: La. -2; Mumps: Ind. +2, Mo. -1; Pertussis: N.J. +1, Ind. +2, Mo. +5, N. Mex. +2; Rubella: Ohio -86, Ind. +9, La. -3, Guam +2.

Free	THRE	RCULOSIS	TULA	TYP	ного	TYPHU	SFEVER		VENERE	AL DISEASES ((Civilian)			RABIES
REPORTING AREA	1000	htutuala	REMIA	FE	VER		borne) ASF)		GONORRHEA		SY	PHILIS (Pri.	& Sec.)	(in Animals)
-	1978	CUM. 1978	CUM. 1978	1978	CUM. 1978	1978	CUM. 1978	1978	CUM. 1978	CUM. 1977*	1978	CUM. 1978	CUM. 1977"	CUM. 1978
UNITED STATES	673	16,548	57	10	238	51	501	22,490	527,339	530,099	393	11,373	11.418	1,65
NEW ENGLAND	16	535	-	2	38	-	9	396	13,568	13,725	12	337	479	6
Mainet N.H.	2	38	-	-	-	-	-	40	1,031	987	-	8	14	
Vt.	-	10 23	-	-	5	2	-	24	626 314	545	1	4	3	
Mass.	6	305	_	2	23	-	4	136	5,949	362 5,410	7	207	6 338	
R.1.	4	39	-	-	4	-	i	44	966	1,154	2	16	7	
Conn.	4	120	-	-	5	-	4	147	4,682	4.767	3	99	111	
MID. ATLANTIC	109 15	2,843	3	3	26 7	3	28	2,316	56,681	53,706	44	1,528	1,598	
N.Y. City t	53	1,004	ĩ	3	13	-	2	841	9.454 22.220	8,914 21,323	33	111	151	
N.J.	14	711	-	_	-4	2	4	596	10,717	9,268	ŝ	170	207	
Pa,	27	701	-	-	2	-	7	601	14.290	14,201	6	161	236	
E.N. CENTRAL Ohio	119	2,528	1	-	11	1	14	3,733	78,819	82,272	34	1,227	1,225	
Ind,	16	463	I	-	5	1	9	906	20,680	21,394	2	228	279	
10.	11 40	305 962	-	-	-	-	1	621	8,121	7,638	8	67 770	92	
Mich.t	52	692	_	-	5	_	4	1,125	24,572 18,231	26,750 18,919	16	123	660 138	
Wis.	-	106	-		-	-	-	317	7,215	7,571	2	39	56	
W.N. CENTRAL	15	557	10	-	11	1	12	1,170	26,631	27,565	7	272	255	362
lowa	1	105	-	-	4	-	-	282	4,630	4,985	2	113	81	121
Mo, t	3	61	_	-	2	-	-	101	3,002	3,214	-	32	24	
N. Dak.	2	233	9	_	3	1	8	517 27	11.397	11,684	4	74	88	
S. Dak.	î	47	-	_	_	-	1	59	489 954	513 743		2	2	
Nebr. t		11	-	-	-	-	-	34	1,940	2,363	ı	8	24	
Kans.	7	73	ì	-	2	-	3	150	4,219	4,063	-	42	34	
S ATLANTIC	167	3,580	5	4	34	31	289	5.544	127,330	131,764	107	3,027	3,250	221
Md.t	3	29		-	1	-	4	57	1,750	1,810	1	6	17	
D.C	26 3	549 192	4	2	5	5	63	741	16,308	16,386	4	240	211	
Va.t	32	383	1		1	14	66	314 458	8,378 12,056	8,707 13,451	8	241 255	339 323	
W. Va. N.C. t	6	156	-	-	2	ì	9	78	1,842	1,814	-	6	1	
S.C. t	33	542		-	2	8	94	1,100	18,382	19,440	12	291	465	
Ga.	6	312	-	1	4	3	30	540	12,622	12,108	9	156	143	
Fla.†	19 39	483 934	-	1	3 10		23	1,406 850	23,674 32,318	25,858 32,190	21 47	733 1,097	651 1,100	
E.S. CENTRAL	66	1,552	5		5		90	1,933	45,662	47,714	20	578	401	
Ky. Tenn.	16	352	ź	_	2	2	28	176	5,580	6,557	20	74	50	
Ala.	10	463	3	-	ī	8	54	615	10,715	19,283	ś	199	124	
Miss.	16	374		-	1	1	5	526	13,235	12,981	1	66	73	
W.S. CENTRAL	24	363	-		1	-	3	616	10,132	8,893	9	217	154	
	80	1,939	28	-	27	4	54	2,968	73,081	67,381	112	1,827	1,571	543
La.t	3	213	19	-	2	-	8	336	5,530	5:056	-	46	37	76
Okla, †	20 8	324	5	_	2	1	33	487	12,065	10,105	44 2	396	362	
Tex.t	49	1,200	1	-	21	-	12	1,911	48,615	6,291 45,929	66	53 1,332	43 1,129	
MOUNTAIN	30	486	3	-	12	_	4	900	19,668	21,444	9	223	227	21
daho	_	31	1	-	-	-	2	49	1,163	1.067	_	1	3	
Nyo.	-	20	2	-	5	-	ī	36	733	1,007	-	6	5	
Glat	1	12	-	-	-	-	-	17	440	517	-	4	2	-
N. Mex.	1	43		-	2	_	-	229	5.442	5,519	5	65	70	
Ariz, Utah	22	238		12	1 2	_	-	101 324	2,861 5,083	3,188	-	54 48	40 94	
Vev.	2	25	1	-	ĩ	_	_	27	1,059	1,186	-	48	94	12
	2	41	-	-	i	-	1	117	2,887	2,796	4	28	8	-
ACIFIC Vash. t	71	2,528	2	1	74	_	1	3,530	85,849	84,528	48	2,354	2,412	
Dreg	-	112	-	-	6	-	-	233	61685	6,276	-	2,354	127	
alif	7	111	-	-	ĩ	-	-	228	5,921	5,802	-	79	68	
Alaska t	51	1,930	2	1	60	-	1	2,924	68,945	07.878	49	2,165	2,177	
Hawaii	- ū	46 329	1	120	7	ΞĘ.	_	83 62	2.749 1.599	2,770	-	7 23	15	e
	-				•			95					24	
Guant P.R.	NA	33	-	NA	-	NA		NA	108	1 30	NA	-	1	
V.I.	5	240	-	-	1	-	-	29	1,268	1,788	6	249	310	15
NA: Not available	ΝA	4		NA	2	NA	-	ΝA	112	115	NA	9	5	-

TABLE III (Cont.'d). Cases of specified notifiable diseases, United States, weeks ending July 22, 1978, and July 23, 1977 (29th week)

NA: Not available "Delayed reports received for 1977 are not shown below but are used to update last year's weekly and cumulative totals. The following delayed reports will be reflected in next week's cumulative totals: T8: NYC +71, Mich. -2, Nebr. +1, Md. -3, N.C.-2, S.C. -1, Fia. -3, Guan, 42, T. fetter: Mo. +1, Va. -1, Tex. -1, Colo. +1; RMSF. Md. -1, Va. -1; GC: Md. +2; Cuiv. La. -2; civ., Okla +100 mil., Wash. +84 mil., Alaska -1 civ +1 mil., Guam +5; Syphilis: Maine -1 civ., Md. -5 civ., Wash. +22 civ. +2 mil.; An. rables: Wash. -1.

TABLE IV. Deaths in 121 U.S. cities,* week ending July 22,1978 (29th week)

		ALL CAUSE	S, BY AGE	(YEARS)					ALL CAUS	SES, BY AG	E (YEARS)		
REPORTING AREA	ALL AGES	>65	45-64	25-44	<1	P&I** TOTAL	REPORTING AREA	ALL AGES	>65	45-64	25-44	<1	P&I** Total
NEW ENGLAND	596	351	1 32	35	17	24	S. ATLANTIC	1,151	645	338	85	29	61
Boston, Mass.	175	97	47	14	9	9	Atlanta, Ga.	128	55	52	15	1	6
Bridgeport, Conn.	41	31	9	1	-	3	Baltimore, Md.	259	142	80	23	6	- 7
Cambridge, Mass. Fall River, Mass.	14	5	9	-	-	1	Charlotta, N.C. Jacksonville, Fla.	55	35	9	2	1	- 1
Hartford, Conn.	23 43	21	2	- ĩ	3	1	Miami, Fla.	112	63 34	31 28	6	4	6
Lowell, Mass.	18	12	ŝ	1	_	-	Norfolk, Va.	64	32	20	4	6	7
Lynn, Mass.	23	18	4	_	-	1	Richmond, Va.	68	36	22	6	2	11
New Bodford, Mass.	23	14	6	1	1	-	Savannah, Ga.	39	29	5	-	-	3
New Haven, Conn.	42	30	5	4	1		St. Petersburg, Fla.	88 67	66	18	1	1	45
Providence, R.I. Somerville, Mass.	58 6	41	11 3	4	2	5	Tampa, Fla. Washington, D.C.	154	43 86	16	14	1 4	1
Springfield, Mass.	52	35	10	4	1	1	Wilmington, Del.	46	24	13	3	ī	2
Waterbury, Conn.	32	27	2	ż	î	ź					-	-	
Worcester, Mass.	46	31	8	3	-	L							
							E.S. CENTRAL	772	459	214	54	15	28
MID. ATLANTIC						0.5	Birmingham, Ala.	137	60 35	35	9	7	1
Albany, N.Y.	21456	1,535	605 12	167	69 1	85	Chattanooga, Tenn. Knoxville, Tenn.	50	40	9	1	1	3
Allentown, Pa.	29	9	11	7	-	-	Louisville, Ky.	120	72	34	3	2	7
Buffalo, NLY.	121	70	25	14	4	9	Memphis, Tenn.	158	87	39	21	2	5
Camden, N.J.	39	21	12	1	1	-	Mobile, Ala.	68	52	27	7	-	4
Elizabeth, N.J.	22	16	5	l	-	-	Montgomery, Ala.	35	22	11	1	1	1
Erie, Pa. Jersey City, N.J.	24	16	.7	-	-	-	Nashville, Tenn.	127	71	42	9	2	4
Newark, N.J.	37	22 23	11	3	1 9	1							
N.Y. City, N.Y.	1,251	795	307	65	24	36	W.S. CENTRAL	1,364	754	354	113	61	31
Paterson, N.J.	33	22	1	2	2	3	Austin, Tex.	62	43	13	3	-	4
Philadelphia, Pa.†	339	198	94	24	12	12	Baton Rouge, La.	39	21	11	4	1	1
Pittsburgh, Pa. Reading, Pa.	68	45	19	1	2	2	Corous Christi, Tex.	33	14	9	5	4	1
Rochester, N.Y.	42 125	33 85	6 25	3	-	1	Dallas, Tex.	216 61	118 28	59 15	17	9	2
Schenectady, N.Y.	25	17	25	2	6	6 1	El Paso, Tex. Fort Worth, Tex.	106	20 60	25	10	5	1
Scranton, Pa.	31	21	9	-	1	3	Houston, Tex.	291	134	86	34	15	4
Syracuse, N.Y.	85	54	16	4	5	-	Little Rock, Ark.	57	34	13	4	3	6
Tronton, N.J.	32	16	13	3		4	New Orleans, La.	155	82	51	7	8	-
Utica, N.Y. Yonkers, N.Y.	18	17	-		1	3	San Antonio, Tex.	187	115	39	13	6	43
Galkera, N. T.	26	20	5	1	-	1	Shreveport, La. Tulsa, Okla.	61 96	40 65	14	5	2	5
E.N. CENTRAL	2.174	1,291	546	154	78	49							
Akron, Ohio	71	40	21	5	ī		MOUNTAIN	559	329	142	38	26	16
Canton, Ohio	33	22	8	2	1	1	Albuquerque, N. Mex.	61	27	16	8	4	3
Chicago, III.	\$32	302	140	50	17	12	Colo. Springs, Colo.	24	14	6	3	1	3
Cincinnati, Ohio	157	109	37	6 10	4	3	Denver, Colo. Las Vegas, Nev.	107	66 26	21	8	5	-
Cleveland, Ohio Columbus, Ohio	136	83	34	9	5	8	Ogdan, Utah	22	16	- 4	ĩ	1	3
Dayton, Ohio	117	60	37	- ní -	5	1	Phoenix, Ariz.	149	87	39	12	6	1
Detroit, Mich.	247	131	74	21	4	5	Pueblo, Colo.	17	10	7	-		-
Evansville, Ind.	32	23	6	2	-	-	Salt Lake City, Utah	71	41	22	-	6	2
Fort Wayne, ind.	56	37	9	2	3		Tucson, Ariz.	62	42	13	2	3	
Gary, Ind. Grand Rapids, Mich.	24 40	14 25	13	4	1	1							
Indianapolis, Ind.	131	23	34	Ĝ	6	_	PACIFIC	1,562	965	367	108	58	31
Madison, Was.	39	25	8	ī	3	4	Berkeley, Calif.	15	13	2	-		1
Milwaukee, Wis.	121	89	21	4		1	Fresno, Calif.	65	35	19	3	2	1
Peoria, III.	47	27	9	4	3	3	Glendale, Calif.	10	9	1		-	1
Rockford, III. South Bend, Ind.	44 35	28 16	9 13	4	1	1	Honolulu, Hawaii Long Beach, Calif.	36	15	12	4 7	4	3
Tolado, Ohio	55	61	20	8	ž	ī	Los Angeles, Calif.	505	330	1 02	37	- 11	6
Youngstown, Ohio	61	43	8	3	5	-	Oakland, Calif.	72	44	18	5	4	-
							Pasadena, Calif.	20	16	2	-	2	1
WAL AFTITOA:							Portland, Oreg	119	82	24	6	4	1
W.N. CENTRAL Des Moines, Iowa	750	468	167	47	36 3	22	Sacramento, Calif. San Diego, Calif.	55 126	30 81	16 28	5	-	3
Duluth, Minn.	26	20	13	2	_	1	San Diego, Calif. San Francisco, Calif.	120	93	40	12	5	3
Kansat City, Kans.	33	19	6	ź	3	1	San Jose, Calif.	56	30	14	- 6	2	2
Kansas City, Mo.	113	78	22	6	3	ż	Seattle, Wash.	143	80	34	6	16	7
Lincoln, Nebr.	21	11	7	2	-	1	Spokane, Wash.	57	34	14	4	2	4
Minneapolis, Mirsn.	82	55	26	5	8	2	Tacoma, Wash.	36	21	11	1	2	
Omaha, Nebr.	68 187	41	13 35	5 16	8 8	2							
St. Louis, Mo. St. Paul, Minn.	187	42	20	10	1	2	TOTAL	11,384	6.837	2.845	801	389	347
Wichita, Kans.	65	36	21	2	ż	6				_,,			
							Expected Number	10,923	6, 509	2,918	716	426	353

*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 +Data not availabe this week. Figures are estimates based on average percent of regional totals.

July 28, 1978

MMWR

RSV Infection -- Continued

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Epidemiologic Notes and Reports

Human Rabies – Texas

On June 17, 1978, a 25-year-old male Mexican national died of rabies in Fort Bend County, near Houston, Texas. The diagnosis was confirmed by fluorescent microscopy by the Texas Department of Health Regional Laboratory in Houston.

The patient and 3 of his companions, residents of the State of Guanajuato, Mexico, had departed their home on approximately May 22 and had reached the Fort Bend area approximately June 1. There they found employment on a ranch.

On June 13 the patient complained of sore throat and vomiting; symptoms worsened the next day, and he was taken to a physician. At that time he was perspiring profusely, salivating excessively, gagging, and complaining of stomach pains and air blowing on his face. The physician noted an inflamed throat but no other specific signs. The following day the patient again visited the physician at which time he appeared apprehensive and unable to swallow. No fever or other positive findings except inflamed throat were noted. He was treated symptomatically.

By June 16, he had convulsive-type seizures, was at time irrational, complained of severe headache, and had difficulty breathing. He was taken to the emergency room of a nearby hospital where he was afebrile and found to have an inflamed throat and enlarged cervical lymph nodes. He was treated symptomatically and released. By evening the patient became violent and was exhibiting bizarre behavior. He became comatose the next day and died at 3:45 PM.

Fourteen persons are receiving antirables treatment as a result of exposure to this individual: 9 ranch employees with whom he shared eating utensils and had continuous close contact for the 2 weeks before his death, as well as 3 physicians and 2 deputy sheriffs who were exposed during treatment and restraint of the patient.

Extensive questioning of the patient, his companions who traveled with him from Mexico, and fellow ranch hands in Texas, failed to identify any animal bite or other kind of possible rabies exposure.

Reported by RE Pope, DVM, NM Sisley, MD, Public Health Region 8, CR Webb Jr, MD, State Epidemiologist, Texas Dept of Health; Viral Diseases Div, Bur of Epidemiology, CDC.

Editorial Note: Since 1966 (12 years) a total of 19 other human cases of rabies have been reported in the United States. In 1 instance the source of exposure, as in this case, was unknown. For the other cases the sources of exposure were dog bites (7), wildlife (8), a cat (1), and laboratory accidents (2).

Recommendation of Public Health Service Advisory Committee on Immunization Practices

Yellow Fever Vaccine

INTRODUCTION

At present, cases of yellow fever are reported only from Africa and South America. Two forms of yellow fever—urban and jungle—are distinguishable epidemiologically. Clinically and etiologically, they are identical.

Urban yellow fever is an epidemic viral disease of man transmitted from infected to susceptible persons by a vector, the *Aedes aegypti* mosquito. With the elimination or suppression of *A. aegypti*, urban yellow fever has disappeared from previously epidemic foci.

Jungle yellow fever is an enzootic viral disease transmitted among non-human hosts by a variety of mosquito vectors. It is currently observed only in the jungles of South America and Africa, but in the past it extended into parts of Central America as well. Human cases are rare. The disease can ostensibly disappear from an area for years and then reappear. Delineation of affected areas depends upon surveillance of animal reservoirs and vectors, accurate diagnosis, and prompt reporting of all cases.

Urban yellow tever can be prevented by eradicating *A. aegypti* mosquitos or by suppressing their numbers to the point that they no longer perpetuate infection. At the present time, jungle yellow fever can most effectively be prevented in humans by immunization.

YELLOW FEVER VACCINE

Yellow fever vaccine^{*} is a live, attenuated virus preparation made from 1 or 2 strains of virus: 17D and Dakar (also referred to as French neurotropic). The Dakar strain has been associated with a significant (0.5%) incidence of meningoencephalitic reactions and is not recommended. The 17D strain has caused no significant complications.

Licensed vaccine available in the United States is prepared from the 17D strain, which is grown in chick embryo inoculated with a seed virus of a fixed passage level. The vaccine is freeze-dried supernate of centrifuged embryo homogenate.

Vaccine should be stored at the temperature recommended by the manufacturer (<5 C) until it is reconstituted by the addition of sterile physiologic saline. Unused vaccine should be discarded within approximately 1 hour of reconstitution.

VACCINE USAGE

Vaccine Recipients

- Persons 6 months of age or older traveling or living in areas where yellow fever infection exists—currently parts of Africa and South America. (These are listed in the Weekly Summary of Countries with Areas Infected with Quarantinable Diseases—available in most health departments.)
- 2. Laboratory personnel who might be exposed to virulent yellow fever virus.

Vaccination for International Travel

For purposes of international travel, the yellow fever vaccines used must be approved by the World Health Organization and administered at an approved Yellow Fever Vaccination Center. (Centers can be identified by contacting state and local health departments and/or the Public Health Service.) Vaccinees should have an international Certifi-

July 28, 1978

MMWR

Yellow Fever Vaccine - Continued

cate of Vaccination filled in, signed, and validated with the stamp of the center where the vaccine is given.

Vaccination for international travel may be required under circumstances other than those specified herein. Some countries in Africa require evidence of vaccination from all entering travelers. Some countries may waive the requirements for travelers coming from non-infected areas and staying less than 2 weeks. These requirements may change, so all travelers should seek current information from health departments, travel agencies, international airlines, or shipping lines.

Some countries require an individual, even if only in transit, to have a valid International Certificate of Vaccination if he or she has been in countries either known or thought to harbor yellow fever virus.

Primary Immunization

For both adults and children, a single subcutaneous injection of 0.5 ml of reconstituted vaccine is used.

Booster Doses

Yellow fever immunity following vaccination with 17D strain virus persists for more than 10 years; the International Health Regulations do not require revaccination more often than every 10 years.

PRECAUTIONS AND CONTRAINDICATIONS

Reactions to 17D yellow fever vaccine are generally mild. Five to 10% of vaccinees have mild headache, myalgia, low-grade fever, or other minor symptoms 5-10 days after vaccination. Fewer than 0.2% curtail regular activities. Although more than 34 million doses of vaccine have been distributed, only 2 cases of encephalitis have been reported in the United States.

Contraindications

Pregnancy: Although specific information is not available concerning adverse effects of yellow fever vaccine on the developing fetus, it is prudent on theoretical grounds to avoid vaccinating pregnant women and to postpone travel until after delivery.

The morbidity and mortality from yellow tever disease are not altered by pregnancy. Therefore, pregnant women who **must** travel to areas where the risk of yellow fever is high should be vaccinated. It is believed that under these circumstances, the small theoretical risk for mother and fetus from vaccination is far outweighed by the risk from Yellow fever infection.

Altered immune states: Infection with the yellow fever vaccine virus could pose excessive risk to patients with leukemia, lymphoma, or generalized malignancy or to those Whose immunologic responses are suppressed by steroids, alkylating drugs, antimetabolites, or radiation.

Hypersensitivity: Live yellow fever vaccine is produced in chick embryo cell culture and should not be given to persons clearly hypersensitive to eggs. Furthermore, it should

The Morbidity and Mortality Weekly Report, circulation 78,750, is published by the Center for Disease Control, Atlanta, Georgia. The data in this report are provisional, based on weekly telegraphs 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. Send reports to: Center for Disease ^{Control}, Attn: Editor, Morbidity and Mortality Weekly Report, Atlanta, Georgia 30333.

Send mailing list additions, deletions, and address changes to: Center for Disease Control, Attn: Distribution Services, GSO, 1-SB-36, Atlanta, Georgia 30333. When requesting changes be sure to give your former address, including zip code and mailing list code number, or send an old address label.

Yellow Fever Vaccine - Continued

not be given to persons hypersensitive to other vaccine components such as trace amounts of particular antibiotics. (See the manufacturer's labeling.)

If international travel regulations are the only reason to vaccinate a patient hypersensitive to eggs, efforts should be made to obtain a waiver. A physician's letter clearly stating the contraindication to vaccination has been acceptable to some governments. (Ideally, it should be written on letterhead stationery and bear the stamp used by health departments and official immunization centers to validate International Certificates of Vaccination.) Under these conditions, it is also useful for the traveler to obtain specific and authoritative advice from the country or countries he or she plans to visit. Their embassies or consulates may be contacted. Subsequent waiver of requirements should be documented by appropriate letters.

SIMULTANEOUS ADMINISTRATION OF LIVE VIRUS VACCINES

See the Advisory Committee on Immunization Practice Statement, "General Recommendations on Immunization," in MMWR 25:349-350, 355, 1976.

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270