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EXPERIMENTAL TRANSMISSION OF ENDEMIC TYPHUS FEVER OF THE UNITED STATES BY THE RAT FLEA (XENOPSYLLA CHEOPIS)

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The importance of the rat flea as a vector of endemic typhus fever in the United States has been shown by the recovery of the virus of this disease from fleas taken from wild rats trapped at typhus foci in Baltimore and Savannah (1) (2). The importance of these observations has been emphasized by the recovery of typhus virus from wild rats by Mooser, Castaneda, and Zinsser (3).

Kemp has recently confirmed our observations by the recovery of typhus virus from rat fleas taken at typhus foci in Texas (4).

A preliminary report on the transmission of endemic typhus virus from rat to rat by means of the rat flea (Xenopsylla cheopis) has been made (5). In the experiments described in that report fresh noninfected white rats were exposed in glass boxes to rats infected with endemic typhus and to rat fleas. The brains and spleens from the fresh rats, on inoculation into guinea pigs, produced a reaction clinically identical with the reaction of endemic typhus. strains have since been shown to be identical with endemic typhus by the production of agglutinins for proteus X19, type O, in monkeys and rabbits, by the presence of rickettsiae in smears from the tunica vaginalis of guinea pigs, by the presence of the typical lesions in the brains of guinea pigs, and by cross-immunity tests with known strains of endemic typhus. Since the publication of the preliminary report above mentioned, we have recovered the virus of endemic typhus from white rats exposed to infected rat fleas but not exposed to infected rats. In these experiments fleas (Xenopsulla cheopis) infected with typhus virus were placed in freshly sterilized boxes with That the virus recovered from these fresh rats is the fresh white rats. virus of endemic typhus has been shown by the production of agglutinins for proteus X₁₉, type O, in monkeys and rabbits, by the presence of rickettsiae in smears from the tunica vaginalis of guinea pigs, by the presence of the typical pathological lesions of endemic typhus in the brains of guinea pigs, and by cross-immunity tests with known strains of endemic typhus.

It has further been found that the typhus virus is present in the flea for at least nine days after feeding on infected rats.

Typhus virus has also been repeatedly recovered from the feces of infected fleas. Rickettsiae have been observed in smears from infected fleas.

CONCLUSION

The virus of endemic typhus has been experimentally transmitted from rat to rat by means of the rat flea (Xenopsylla cheopis).

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AGGLUTININ ABSORPTION IN UNDULANT FEVER (BRUCELLOSIS)¹

By Edward Francis, Medical Director, United States Public Health Service

Likeness is the quality which has combined into the one genus, *Brucella*, the three microorganisms which affect primarily goat, cow, and hog, but which are transmissible to man and other animals.

TABLE 1.—Comparison of two methods of classification

Culture	Pathological source of cul- ture	Classification of culture by the National Institute of Health by reciprocal ag- glutinin absorption	Classification of cul- tures by Dr. I. F. Huddleson by the bacteriostatic action of dyes
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41-1	do	do	Do.

¹ From the National Institute of Health, formerly the Hygienic Laboratory, Washington, D. C.

Differences must necessarily be found as a basis for separation into species. Hence, the striving to find differential characters which might serve to trace each organism, wherever found, back to its original source—goat, cow, or hog. Differences have been sought in cultural characteristics, serological reactions, pathological changes, hydrogen sulphide metabolism, bacteriostatic action of dyes, and utilization of dextrose.

Table 2.—Absorption of agglutinins from melitensis type serum and from abortus type serum by cultures to be classified

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melitensis type serum 428				İ																	l
Not absorbed Absorbed by— Melitensis 428. Culture McC. Culture H-1 Culture 41-2 Culture H-5 Culture B-8 Culture B-9 Culture B. H. W Culture E. F. Culture C-10. Culture C-10. Culture C-10. Culture C-10. Culture C-10. Culture 634. Culture 635. Culture 635. Culture 426. Culture L. Z. Culture L. M. Culture L. C. Culture L. M. Culture L. C. Culture L. C. Culture L. C. Culture A. B. C. Culture 41-1	000000000000000000000000000000000000000	000000000000000000000000000000000000000	000000000000000000000000000000000000000	000000000000000000000000000000000000000	000000000000000000000000000000000000000	000000000000000000000000000000000000000			10000000000000000000000000000000000000	00000000000000	000000000000000000000000000000000000000	000000000000004444444444444444444444444	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0			McC	000000000000000000000000000000000000000	000000000000000000000000000000000000000	000000000000000000000000000000000000000	0000000 10 1000000
ABORTUS TYPE SERUM 456 Not absorbed Absorbed by Melitensis 428 Culture McC Culture H-1 Culture 41-2 Culture H-5 Culture B-8 Culture B-8 Culture R. H. W Culture E. F. Culture W. T. H. Abortus 456 Culture 83 Culture 634 Culture 634 Culture 634 Culture 426 Culture 426 Culture E. M Culture E. M Culture E. M Culture A. B. C Culture A. B. C Culture A. B. C Culture 41-1	434444444444444444444444444444444444444	44444 00000000000	444444444444444444444444444444444444444	444444444444444444444444444444444444444	4 4 1 4 4 0 4 3 4 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0	0	000000000000000000000000000000000000000	000000000000000000000000000000000000000	0000000 0 00000000	4 000000000 0 000000000000	4 0000000000000000000000000000000000000	000000000000000000000000000000000000000			H-5 B-8 C-15 89 R. H. W. E. F W. T. H.	130200010010010010010010010010010010010010	100200000 10 1 100000	10000000 10 1000000	000000000000000000000000000000000000000

With the view of testing the agreement of two methods of classification, an exchange of cultures under key numbers was effected between Dr. I. Forest Huddleson, of East Lansing, Mich., and the National Institute of Health, each being furnished with the history of a culture only after rendering a report upon its key number.

Table 1 shows the classification of 22 Brucella cultures by the method of agglutinin absorption at the National Institute of Health, in comparison with the classification of the same cultures by the bacteriostatic action of dyes in the hands of Doctor Huddleson. The results of the two methods are not exactly comparable, because the agglutinin absorption test makes no claim of being able to separate bovine abortus from porcine abortus, of which latter there were four cultures, according to the dye method. Of the remaining 18 cultures there was agreement in the classification of 9 and disagreement in the classification of 9. The latter 9 are therefore the cultures of greatest interest, 6 of which reacted as melitensis by agglutinin absorption but reacted as bovine abortus to the dyes, while the remaining 3 reacted as abortus by absorption, but reacted as melitensis to the dyes. Which method is entirely correct, if either, is not clear.

TECHNIQUE

The method employed in performing agglutinin absorption tests was essentially that used by Alice C. Evans (1).

Antiserums.—Type serums were prepared from rabbits by the intravenous injection of type cultures of Brucella abortus 456 and Brucella melitensis 428. For the purpose of reciprocal absorption tests, rabbit serums were prepared with each of the cultures to be classified. A uniform titer of 1:640 against the organism injected is desirable in all serums which are to be absorbed, because this contributes very materially to uniformity in results, thus aiding in the interpretation of results.

Six days after a single intravenous injection of a killed culture, a preliminary test of aggultinins was made on a sample of blood taken from the ear. This test usually showed a titer of 1:640, or a little higher, against the organism injected; in this case the rabbit was etherized and bled to death from the heart or carotid artery on the sixth day. If the titer on the sixth day was only 1:320, killing was postponed two or three days. In the latter case the titer frequently reached 1:1280 or 1:2560.

An occasional poor agglutinogenic culture required two or three injections, separated by intervals of two days, in order to produce a titer of 1:640 in a rabbit. The material for each injection consisted of 0.2 c. c. of a formalin-killed concentrated stock suspension of organisms having a turbidity of 25,000 according to the silica standard. This amount of antigen was diluted with physiological

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saline solution to a volume of 2 c. c. and injected slowly into the marginal vein of a rabbit's ear.

Clearness of a serum and freedom from opalescence are promoted by withholding all food from a rabbit for 24 to 48 hours before bleeding. Serums used in these studies were preserved by adding only a small platinum loopful of pure tricresol to each 5 c. c. of serum. Glycerin is not only a good preservative but has the advantage of clearing a cloudy serum. An equal amount of pure undiluted neutral glycerin should be added to a serum if glycerin is used as a preservative. Serums were not inactivated. Preserved serums were stored at 10° C.

Preparation of stock antigens.—All cultures were grown on beef infusion glucose agar in Blake bottles for three days at 37° C. Each bottle was inoculated with the entire growth from a glucose agar slant suspended in 1 c. c. of saline solution. In case an organism required carbon dioxide, several cotton-stoppered Blake bottles were placed within a large desiccator, the air was exhausted from the desiccator, a volume of carbon dioxide was admitted approximately equal to 10 or 15 per cent of the volume of the desiccator, and finally air was admitted to replace all vacuum. At the end of 72 hours the growth was washed off in saline solution containing 1 per cent commercial formalin, using 20 c. c. per Blake bottle. After standing six days the formalinized suspension of organisms was thrown down in the centrifuge, the clear supernatant fluid was poured off, and the bacterial mass was resuspended in saline solution containing 0.5 per cent of formalin and standardized to a turbidity of 25,000. malinized, concentrated, stock antigens which had been stored for two years at 10° C. have given satisfactory results in absorption tests and in ordinary agglutination tests. Absorption tests were not done with heat-killed or with living antigens.

If virulent cultures are used in making antigens, great care should be exercised in determining the death of the organisms in the 0.5 per cent formalin suspension, by sterility tests on glucose agar slants, before allowing laboratory workers to use the antigen for agglutination tests or for intravenous injection of rabbits.

Turbidity standard.—Stock antigens suspended in physiologic saline solution containing 0.5 per cent formalin were standardized to a turbidity of 25,000 according to the Standard Methods of Water Analysis, published by the American Public Health Association (1925).

An agglutinin absorption test.—In setting up an absorption test a balance must be observed between the titer of the serum, the amount of the serum used, the amount of bacteria (absorbing dose), and the total volume of the whole mixture. The unit amount of serum used in a test was 0.5 c. c. of a serum having a titer of 1:640; the unit amount of bacteria employed was the bacterial mass contained in

6 c. c. of a suspension having a turbidity of 25,000, and the total volume of the test was 2.5 c. c. In this combination the ingredients are so balanced that the serum, after absorption by the bacterial dose, will no longer agglutinate the absorbing culture. This phase of the test is shown in each test in Tables 2 and 3.

Preserved antiserums having a natural titer of 1:640 were ready for absorption without adjustment of titer, but if the titer was 1:1280 or 1:2560, then an amount of such a serum necessary for the test was diluted at the time of testing with sufficient saline solution to reduce its titer to 1:640. The unit amount of a 640 serum used in a test was 0.5 c. c., but this was diluted 1:5 with saline solution, which gave it a volume of 2.5 c. c.

Stock antigens killed with formalin and standardized to a turbidity of 25,000 were measured out, 6 c. c. to a centrifuge tube, to which was then added about 25 c. c. of saline solution for the purpose of "washing" the bacteria. The bacteria were thrown down in the centrifuge, the clear supernatant fluid was poured off, and to the bacterial sediment were added 2.5 c. c. of a 1:5 dilution of a serum whose titer was 1:640. Thorough mixing of serum and bacterial sediment in the centrifuge tubes was obtained by stirring with a capillary pipette, into which the mixture was alternately sucked and rapidly expelled. The centrifuge tubes were then covered with rubber caps to prevent evaporation and were incubated in a water bath at a temperature of 37° to 42° C. for six hours, after which they were transferred to the cold room at 10° C. overnight. During absorption in the water bath the mixtures were agitated several times. The centrifuge tubes were not calibrated nor was any correction made for saline remaining in the packed bacteria mass, as the error from that source was considered to be not only very small but constant for all tests.

Ordinary agglutination tests were performed the next morning, testing the absorbed serum for agglutination of the type abortus culture, the type melitensis culture, and the culture to be classified as follows: The rubber-capped tubes were centrifuged for about an hour and a quarter. The clear absorbed serum was pipetted off and set up in 0.5 c. c. amounts in agglutination tubes in dilutions of 1:5, 10, 20, 40, 80, and 160, remembering that the absorbed serum was already in dilution of 1:15. To each tube was then added 0.5 c. c. of the formalinized stock antigen, the turbidity of which had been reduced from 25,000 to 500 by adding 1 part of the concentrated antigen to 49 parts of saline solution so that the final turbidity in the agglutination tubes was 250. It was kept in mind that the absorbed serum was a 1:5 dilution of a 640 serum, and therefore the addition of 0.5 c. c. of diluted antigen to 0.5 c. c. of the diluted absorbed serum produced

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a final dilution of serums in the agglutination tubes to 1:10, 20, 40, 80, 160, and 320.

Incubation of the agglutination tubes was at 37° to 42° C. in the water bath for two and one-half hours. They were then placed in the cold room at 10° C. overnight and recorded the next morning. Since the agglutination phase is usually not completed by the next morning it is advisable to allow the test to remain at room temperature for 24 additional hours if complete results are desired. Complete sedimentation of bacteria and water-clear supernatant fluid were indicated by 4. Lesser degrees of clearing were indicated by 3, 2, and 1. All serums in a test were absorbed only once. No sample of serum was reabsorbed.

INTERPRETATION OF ABSORPTION TESTS

By reference to Table 2 one sees the following reactions between type serums 428 and 456 and their type cultures 428 and 456: (1) Either type serum (unabsorbed) agglutinated both type cultures completely or partially in dilution of 1:640. (2) Either type serum after absorption by its homologous culture lost all agglutinins for both type cultures. (3) Either type serum after absorption by the heterologous type culture lost all agglutinins for the heterologous culture but still agglutinated its homologous culture to a considerable degree, thus showing that the two type cultures were serologically different.

Using type melitensis serum 428 and type abortus serum 456 as a basis for classification of the cultures to be studied, the following results were obtained: (1) A culture which absorbed from type serum 428 all agglutinins for type cultures 428 and 456 and for itself was regarded as similar serologically to type culture 428. (2) A culture which absorbed from type serum 456 all agglutinins for type cultures 456 and 428 and for itself was regarded as similar serologically to type culture 456. On this basis of classification the first 10 cultures of Table 2 down to and including culture W. T. H. are similar serologically to Brucella melitensis 428, and the last 10 cultures at the end of Table 2 are similar serologically to Brucella abortus 456.

Culturally, the members of either group of 10 cultures differed among themselves in regard to the carbon dioxide requirement of isolation.

INTERPRETATION OF RECIPROCAL ABSORPTION TESTS

Table 2 presents the results of absorption of agglutinins from type serums by the cultures to be classified. Table 3 presents the absorption of agglutinins by the type cultures from antiserums of the cultures to be classified.

The first 10 cultures of Table 2 are represented in Table 3 by rabbit serums 1, 4, 8, 9, 10, 11, 12, 14, 15, and 17, all of which reacted as

melitensis serums as follows: (1) Each serum (unabsorbed) agglutinated both type cultures and its own homologous culture completely or partially in dilution of 1:640. (2) Each serum after absorption by its homologous culture lost all agglutinins for both type cultures and for its homologous culture. (3) Each serum after absorption by type abortus culture 456 lost all agglutinins for 456 but still agglutinated to a considerable degree type melitensis culture 428 and its own homologous culture, thus showing that it was different from the type abortus serum. (4) Each serum after absorption by type melitensis culture 428 lost all agglutinins for melitensis 428, for abortus 456, and for its own homologous culture, thus showing that each serum was similar to type melitensis serum 428.

The last 10 cultures at the bottom of Table 2 are represented in Table 3 by rabbit serums 18 to 27, all of which reacted as abortus serums as follows: (1) Each serum (unabsorbed) agglutinated both type cultures and its own homologous culture completely or partially in dilution of 1:640. (2) Each serum after absorption by its homologous culture lost all agglutinins for both type cultures and for its homologous culture. (3) Each serum after absorption by type melitensis culture 428 lost all agglutinins for 428 but still agglutinated to a considerable degree type abortus culture 456 and its own homologous culture, thus showing that the serum was different from the type melitensis serum. (4) Each serum after absorption by type abortus culture 456 lost all agglutinins for abortus 456, for melitensis 428, and for its own homologous culture, thus showing that each serum was similar to type abortus serum 456.

Table 3.—Reciprocal absorption of agglutinins from antiserums of cultures to be classified

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Table 3.—Reciprocal absorption of agglutining from antiserums of cultures to be classified—Continued

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Table 3.—Reciprocal absorption of agglutinins from antiserums of cultures to be classified—Continued

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Table 3.—Reciprocal absorption of agglutinins from antiserums of cultures to be classified—Continued

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(19) 88 rabbit serum: Not absorbed	. 4	4	4	4	4	4		•	4	4	4	4	4	4		*	*	4	-	-	7	7	-
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Absorbed by— Abortus 428. Melitensis 428. Culture 88.	044	040	040	040	040	000	$- \!$		000	000	000	000	000			044	040	040	040	040	0-0		
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(20) 633 rabbit seeme	Not absorbed. Absorbed by- Abortus 458. Melitensis 428. Culture 633.	(21) 634 rabbit serim:	Not absorbed Absorbed by— Absorbed by— Abritus 438. Culture 634.	(99) 638 sabbit samen.	Absorbed by Abroria 456	Melitonsis 428. Culture 635.	(92) And well-hit agentum.	Not absorbed by—	Molitensis 428. Culture 429.	(9A) I 7 ashkit commen	Absorbed by—	Abortus 456

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TABLE 3.—Reciprocal absorption of agglutinins from antiserums of cultures to be classified—Continued

					1			-		Agglu	tinst	Jo uo	Agglutination of cultures	8				1					1
Antisorums of cultures to be classified	Ā	rucell	a abo	rtus t	Brucella abortus type culture 466	Iture 4	3		Bruc	Brucella melitensis type culture 428	eliter	sis ty	pe cul	uro 6			ľ	Culture to be classified	to be	class	peg		1
	10	92	9	80 1	160 32	320 640	0 1280	. 08 10	8	\$	8	160	330	\$	128 0821	2	8	3	88	160	828	940	<u>82</u>
			! 						<u> </u>									S	Culture E. M.	. K			
(25) E. M. rabbit serum: Not absorbed	4	4	4	4	4	4		•	-				_	•	•	*	4	7	4	-	-	8	•
Abortus 456 Melitensis 428 Culture E. M.	044	040	040	040	0=0	000			000	000						60460	040	040	040	000	600	$\frac{1}{1}$	
•																		Cult	Culture A.	m.	o.		l
(2c) A. B. C. rabbit serum: Not absorbed		4	4	4	. 4		4	=	<u> </u>	4		4	-		۰		4	*	7	-	-	-	l °
Abortus 466. Melitensis 428. Culture A. B. C.	040	040	040	040	04	000			000	000		000	0	Щ		040	. 4	4	4	4	69	╫	
								·										_ Ö	Culture 41-1	1	-	-	1
(27) 41-1 rabbit serum: Not absorbed	4	4	4	4	4	4		•	4				*	, ,	•	-	4	7	4	-	-	-	-
																		บี	Culture 41-2	41-2			
																4	7	4	+	1	-	7	•
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Absorbed by Abortus 466. Melitensis 428. Culture 41-1.	80	040	040	040	000	000	$\dashv \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$		000		000	000	•••			1 0	040	040	040	040	000		
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			Culture 41-2		

SERUMS OF COW, GUINEA PIG, AND MAN

The rabbit is the animal of choice for the preparation of antiserums for the study of cultures by agglutinin absorption, but in this study serums of cow, guinea pig, and man were available in connection with a few cultures.

Culture McC. was isolated from the milk of a cow by guinea pig inoculation, hence the serums of this cow and guinea pig were absorbed. (See Table 3, serums 2 and 3.) The results were in no way different from those obtained with the McC. rabbit serum; all reacted as melitensis serums.

Culture H-1 was isolated from a guinea pig after inoculation with the cervical seal of a cow whose milk had caused undulant fever in J. J. H. Serums of the man, cow, and guinea pig (see Table 3, serums 5, 6, and 7) reacted after absorption, as did the H-1 rabbit serum; all were *melitensis* serums.

Culture E. F. was isolated from the blood of E. F. directly on culture medium. The human serum E. F., after absorption with *melitensis* type culture 428 failed to agglutinate *melitensis* 428, abortus 456, and its homologous culture E. F., in which respect it reacted, as did the E. F. rabbit serum.

DISSIMILAR CULTURES ISOLATED FROM ONE INDIVIDUAL

(1) Cultures 41-1 and 41-2 (see histories of cultures) were isolated from different portions of the same sample of human blood, the former growing directly in air, the latter requiring carbon dioxide for isolation.

Culture 41-1 was classified as *abortus* by agglutinin absorption and as porcine *abortus* by the bacteriostatic action of dyes, but the absorption test does not distinguish between bovine *abortus* and porcine *abortus*.

Culture 41-2 was classified as melitensis by agglutinin absorption, as Brucella abortus (Bang) by the carbon dioxide requirement of isolation, and as bovine abortus by the bacteriostatic action of dyes.

The isolation from the same blood sample of two cultures differing from each other serologically and in the CO₂ requirement of isolation raised the question as to whether either culture might also contain organisms belonging to the other culture. The purity of both cultures, however, was established by the monovalent character of their respective rabbit antiserums. Neither antiserum after absorption by its homologous culture showed agglutinins for the other culture. (See Table 3, serums 8 and 27.)

(2) Cultures 88 and 89 (see histories of cultures) were isolated from the blood of human case C. G., the former requiring CO₂ for isolation while the latter grew directly in atmospheric air.

Culture 88 was classed as *abortus* by agglutinin absorption and as bovine *abortus* by the bacteriostatic action of dyes. Culture 89 was classed as *melitensis* both by absorption and by the dyes.

Neither culture contained a mixture of both organisms, as shown by the monovalent character of the rabbit serums prepared from each culture. (See Table 3, serums 12 and 19.)

The serum collected August 28, 1930, from the patient C. G., from whom both cultures were isolated June 10, 1930, reacted by agglutinin absorption as a *melitensis* serum. (See Table 3, serum 13.)

CONCLUSION

Chief interest in these studies centers about the serological reactions of certain cultures of *Brucella abortus* (Bang).

A Brucella organism which manifests the cultural character of requiring carbon dioxide for its isolation is Br. abortus (Bang), and yet certain ones of such cultures (McC., H-1, 41-2, H-5, and C-15) are shown by agglutinin absorption to give the melitensis A serological reaction of Brucella melitensis (Bruce). Any contention that agglutinin absorption is a reliable test for the differentiation of Brucella abortus of Bang from Brucella melitensis of Bruce is not supported by these studies.

PRESUMPTIVE DIFFERENTIAL TEST

Abortus and melitensis cultures may be quickly but only tentatively separated serologically by their agglutination in one or the other of the following previously absorbed serums: (1) A type abortus serum which has been absorbed by a type melitensis culture or (2) a type melitensis serum which has been absorbed by a type abortus culture.

The result of such an incomplete test is not to be taken as final evidence but only as suggestive.

HISTORIES OF CULTURES

Culture 428 is the type culture of the group designated Brucella melitensis variety melitensis A by Alice C. Evans (1), who received this strain in 1921 from Feusier and Meyer (2), in whose article it bears the designation of Group III, No. 7.

Culture McC. was isolated November 15, 1930, by Edward Francis, at the National Institute of Health, Washington, D. C., from the spleen of a guinea pig which had been inoculated intraperitoneally October 5 with 2 c. c. of "clear milk" from a cow belonging to Doctor McC., of Chillum, Md. The "clear milk" sample was expressed from the four quarters when the cow was two months dry and two months pregnant. The same organism was also isolated from nine other guinea pigs inoculated October 5 with the same milk sample from the McC. cow. From the 10 guinea pigs 71 cultures were isolated in an atmosphere containing approximately 10 per cent of carbon dioxide, while 71 control culture tubes similarly inoculated from the 10 guinea pigs, but incubated in atmospheric air,

remained sterile. Mrs. McC. became ill from drinking the cow's milk, and her blood serum was reported positive for undulant fever in dilution of 1:320 by the Maryland State Board of Health. Milk from the McC. cow for the inoculation of guinea pigs on October 5, 1930, was kindly furnished by Dr. W. E. Cotton, Director United States Agricultural Experiment Station, Bethesda, Md. For the serological reactions of cow, guinea pig, and rabbit serums, see tests (2), (3), and (1), Table 3.

Culture H-1 was isolated in November, 1929, by Dr. F. P. Mathews, of Purdue University, Lafayette, Ind., from the spleen of a guinea pig which he injected with a portion of the cervical seal of a cow of the H. herd, which was about to calve. Doctor Mathews stated that "the organism was isolated in an atmosphere of carbon dioxide and did not grow in ordinary aerobic conditions." On receipt of the culture, December 6, 1929, at the National Institute of Health, it gave scant growth in atmospheric air but luxuriant growth in air containing 10 per cent CO₂. For the serological reactions of cow, owner (J. J. H.), guinea pig, and rabbit serums, see tests (6), (5), (7), and (4), Table 3. This unusual combination of material bearing on the identity of the H-1 culture is due to the interest and prompt action of Dr. Walter W. Lee (3), then assistant secretary of the Indiana State Board of Health.

Cultures 41-1 and 41-2 were isolated by Dr. I. H. Borts (4) in June, 1929, at Iowa City, Iowa, from blood of a case of undulant fever (WB) which terminated in death in August, 1929. Two blood specimens collected in June, 1929, both agglutinated Brucella melitensis variety abortus in dilution of 1:320. Culture 41-1 was isolated from that portion of a blood sample which was incubated in atmospheric air, while 41-2 was isolated from another portion of the same blood sample but which was incubated in a desiccator from which 10 per cent of the air had been displaced by carbon dioxide. On receipt of these cultures at the National Institute of Health, August 25, 1930, 41-1 grew well in atmospheric air, but 41-2 grew only in air containing 10 per cent carbon dioxide.

Culture H-5 was isolated in an atmosphere of 10 per cent carbon dioxide by Dr. I. F. Huddleson, Michigan State College, East Lansing, Mich., September 20, 1930, from milk of a cow in the M. herd. On receipt of the culture at the National Institute of Health, October 6, 1930, it failed to grow in air but grew luxuriantly in 10 per cent CO₂. In January, 1931, the growth in atmospheric air occurred only at points of heaviest inoculum and not between these points.

Culture B-8 was isolated January 10, 1924, by Dr. F. P. Mathews, Purdue University, Lafayette, Ind., from the stomach contents of a fetus of a cow by direct culture, and Doctor Mathews adds, "presumably by the use of increased CO₂ content, although no record is available in regard to this feature." On receipt of the culture at the National Institute of Health, November 10, 1929, it grew well in atmospheric air.

Cultures C-10 and C-15 were isolated under increased CO₂ content of the atmosphere in April, 1928, by Dr. F. P. Mathews (5), Purdue University, Lafayette, Ind., from two guinea pigs which had been inoculated, respectively, with milk from cow 10 and cow 15 of the Earlham College herd at Richmond, Ind. On June 11, 1928, Doctor Mathews forwarded these cultures to the National Institute of Health with the statement that "C-10 had not yet been trained to grow under aerobic conditions and C-15 had been trained to grow under aerobic conditions only with great difficulty."

Cultures 88 and 89 were isolated by Jordan and Borts (6), June 10, 1930, at Iowa City, Iowa, from the blood of a case of undulant fever (C. G.), a Mexican who had been in the United States but a few weeks. One sample of blood was divided into two portions. One portion, after inoculation into culture medium, was incubated under 10 per cent carbon dioxide tension and yielded culture 88,

which failed to grow in ordinary air. The other portion, after inoculation into culture medium, was incubated in ordinary air and yielded culture 89, which would grow as well in 10 per cent CO₂. On receipt at the National Institute of Health, July 1, 1930, 88 grew luxuriantly in air containing 10 per cent CO₂, but in atmospheric air growth occurred only at points of heaviest inoculum and not between such points. Culture 89 grew luxuriantly in atmospheric air. The patient's blood serum collected August 28, 1930, completely agglutinated abortus 456 and melitensis 428 in dilution of 1:320 and agglutinated melitensis 428 partially in dilution of 1:640.

Culture R. H. W. was isolated April 17, 1928, without the use of CO₂ by W. G. Carhart, pathologist, United States Veterans' Hospital, Whipple, Ariz., from the blood of a patient (R. H. W.) planted on blood agar, growth appearing on the third or fourth day after planting. R. H. W. was a rancher and stated that he probably contracted the infection by delivering aborting goats, since of 1,800 goats due to kid, 400 aborted. The patient's serum, collected May 29, 1928, agglutinated abortus 456, melitensis 428, and his own culture, R. H. W., in dilution of 1:640.

Culture E. F.-1 was isolated from blood drawn November 13, 1928, from E. F. at Washington, D. C., by Surg. W. T. Harrison, who planted the blood clots into flasks of glucose (1 per cent) bouillon (200 c. c.) which were incubated in air at 37° C., and later subcultured to glucose agar slants in air. Onset of illness was November 6, 1928. The source of infection was probably recently isolated cultures with which E. F. was working in the laboratory and which were of melitensis, bovine, and porcine types. The patient's serum collected December 3, 1928, agglutinated abortus 456, melitensis 428, and his own culture, E. F., in dilution of 1:10240. The following cultures were similarly isolated by Doctor Harrison from blood drawn from E. F. on the dates indicated:

- E. F.-2, November 23, 1928.
- E. F.-3, December 3, 1928.
- E. F.-4, December 11, 1928.
- E. F.-5, December 19, 1928.
- E. F.-6, January 4, 1929.

These cultures showed spontaneous sedimentation in 0.85 per cent sodium chloride solution.

Culture W. T. H.-1 was isolated from blood drawn March 18, 1929, from W. T. H. at Washington, D. C., by Medical Director G. W. McCoy, who planted the blood clots into flasks of glucose (1 per cent) bouillon (200 c. c.) which were incubated in air at 37° C. and later subcultured to glucose agar slants in air. Onset of illness was March 10, 1929. The source of infection was probably one of six cultures which W. T. H. had isolated from blood which he drew from E. F. between November 13, 1928, and January 4, 1929, because W. T. H. had made no other contacts with other Brucella cultures at any time. The patient's serum collected March 18, 1929, agglutinated abortus 456 and melitensis 428, in dilution of 1:160. The following cultures were similarly isolated by Doctor McCoy from blood drawn from W. T. H. on the dates indicated:

- W. T. H.-2, April 2, 1929.
- W. T. H.-3, April 15, 1929.
- W. T. H.-4, May 6, 1929.

Culture 456 was isolated in September, 1917, from a cow's fetus at Laurel, Md., and was received from the Bureau of Animal Industry, Department of Agriculture, Washington, D. C. (See Alice C. Evans (1).)

Cultures 633, 634, and 635 were isolated by Doctor Zeller, in Germany, from aborted bovine fetuses in March and April, 1930. When received July 9, 1930,

at the National Institute of Health from Dr. I. F. Huddleson, East Lansing, Mich., the cultures grew well in atmospheric air.

Culture 426 is the type culture of the group designated Brucella melitensis, variety abortus, by Alice C. Evans (1), who received this culture in 1921 from Feusier and Meyer (2), in whose article it bears the designation of group I, No. 20.

Culture L. Z. was isolated by Edward Francis from the blood of L. Z., patient of Dr. Harry V. Paryzek, Cleveland, Ohio. On receipt of the whole-blood sample at the National Institute of Health, Washington, D. C., August 4, 1928, the clot was planted into 200 c. c. of glucose bouillon and incubated in atmospheric air. Subcultures were made on glucose agar slants and incubated in air. The patient's serum received August 4, 1928, agglutinated abortus 456 in dilution of 1:5120 and melitensis 428 in dilution of 1:2560. At the time he became ill the patient was employed in a sausage factory cutting up pork more frequently than he did beef.

Culture E. M. was isolated June 15, 1929, from E. M., a patient at the North Hudson Hospital, Weehawken, N. J. At the time of admission to hospital (April 2, 1929) the patient was employed on the hog feed farms at Secaucus, N. J. The source of the culture was the pus from a "fixation abscess" which had been produced on the abdominal wall of the patient in the region of the appendix. The pus was planted on culture medium June 15, 1929, and was incubated in atmospheric air.

Culture A. B. C. was isolated by Edward Francis, March 10, 1928, from the blood of A. B. C., a patient at the United States Naval Hospital, Washington, D. C. Twenty cubic centimeters of blood collected March 10, 1928, were allowed to clot in four portions of 5 c. c. each. The clots were planted each into 200 c. c. of glucose bouillon and incubated in atmospheric air. Subcultures were made to glucose agar slants and incubated in air. The serum collected March 10, agglutinated cultures 456 and 428 in dilution of 1:160, partial in 1:320. The same organism was isolated from the patient by the same technique on March 17, April 24, and June 1, 1928.

Culture 41-1. (See above.)

Note: While this article was in press, Culture C. S. was isolated from human blood. It adds one more to the list of Brucella abortus (Bang) cultures which by agglutinin absorption give the melitensis A serological reaction of Brucella melitensis (Bruce).

Culture C. S. was one of 12 cultures isolated by Edward Francis, National Institute of Health, Washington, D. C., from the blood of C. S., a patient in the United States Marine Hospital, Detroit, Mich. (Dr. J. H. Linson in charge). Every step in the isolation of the culture was so controlled as to determine whether primary growth was dependent on the presence of carbon dioxide.

Samples of blood received September 15, 18, and 22, 1931, were planted on duplicate sets of culture media, one set being incubated in an atmosphere of 10 per cent carbon dioxide and the other in atmospheric air. Four cultures were isolated from each of the three blood samples, the 12 successful isolations being onmedia incubated in 10 per cent CO₂, whereas there were 12 failures to isolate a culture on the duplicate sets of media incubated in atmospheric air. The patient's serum collected September 2, 1931, agglutinated Brucella abortus 456 in dilution of 1:640 and Brucella melitensis 428 in dilution of 1:1,280. By agglutinin absorption, the patient's serum reacted as a melitensis serum.

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DOUBLE INFECTION BY ORGANISMS OF THE BRUCELLA GROUP

Report of a Case

By Carl F. Jordan, M. D., Assistant Professor of Hygiene and Preventive Medicine, State University of Iowa, Iowa City, Iowa; and I. H. Borts, M. D., Chief Bacteriologist, State Hygienic Laboratories, Iowa City, Iowa

In the literature on undulant or Malta fever there are numerous case reports. With the exception of one case reported by Hardy, Jordan, Borts, and Hardy (1), all have been regarded as infections by a single variety or species of the *Brucella* group. We present here a report of another patient with undulant fever, from whose blood culture two varieties (abortus and melitensis) of Brucella melitensis were isolated.

CASE REPORT

C. G., male, age 30, Mexican laborer, admitted to the Santa Fe Hospital, Fort Madison, Iowa, on May 9, 1930.

History.—For one week previous patient had complained of weakness, general aching, headache, and feverishness. The signs of jaundice had also developed. There were marked anorexia and occasional vomiting. Owing to language difficulties the attending physician was not able to obtain the details of the history.

The past medical history was essentially negative.

Physical findings.—

General: Temperature 104.6°; pulse, 80; respiration, 24; blood pressure, 108/50. Anemic in appearance, weak, and poorly nourished.

Head: Eyes, ears, nose, and throat essentially negative.

Neck: There was tenderness over the anterior cervical glands.

Lungs: Clear.

Heart: No enlargement, rhythm regular, a faint systolic murmur was heard at apex and base.

Abdomen: Moderate tenderness in epigastrium just above the umbilicus. The liver and spleen were not felt.

Skin: More vellow than normal.

Extremities: Normal. Reflexes: Normal.

Provisional diagnosis.—Typhoid, typhus, or undulant fever.

Laboratory findings.—

Urine: Free from albumin, casts, and sugar.

Blood: R. B. C. 3,800,000; W. B. C. 4,500; Hb. 70 per cent.

Agglutination tests on blood serum received at the State laboratory May 14, 1930, showed no reaction for typhoid fever, but Br. melitensis was agglutinated in dilutions to 1:320. On July 2, the agglutination titer was 1:1280.

A blood culture, consisting of 100 c. c. of fresh beef liver infusion broth containing 3-5 c. c. of the patient's blood, received by us May 23, was incubated under atmospheric conditions, subcultures being made May 27, May 31, and June 3, 1930, to beef liver infusion agar pH 6.6 in Petri plates in duplicate sets. One set was incubated under atmospheric conditions and the other under an atmosphere of 10 per cent carbon dioxide tension. Growth appeared June 10 on a plate incubated under atmospheric conditions and also on one under an atmosphere of 10 per cent carbon dioxide tension. The organisms on both plates were Gram negative, resembled *Brucella* morphologically, and were agglutinated by specific immune serum.

Several stool specimens cultured by the Amoss method (2) did not reveal organisms of the *Brucella* group.

Identification of strains.—In view of the patient's recent entrance from Mexico into the United States and his contact with goats, we had entertained the possibility of his suffering from a *Brucella* infection of the *melitensis* variety. For this reason one of us (I. H. B.) attended to the technical work involved.

When cultured according to Huddleson's dye method (3), the organism which grew under atmospheric conditions proved to be *Brucella melitensis*, variety *melitensis* (Lab. strain No. 89); that isolated under 10 per cent carbon dioxide tension, *Brucella melitensis*, variety *abortus* (Lab. strain No. 88).

These strains were forwarded to Dr. Edward Francis, of the National Institute of Health, at Washington, D. C., and to Dr. I. Forest Huddleson, Michigan State Agricultural College, East Lansing, Michigan. Both Francis (4) and Huddleson corroborated our findings. They used the agglutinin absorption and the dye test, respectively. From a second blood culture taken June 28, but one organism was isolated, namely, *Brucella melitensis*, variety melitensis.

Animal inoculations.—In order to determine the pathogenicity of the organisms isolated from the blood culture of this case, one-fourth 2439 October 9, 1931

c. c. of a dilute suspension (1:100 dilution of a suspension having a turbidity corresponding to 500 parts per million by the silica standard) was made from 72-hour cultures of laboratory strains Nos. 88 and 89 and inoculated into each of two guinea pigs. Prior to inoculation all of the guinea pig sera failed to agglutinate our standard Brucella antigen. After six weeks these pigs were again bled, the serum in each instance agglutinating Br. abortus antigen, in dilutions ranging from 1:320 to 1:5,120.

One of the pigs inoculated with Culture No. 89 was autopsied after eight weeks. The spleen was three times the normal size, contained several pin-head sized abscesses, and was bound down to the posterior wall by dense fibrous adhesions. The liver was somewhat enlarged. and diffusely studded with pin-point abscesses. Lymph glands in the groin and axilla, and of the mesenteric, iliac, and bronchial groups were enlarged to the size of a large navy bean. The glands were firm and on section contained thick creamy pus. The costo-sternal articulations were markedly arthrosed, being involved in a dense mass of fibrous tissue. The lungs contained many small pea-sized abscesses. The serum of blood taken at autopsy agglutinated Brucella antigen in dilutions through 1:5,120. Brucella melitensis, variety melitensis, was isolated from all organs. The mate of this pig was posted with similar findings with the exception that the costosternal articulations were not involved and the serum titer was 1:2,560. It is interesting to note that these findings closely simulate the pathology found in guinea pigs experimentally inoculated with freshly isolated strains of the suis variety of Br. melitensis.

One of the pigs inoculated with Culture No. 88 showed several pinpoint abscesses in the liver. The spleen was twice the normal size, and no abscesses could be demonstrated. There was no evidence of lymph gland involvement. The remaining organs were apparently normal. The blood serum agglutinated *Brucella* antigen through 1:640. *Br. melitensis*, variety abortus, was isolated from the spleen and liver. The mate of this pig presented identical findings.

It is quite evident from the pathological standpoint as well as from the agglutinin absorption tests as carried out by Francis and the dye method of Huddleson that we were dealing with two distinct varieties of organisms of the *Brucella* group.

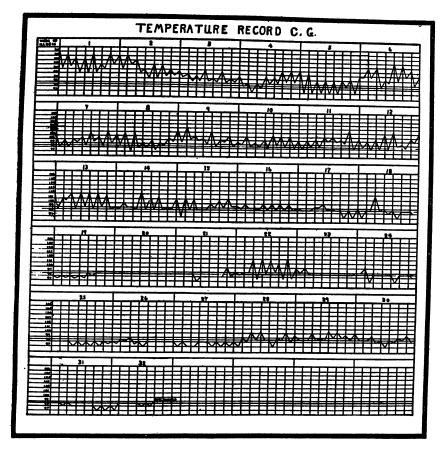
Epidemiological data.—On June 28 the patient was seen in the Santa Fe Hospital, an American woman who had lived for years among Mexicans ably assisting as interpreter. Remarks were at times incoherent, due to the feverish state.

The patient's home was in Juriria, Guanajuato, Mexico. He had made several trips to the United States from Mexico, between the years 1924 and 1930. While in Mexico, he did farm work, and had eight goats, which were usually milked by his mother and sisters.

He did not drink goat's milk, but ate cheese made from this milk. It was impossible to elicit accurate information relative to infectious abortion in animals.

The patient drank cow's milk at times. He had one cow which was sold to enable him to come to the United States.

He left Mexico in February, 1930, along with other Mexicans and a labor agent, going through Laredo to Fort Worth, from there to Kansas City, and finally to Sibley, Mo. He arrived at Sibley about



a month before admission to the hospital and about three weeks before the onset of his illness. While there, he had no contact with livestock, used only canned milk, and ate no butter—Mexicans, it was stated, eat lard instead of butter.

Course and treatment.—The patient's temperature (see temperature chart) was 106° on May 9, the day of admission. He continued to run a septic type of temperature, ranging from 98° to 99° in the morning to 101° and 103° every afternoon until September 1. Vaccine therapy was instituted July 2 and continued until July 9. The patient's condition improved slightly, but there was no appreciable

change in the temperature. On August 31, 0.1 gram acriflavine was given intravenously, and on September 1, 0.2 gram. The temperature became normal and remained so until September 6, when it again rose to 101.8°. On September 7, 0.4 gram of acriflavine was administered, following which the temperature again became normal and remained so several weeks.

The patient was "weak and emaciated" on September 24, but "gaining ground slowly". During the period October 1 to 8 another exacerbation of fever occurred, the temperature reaching 102° on the 5th and 6th, and then returning to normal. On November 12, patient was "gaining rapidly in weight and general appearance." He was granted a pass to Mexico and left the hospital December 17, 1930.

COMMENT

The following evidence indicates that the melitensis variety of Br. melitensis was in all likelihood acquired in Mexico.

(1) Undulant fever cases due to *Br. melitensis* occurred in Texas and were described in 1911 by Ferenbaugh (5) and by Gentry and Ferenbaugh (6). Cases occurring in Arizona were reported by Yount and Looney (7), those of 1922 by Watkins and Lake (8), and those in Southwestern United States by Lake (9). Goats were regarded as the source of infection.

A case of undulant fever with isolation of *Br. melitensis*, variety *melitensis*, from blood and urine was reported in 1918 by Woolsey (10). The patient was a Mexican and apparently developed the infection in Mexico.

- (2) From 69 cases of undulant fever in Iowa in which blood cultures were positive, 71 organisms have been isolated. Forty-five strains were of variety suis, 25 of variety abortus, while the only melitensis variety is that isolated from the case here reported. Variety melitensis infection is not endemic in Iowa.
- (3) Br. melitensis variety melitensis infection is known to be endemic in Mexico.

Letters were directed June 30, 1930, to health officers of Tampico, Mexico City, and Vera Cruz requesting information relative to the incidence of undulant fever and infectious abortion in goats and cows. Dr. E. Garcia, writing from Tampico, July 7, stated that no cases of undulant fever had been reported, but that infectious abortion occurred in goats and frequently in cows. Undulant fever cases were reported in the vicinity of Mexico City by Dr. E. Lando, officer in charge of the bureau of interchange.

Dr. Miguel E. Bustemante, director of the sanitary bureau of Vera Cruz, in a letter dated August 5, 1930, referred to undulant fever studies carried out in 1919-1920 by Dr. Ferando Ocaranza, director of the medical faculty at the National University of Mexico, and Dr. Gerar de Varela, bacteriologist of the Institute of Hygiene, Poptla

Tacuba. Doctor Bustemante stated that undulant fever cases were for the most part confined to the central plateau of Mexico, in the States of Pueblo, Mexico, Guanajuato, and San Luis Potosi.

SUMMARY

A Mexican laborer, aged 30, left his native country in February, 1930, took sick during April in Missouri, and was treated for undulant fever in a hospital in Iowa for 32 weeks. The blood culture yielded two strains of Brucella melitensis—variety melitensis and variety abortus. The melitensis variety of Brucella infection was in all likelihood acquired in Mexico because (1) with this one exception, all of the undulant fever cases in Iowa have, so far as known, been due to Brucella melitensis, variety abortus or variety suis, variety melitensis not being endemic in Iowa; (2) Brucella melitensis, variety melitensis infection is known to be endemic in Mexico; (3) the patient had contact with and used dairy products from goats in Mexico but not in the United States.

The source of the abortus variety of organism is not so clear. A double infection may have developed before the patient left Mexico, as he used milk in addition to caprine dairy products. On the other hand, it is possible that the bovine infection was superimposed after the patient's arrival in Iowa. Pasteurized milk was used, but several cases of undulant fever are known to have occurred in the same community, with dairy products as the probable source of infection, one other case occurring within the same period.

ACKNOWLEDGMENTS

We desire to acknowledge the collaboration of Dr. F. D. Ullrich, of Fort Madison, Iowa, who supervised the care of the patient, forwarded blood specimens, and made available the clinical and in part the epidemiological data relative to this case. We wish also to acknowledge the services rendered by Doctors Francis and Huddleson in corroborating our findings as to identification of the organisms isolated from the blood culture.

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(9) Lake, G. C.: Malta fever in Southwestern United States. Pub. Health Rep., 37: 2895-2899 (Nov. 24), 1922.

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COURT DECISION RELATING TO PUBLIC HEALTH

Conviction for unlawful possession of plants known as "marajuana" upheld.—(Louisiana Supreme Court; State v. Bonoa, 136 So. 15; decided May 25, 1931.) Act 41 of 1924 provided in section 1 as follows:

That no person shall possess, sell, dispose of, transport, deliver, in any form whatever in the State of Louisiana, the plant known as marajuana or any of its derivatives, either dried or in the form of cigarettes, tobacco, or any other way whatsoever.

Violation was made a misdemeanor, punishable by fine and imprisonment. The defendant was charged with unlawfully possessing plants known as marajuana, in that he had a number of the plants growing on his premises. These plants were growing in what was termed a second back yard immediately in the rear of the first. The two yards were separated by a shed through which one had to go to enter the second yard from the first. The defendant was convicted and appealed to the supreme court.

One of the defendant's contentions was that section 1 of the act involved was unconstitutional and void in so far as it attempted to prohibit the possession of plants termed "marajuana," as the section by so doing sought to prohibit the possession of something unknown. It was urged that there was no such plant known as marajuana and that hence the terminology used conveyed no conception of what was Concerning this the supreme court stated that it did not prohibited. find any difficulty in holding that the use of the word "marajuana" in connection with the word "plant" conveyed to the mind exactly what the legislature intended to convey, namely, the plant scientifically known as Cannabis indica or Cannabis americana. sides," said the court, "whatever doubt there may be as to what was meant by the use of the word is removed not only by the title of the act, where the plant is referred to as 'Cannabis indica, Cannabis america, or marajuana,' but also by section 4 of the act, where the plant is similarly designated, the name 'Cannabis indica' being well known scientifically."

Another ground urged by the defendant against the validity of section 1 was that, in so far as it prohibited the possession of the

marajuana plant in any form whatsoever, it was an infringement upon liberty and the rights to property in violation of the State and Federal constitutions. The court stated the defendant's views in this respect as follows:

The theory of the accused seems to be that, although the marajuana plant may be used in forms, such as cigarettes or tobacco, injurious to the public health, morals, and safety, yet it may be used for valuable purposes, such as the manufacture of hemp rope and twine, in the preparation of useful drugs, and for the production of seed which forms a large part of the rations of the millions of pet canary birds in this country, and that only in so far as the plant is sold, used, and possessed for deleterious purposes may such sale, use, or possession be prohibited without infringing, in violation of the State and Federal constitutions, upon the liberty of the people.

The court's holding with regard to this contention was adverse to the defendant, the view being taken that the legislature had not exceeded its powers by enacting section 1 of the act. In disposing of this point the court stated, in part, as follows:

One who has upon his premises to his knowledge a growing crop of Cannabis indica or Cannabis americana or marajuana, or any number of the plants growing thereon, possesses these plants within the meaning of section 1 of the statute. * * *

The act was passed under the police power of the State. In State v. Mc-Cormick (142 La. 580, 77 So. 288, 289, L. R. A. 1918C, 262) it was said: "The legitimate exercise of the police power is not subject to restraint by constitutional provisions for the general protection of rights of individual life, liberty, and property.' State v. Schlemmer (42 La. Ann. 1166, 8 So. 307, 10 L. R. A. 135). And the fourteenth amendment to the Constitution of the United States does not interfere with the proper exercise of that power. (6 R. C. L. pars. 193, 194; L'Hote v. New Orleans, 177 U. S. 596, 20 S. Ct. 788, 44 L. Ed. 903.)"

The marajuana plant is a plant possessing properties deleterious to health and dangerous to the public safety and morals. * * * To permit the plant to be possessed in the State, even in its growing form, is virtually as unsafe as to permit its possession in the manufactured forms of cigarettes and tobacco, so readily and easily may it be converted into those forms.

The marajuana plant is not one of the crops of this State. While the plant may be put to valuable uses, nevertheless its deleterious properties may be fairly considered as outweighing those uses.

DEATHS DURING WEEK ENDED SEPTEMBER 19, 1931

Summary of information received by telegraph from industrial insurance companies for the week ended September 19, 1931; and corresponding week of 1930. (From the Weekly Health Index, issued by the Bureau of the Census, Department of Commerce)

Policies in force	Week ended Sept. 19, 1931 74, 883, 159	Corresponding week, 1930 75, 532, 011
Number of death claims		13, 466
Death claims per 1,000 policies in force, annual rate.		9. 3
Death claims per 1,000 policies, first 38 weeks of		
year, annual rate	9. 9	9. 7

Deaths¹ from all causes in certain large cities of the United States during the week ended September 19, 1931, infant mortality, annual death rate, and comparison with corresponding week of 1930. (From the Weekly Health Index, issued by the Bureau of the Census, Department of Commerce)

[The rates published in this summary are based upon midyear population estimates derived from the 1930 census]

1990 General									
	Wee	ek ended	Sept. 19	, 1931	Corresponding Death rate the first weeks				
City	Total deaths	Death rate ²	Deaths under 1 year	Infant mor- tality rate 3	Death rate ¹	Deaths under 1 year	1931	1999	
Total (82 cities)	7, 453	10.9	732	4 57	10.3	728	12. 2	12. 1	
Akron	30 32 60 37	6. 1 12. 9 11. 3	5 2 15 8	49 40 153 127	7. 1 11. 4 15. 2	3 2 10 8	7. 9 13. 9 15. 2	7. 9 15. 0 16. 9	
Colored Baltimore 5 White	23 206 152	(⁵) 13. 2	8 7 22 14	201 75 61	(9) 14. 5	17 17 11	(9) 14. 6	(⁶) 14. 1	
ColoredBirminghamWhite	54 53 29	(6) 10. 3	8 1 1	125 10 17	(6) 11. 2	6 7 3	(9) 13.8	(⁶) 13. 9	
Colored Boston Bridgeport Bridgelo	24 202 34 129	(9) 13. 4 12. 1 11. 6	0 20 1 20	0 57 17 82	(6) 13. 2 7. 8 11. 3	4 25 0 19	(9) 14. 3 11. 3 13. 3	(6) 14. 3 11. 2 13. 1	
Buffalo. Cambridge. Camden Canton. Chicago 3	17 25 16	7.8 11.0 7.8	0 1 2	0 17 4 6	11.0 10.5 6.4	3 3 4	12. 3 14. 5 10. 3	11.8 13.7 10.1	
Cleveland Columbus	632 147 198 79	9. 5 16. 8 11. 3 13. 9	54 22 15 4	48 132 44 39	9.9 13.1 10.0 13.4	52 8 19 5	10.9 16.2 11.4 13.9	10.5 15.7 11.3 15.8 11.7	
Dallas White Colored Dayton	41 29 12 52	7. 9 (6) 13. 1	6 5 1 8	112	7.9 (6) 12.4	7 6 1 10	(°) 11. 9	11. 7 	
Denver Des Moines Detroit	64 24 229	11. 4 8. 7 7. 2	5 3 32	48 53 51	12.1 6.9 8.5	12 3 44	14.1 11.2 8.4	14.9 11.9 9.5	
Duluth. El Paso Erie Fall River 5; Flint	36 24 24 21	18. 4 11. 9 10. 6 9. 5	3 4 1 3	74 19 68	10.3 10.1 12.6 11.3	1 7 2 0	11. 4 16. 2 10. 8 11. 4	11, 2 17, 7 11, 4 12, 1	
Flint Fort Worth White Colored Grand Rapids	21 31 28	6. 7 9. 7	8 1 0	102	7. 6 11. 4	5 6 4	7. 1 11. 0	9. 3 11. 2	
Colored Grand Rapids Houston White	3 28 83 55	(6) 8. 5 14. 0	1 4 7 5	59	(6) 10. 5 12. 5	2 3 10 2	9. 2 11. 3	10. 4 12. 2	
Colored	28 98 79	(6) 13. 8	2 5 4	41 38	(6) 11. 7	8 10 10	(6) 14. 1	14.8	
Kansas City, Kans	19 75 30 26	(6) 12. 3 12. 7	1 6 1	67 53 21 25	(6) 11. 5 11. 1	0 7 4 4	(6) 11. 7 12. 7	(5) 11. 4 11. 6	
Colored Kansas City, Mo Knovville White Colored Long Beach	97 27	(6) 12. 4 12. 9	0 6 7 7	0 46 149	(¢) 10. 3 11. 3	0 6 4 3	(6) 13. 3 12. 5	(6) 13. 4 14. 0	
Colored Long Beach Los Angeles	22 5 28 252	9. 6 10. 0	0 0 15	167 0 0 44	(°) 6. 9 10. 5	1 0 24	(6) 9. 9 10. 8	(⁶) 9. 9 11. 1	
Louisville	70 51 19	11. 8 (6) 9. 3	13 10 3	98 199	8. 8 (6) 7. 8	12 8 4	14. 5 (6) 12. 7	13. 7 (6) 13. 5	
Lowell 7. Lynn Memphis White	18 14 102 42	9. 3 7. 1 20. 6	1 0 8 7	25 0 85 117	7. 8 9. 2 10. 9	2 3 5 1	9. 7 16. 8	10. 6 17. 6	
Colored Miami White Colored	60 20 13	9. 3	1 3 2	29 76 71	(6) 5. 6	2 1 -	(6) 11. 9	11. 2	
Colored	7	(6)	1 1	88	(6)	1	(6)	(6)	

See footnotes at end of table.

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Deaths1 from all causes in certain large cities of the United States during the week ended September 19, 1931, infant mortality, annual death rate, and comparison with corresponding week of 1930. (From the Weekly Health Index, issued by the Bureau of the Census, Department of Commerce)—Continued

	Wee	k ended	Sept. 19,	, 1931	Corresponding week, 1930 Death ra			irst 38
City	Total deaths	Death rate 3	Deaths under 1 year	Infant mor- tality rate 3	Death rate ¹	Deaths under 1 year	1931	1930
Milwaukee Minneapolis	97 105	8. 6 11. 6	14 14	61 90	8. 9 11. 2	14 10	9. 5 11. 5	9. 7 10. 7
Nashville White	57 39	19. 1	13 10	194 199	9. 1	6	17. 1	16.7
Colored	18	(6)	3	177	(6)	5 1	(6)	(6)
New Bedford 7 New Haven	23 31	10.7 9.9	3	80	6.0 12.2	2 2	12.2	10. 9
New Orleans	130	14. 5	14	77	13. 6	10	12.6 17.2	13. 1 17. 5
WhiteColored	78 52		5 9	41		4		
New York	1, 343	(º) 9. 9	121	147 51	(6) 8.8	6 123	(6) 11. 4	(6) 11. 0
Bronx Borough	190 460	7. 4 9. 1	12	27	6.4	10	8.4	8.0
Brooklyn Borough Manhattan Borough	508	14.6	48 41	51 70	8. 5 12. 6	56 42	10. 5 17. 3	10.0 16.3
Uneens Rorongh I	140	6.3	16	44	5. 5	13	7.4	7. 1
Richmond Borough Newark, N. J	45 94	14. 4 11. 0	11	72 58	9. 2 10. 4	6	14. 1 11. 8	14. 5 12. 2
Oakland	52	9.3	5	64	9. 1	5	10.5	11.0
Oklahoma City Omaha	35 52	9.3	4 3	55 34	12. 5 13. 6	7 2	11. 1 14. 1	10. 8 13. 8
Paterson	32	12.5 12.0	4	69	10.5	3	13. 5	12.4
Peoria Philadelphia	18 442	8.7 11.7	3 54	79 78	14.8	48	12. 7 13. 3	12.6 12.8
Pittsburgh	160	12.3	25	86	12.3	25	14.8	13.9
Portland, OregProvidence	62 57	10. 5 11. 7	3 6	36 55	9. 5 11. 1	10	11.7 12.9	12. 2 13. 2
Richmond	54	15. 3	6	87	13.4	5	15.8	15. 2 15. 1
White Colored	32 22		1 5	22 217		2 3		
Rochester	82	(6) 12. 9	5 7	64	(6) 10. 1	4	(6) 12. 1	(6) 11. 6
St. Louis	201 44	12. 7 8. 3	19	64	8. 6 8. 0	13	15.6	14.4
Balt Lake City	35	12.8	2	30	10.4	1 5	10.9 12.3	10. 2 12. 5
lan Antonio	52 41	11. 3 13. 7	12		13. 2	7	14.8	17. 1
an Francisco	146	11.7	6	61 40	12. 2 11. 9	1 7	13. 7 13. 2	14. 5 13. 1
chenectady	13 78	7. 0 10. 9	0	0	13. 1	0	10.6	11.4
omerville	15	7.4	8	81	7. 4 8. 5	2	11. 5 9. 1	10. 9 9. 9
outh Bendpokane	15 19	7. 2 8. 5	2	50	8.4	1	8.1	9. 0
pringfield, Mass	24	8.2	1 5	26 77	9.9 10.7	0 2	12. 4 11. 9	12. 4 12. 2
yracuse 'acoma	46 26	11.3	3	36	9. 2	4	11.8	11.7
'oledo	56	12. 6 9. 9	0	0 55	8.3 12.0	1 12	12.1 12.0	12.6 12.7
renton	37	15.6	2	35	13. 5	1	16.8	16.8
Itica Vashington, D. C White	20 141	10. 2 14. 9	12	52 66	11. 3 12. 0	1 8	14. 1 16. 0	15. 0 15. 2
White	96 _		8	65 -		6 -		
Colored	45 21	(6) 10. 9	4	69 30	(6) 5. 2	2 2 3 2 2	9.8	(6) 9. 9
Vilmington, Del.	36	17.6	5	108	13. 7	3	14. 2	14. 4
Vorcesteronkers	42 18	11. 1 6. 8	3 3	41 79	11. 2 6. 5	2	12.3	13.0
oungstown	32	9.7	i	14	11.9	2	8. 7 10. 3	8. 1 10. 3

¹ Deaths of nonresidents are included. Stillbirths are excluded.

These rates represent annual rates per 1,000 population, as estimated for 1931 and 1930 by the arithmetical method.

Deaths under 1 year of age per 1,000 live births. Cities left blank are not in the registration area for births.

⁴ Data for 77 cities.

Data for 77 cutes.
 Deaths for week ended Friday.
 For the cities for which deaths are shown by color, the percentage of colored population in 1920 was as follows: Atlanta, 31; Baltimore, 15; Birmingham, 39; Dallas, 15; Fort Worth, 14; Houston, 25; Indianapolis, 11; Kansas City, Kans., 14; Knoxville, 15; Louisville, 17; Memphis, 33; Miami, 31; Nashville, 30; New Orleans, 26; Richmond, 32; and Washington, D. C., 25.
 Paralletica Apr. 1, 1020; decreased 1020 to 1930, no estimate made.

PREVALENCE OF DISEASE

No health department, State or local, can effectively prevent or control disease without knowledge of when, where, and under what conditions cases are occurring

UNITED STATES

CURRENT WEEKLY STATE REPORTS

These reports are preliminary, and the figures are subject to change when later returns are received by the State health officers

Reports for Weeks Ended September 26, 1931, and September 27, 1930

Cases of certain communicable diseases reported by telegraph by State health officers for weeks ended September 26, 1931, and September 27, 1930

	Diph	theria	Influ	ienza	Ме	asles		gococcus ngitis
Division and State	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930						
New England States: Maine	30 1 3	1 2 1 50 6 7	4	1	15 1 1 18 6 5	1 50	1 0 0 4 0 2	0 0 0 3 0
Middle Atlantic States: New York New Jersey Pennsylvania East North Central States:	55 15 68	54 46 125	16	1 2 2	55 10 91	45 16 51	5 4 6	7 1 6
Ohio	80 19 63 20 10	63 8 102 44 11	10 6 266	11 5 3	28 12 85 13 8	21 2 20 18 21	1 1 11 8 0	9 3 5 3 1
West North Central States: Minnesota Iowa Missouri North Dakota South Dakota	10 9 55 2	13 5 27 1 8	1		2 2 1	1 2 13 8 9	0 2 0 4 0	0 0 4 3 1
Nebraska Kansas South Atlantic States: Delaware Maryland 11.	14 6 2 40	5 7	8 2	1 4	6 2 1	1 4 3	0 0 0 1 1	0 1 0 0
District of Columbia	11 28 129 28 56	15 21 118 38 21	12 1 113 6	1 14 160 15	5 4 7 2	13 10 5 4	0	0 3 0 0
East South Central States: Kentucky Tennessee. Alabama 1. Mississippi.	147 74 95 112	18 30 23	2 1	5 5	10 1	12 16	0 4 1	1 0 2 0

¹ New York City only.
2 Week ended Friday.
3 Typhus fever, 1931, 10 cases: 1 case in Maryland; 3 cases in Georgia; 3 cases in Florida; 2 cases in Alabama; and 1 case in Texas.

Cases of certain communicable diseases reported by telegraph by State health officers for weeks ended September 26, 1931, and September 27, 1930—Continued

	Diph	theria	Influ	lenza	Me	asles	Menin men	gococcus ingitis
Division and State	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930
West South Central States: Arkansas. Louisiana Oklahoma 4	28 44 76	7 43 35	12	2	2 3	3 2	0 2 0	0 0 1 0
Mountain States:	22	15	2	2	3	2	ŏ	
Idaho Wyoming Colorado New Mexico Arizona	10 2 4	2 1 10 3 6	3		1 6 1	7	0 0 0 0	0 0 2 0 1 2
Utah ² Pacific States: Washington Oregon California	7 1 56	3 5 39	5 15 23	17 39	8 7 37	2 4 16 56	1 0 0 6	2 2 0 4
				Smal	llpox	Typhoi	d fever	
Division and State	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930
New England States: Maine. New Hampshire. Vermont. Massachusotts. Rhode Island. Connecticut.	7 2 4 105 8	21 1 0 32 2 5	6 0 3 75 9	12 2 3 72 3 14	0 0 0 0	0 0 0 0 0	4 3 0 7 1 5	9 9 1 11 11 2
Middle Atlantic States: New York New Jersey Pennsylvania East North Central States:	327 93 49	65 6 10	124 35 143	71 47 138	0	8 0 0	54 16 59	54 21 88
Ohio Indiana Illinois Michigan Wisconsin	14 3 62 138 70	100 6 43 13 20	139 18 110 60 21	184 48 142 101 41	1 3 11 1 4	19 12 12 1 1 6	183 16 53 22 8	75 12 43 24 5
West North Central States: Minnesota Iowa Missouri North Dakota South Dakota Nebraska Kansas	62 9 0 2 1 1	17 21 18 2 4 26 48	39 13 21 10 1 4 20	32 18 36 11 4 12 35	3 8 5 1 1 0 3	4 7 8 1 1 10	19 8 18 5 5 1	7 2 30 4 1 7
Outh Atlantic States: Delaware Maryland 12 District of Columbia West Virginia North Carolina South Carolina Georgia 2 Florida 2	0 5 2 3 5 0 4 1	0 2 0 3 5 2 1	1 33 9 24 75 18 25	4 11 3 19 96 21 16 4	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 7 1 0 0	1 53 3 68 41 49 33 12	3 51 2 53 40 35 35 6
East South Central States: Kentucky. Tennessee Alabama ² Mississippi.	2 7 1 2	1 2 1 2	31 65 28 21	16 30 45 12	0 3 1 3	0 2 0 0	61 82 31 27	54 42 21 35

Week ended Friday.
 Typhus fever, 1931, 10 cases: 1 case in Maryland; 3 cases in Georgia; 3 cases in Florida; 2 cases in Alabama; and 1 case in Texas.
 Exclusive of Oklahoma City and Tulsa.

Cases of certain communicable diseases reported by telegraph by State health officers for weeks ended September 26, 1931, and September 27, 1930—Continued

	Polion	ayelitis	Scarle	t fover	Sma	mallpox Typi		hoid fever	
Division and State	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	Week ended Sept. 26, 1931	Week ended Sept. 27, 1930	
West South Central States: Arkansas. Louisiana Oklahoma 4 Texas 1 Mountain States: Montana. Idaho. Wyoming. Colorado. New Mexico. Arizona. Utah 1 Pacific States: Washington. Oregon. California	0 2 0 1	1 11 9 8 0 1 7 4 2 1 2 3 1 65	14 11 24 30 9 2 5 5 11 2 5 3 3	4 8 34 111 15 1 7 7 9 8 5 5 3 31 13 559	1 0 1 1 1 2 0 0 0 0 0 0 6 6 6	0 1 14 4 0 0 0 0 0 0 0 0	15 56 49 20 10 5 1 7 5 3 2	15 27 59 7 15 0 0 11 20 6 2	

SUMMARY OF MONTHLY REPORTS FROM STATES

The following summary of cases reported monthly by States is published weekly and covers only those States from which reports are received during the current week.

State	Menin- gococ- cus menin- gitis	Diph- theria	Influ- enza	Ma- laria	Mea- sles	Pel- lagra	Polio- myelitis	Scarlet fever	Small- pox	Ty- phoid fever
August, 1951 California	21 17 17 1 37 4	148 12 9 179 69 233 97 19 186 29	49 4 9 17 67 31	7 66 64 62 13 262 8 2 2 3, 424	197 5 6 218 6 793 4 29 413 18	8 1 30 89	22 0 1 116 0 2,638 3 1 34 34	145 6 18 242 46 393 40 20 327	28 1 1 37 6 6 15 35 0	83 13 8 106 245 187 199 21 177
TexasVirginiaWashington	1 4 4	80 123 18	23 505 18	947 91	101 32	9 56	8 13 14	77 125 45	3 47	123 254 29

¹ Exclusive of Oklahoma City and Tulsa.

August, 1931	Cases	Chicken pox—Continued.	Cases
Anthrax:		Oklahoma ¹	9
California	. 1	Oregon	38
Illinois	. 1	Pennsylvania	138
New York	2	Porto Rico	1
Pennsylvania	2	Virginia	84
Chicken pox:		Washington	47
California	129	Diarrhea and dysentery:	
Florida	3	Virginia	1, 691
Idaho	4	Dysentery:	
Illinois	66	California (amebic)	6
Louisiana	5	California (bacillary)	11
New York	1	Florida	1

¹ Exclusive of Oklahoma City and Tulsa.

² Week ended Friday.

³ Typhus fever, 1931, 10 cases: 1 case in Maryland; 3 cases in Georgia; 3 cases in Florida; 2 cases in Alabama; and 1 case in Texas.

⁴ Figures for 1931 are exclusive of Oklahoma City and Tulsa.

Dysentery—Continued.	Cases	Paratyphoid fever—Continued.	• •
Illinois		Texas.	5
Illinois (amebic)		Washington	1
Illinois (bacillary)	7	Puerperal septicemia:	
Louisiana	13	New York	13
New York	21	Pennsylvania	22
Oklahoma 1	41	Washington	1
Porto Rico	17	Rabies in animals:	
Filariasis:		California	36
Porto Rico	3	Louisiana.	4
Food poisoning:		New York 2	3
California	19	Oregon	1
German measles:		Relapsing fever:	-
California	24	California	4
Illinois	9	Rocky Mountain spotted or tick fever:	*
	42	• • • • • • • • • • • • • • • • • • • •	
New York		Oregon	2
Pennsylvania	14	Scables:	
Washington	15	Oklahoma 1	1
Granuloma, coccidioidal:		Oregon	2
California	1	Septic sore throat:	
Hookworm disease:		California	4
California	2	Illinois	3
Louisiana	11	Louisiana	1
Impetigo contagiosa:		New York	11
Oregon	5	Oklahoma 1	22
Washington	3	Oregon	3
Lead poisoning:		Tetanus:	•
Illinois	4	Illinois	10
Leprosy:			13
California	1	Louisiana	3
Louisiana	1	New York	11
Porto Rico	2	Oklahoma 1	2
Lethargic encephalitis:		Pennsylvania	6
California	2	Porto Rico	3
Louisiana	ī	Tetanus, infantile:	
New York	12	Porto Rico	5
Oregon	4	Trachoma:	
Pennsylvania	9	California	7
Washington	5	Illinois	11
	9	Oklahoma ¹	10
Mumps:	140	Porto Rico	2
California	149	Trichinosis:	
Florida	6	Illinois	1
Idaho	5	New York	3
Illinois	127	Pennsylvania	1
Louisiana	12		•
New York	261	Tularaemia:	
Oklahoma 1	2	California	1
Oregon	27	Illinois	2
Pennsylvania	380	Louisiana	1
Porto Rico	4	Virginia	2
Washington	22	Typhus fever:	
Ophthalmia neonatorum:	ı	Florida	3
California	2	New York	2
Illinois	7	Virginia	2
New York	3	Undulant fever:	
Oklahoma 1	1	California	4
Pennsylvania	13	Idaho	2
Porto Rico	4	Illinois	8
Paratyphoid fever:	*	Louisiana	3
California	7	New York	-
Illinois	9	Oklahoma 1	11
Louisiana			1
New York	3	Oregon	2
	16	Pennsylvania	3
Oregon	1	Virginia	2
Porto Rico	3	Washington	7

¹ Exclusive of Oklahoma City and Tulsa.

² Exclusive of New York City.

Vincent's angina:	Cases	Whooping cough—Continued.	Cases
New York 3	76	Louisiana	18
Oklahoma 1	4	New York	1, 754
Oregon	13	Oklahoma 1	35
Whooping cough:		Oregon	43
California	704	Pennsylvania	1, 479
Florida		Porto Rico	165
Idaho		Virginia	518
Illinois		Washington	

¹ Exclusive of Oklahoma City and Tulsa.

Cases of certain communicable diseases reported for the month of March, 1931, by State health officers

State	Chicken pox	Diph- theria	Measles	Mumps	Scarlet fever	Small- pox	Tuber- cu- losis	Typhoid and para- typhoid fever	Whoop- ing cough
Maine New Hampshire 1	216	12	214	262	143	0	47	4	176
Vermont Massachusetts Rhode Island Connecticut	68 1, 050 93 392	201 26 33	25 2, 023 52 3, 065	145 683 128 327	43 1, 637 266 277	0 0 0 0	19 582 55 147	0 9 0 2	96 877 38 396
New York New Jersey Pennsylvania	3, 038 2, 044 4, 795	534 273 415	8, 213 3, 275 15, 170	1, 913 267 2, 228	4, 119 1, 339 2, 464	36 0	1, 849 475 740	38 8 55	2, 223 708 1, 007
Ohio Indiana Illinois Michigan Wisconsin	2, 452 473 1, 563 1, 574 1, 845	199 124 518 160 55	3, 591 3, 026 7, 163 789 1, 887	1, 803 84 1, 497 618 3, 382	2, 285 1, 347 2, 415 1, 752 654	256 447 169 86 24	661 194 1, 079 625 120	21 7 15 11 5	440 232 657 879 497
Minnesota Iowa Missouri North Dakota South Dakota Nebraska Kansas	783 471 448 154 201 390 686	89 25 208 19 43 42 57	455 82 1, 853 156 330 30 118	158 168 117 11 602 522	501 492 1, 591 106 94 226 279	30 335 213 35 112 224 487	210 23 2254 21 17 20 163	5 4 31 4 0 2 2	238 85 107 60 41 79 122
Delaware	21 683 204 840 343 718 304 186 290	11 64 61 110 38 104 132 29 35	388 4, 829 950 3, 358 364 2, 980 485 546 702	101 382 	101 371 127 205 118 219 25 337 26	0 0 10 56 5 12 3 6	18 249 82 158 67 139 130 44	1 11 1 8 16 5 18 27 11	9 124 35 519 208 637 182 131 60
Kentucky ¹	443 248 1,090	57 86 60	1, 651 2, 000 326	159 353 510	561 113 110	73 56 177	229 511 159	28 14 15	169 77 391
Arkansas Louisiana Oklahoma Louisiana Texas	300 86 82	20 88 40 149	127 78 102	82 10 23	92 100 133 142	107 121 260	2 7 2 135 32	12 24 12 15	97 23 61
Montana Idaho Wyoming Colorado New Mexico Arizona	126 87 107 378 113 54	14 13 2 42 20 12	43 57 14 1, 453 289 622	196 83 48 275 95 26	106 121 121 219 43 21	19 31 16 22 16 9	64 13 2 2 54 78 97	6 20 4 2 3 2	173 179 64 283 32 20
Utah 3 Nevada	20	1	263	7	1	0	3.9	0	2
Washington Oregon California	575 297 2, 509	35 18 222	219 331 5, 969	246 276 1, 492	221 93 620	165 112 216	59 45 962	13 8 33	260 50 1, 211

¹ Report not received.
2 Pulmonary.

² Exclusive of New York City.

Reports received weekly.
 Exclusive of Oklahoma City and Tulsa.

Case rates per 100,000 population (annual basis) for the month of March, 1931

							,		,
State	Chicken pox	Diph- theria	Measles	Mumps	Scarlet fever	Small- pox	Tuber- cu- losis	Typhoid and para- typhoid fever	Whoop- ing cough
Maine New Hampshire 1	318	18	315	385	210	0	69	6	259
Vermont	222	13	82	474	140	0	62	0	314
Massachusetts	288 157	55 44	554 88	187 216	448 449	0	159 93	2 0	240 64
Connecticut		24	2, 208	236	200	Ŏ	106	ĭ	285
New York	278	49	752	175	377	3	169	3	201
New Jersey Pennsylvania	580 579	77 50	929	76 269	380 298		135 89	2 7	204 122
Ohio	427	35	626	314	398	45	115	4	77
Indiana	170	45	1,088	30	484	161	70	3	83
Illinois	237	78	1,085	227	366	26	163	2	100
Michigan Wisconsin	372 730	38 22	186 746	146 1, 338	414 259	20 9	148 47	3 2	208 197
Minnesota	357	41	207		228	14	96	2	108
Iowa	224	12	39	75	234	159	11	2	40
Missouri North Dakota	144 265	67 33	597 268	54 201	512 182	69 60	² 82	10 7	34 103
South Dakota	338	72	555	19	158	188	29	10	69
Nebraska	331 426	36 35	25 73	511 324	192 173	190 303	17	2	67
	103		1.901	495	495		101	1	76
Delaware	486	54 46	3, 438	272	264	0	88 177	5 8	44 88
District of Columbia	487	146	2, 269		303	Ŏ	196	2	84
Virginia West Virginia	406 229	53 25	1, 623 243		99 79	5 37	76 45	11	251 139
North Carolina	261	38	1,081		79	2	40	2	231
South Carolina	205 75	89 12	327 221	94 72	17 136	8	94	12	123
Georgia	223	27	541	27	20	5	53 34	11 8	53 46
Kentucky 8								-	
Tennessee	197	25	733	71	249	32	102	12	75
Alabama Mississippi	109 630	38 35	878 188	155 295	50 64	25 102	224 92	6 9	34 226
Arkansas	189	13	80	52	58	67	34	8	61
Louisiana	47	48	43	6	55	67	2 74	13	13
Oklahoma 4 Texas	46	22 29	57	13	75 28	146	18	7 3	84
Montana.	276	31	94	429	232	42	140	- 1	379
Idaho	229	34	150	219	319	82	34	13 53	472
Wyoming	549	10	72	246	621	82	10	21	329
Colorado	425 309	47 55	1, 633 790	309 260	246 117	25 44	61 213	8	318 87
Arizona	142	32	1, 635	68	55	24	255	5	53
Utah 3 Nevada	254	13	3, 340	89	13	0	1114		25
Washington	426	26	162	182	164	122	44	10	193
Oregon	359	22	400	333	112	135	54	10	60
California	496	44	1, 181	295	123	43	190	7	240
			I	<u>i</u>	1			- 1	

Report not received.
 Pulmonary.

RECIPROCAL NOTIFICATIONS

Notifications regarding communicable diseases sent during the month of August, 1931, by departments of health of certain States to other State health departments

Disease	Cali- fornia	Connec- ticut	Illinois	Maine	Minne- sota	Missouri	New Jersey	New York
DiphtheriaGonorrheaMalaria		2						
Pneumonia. Poliomyelitis. Scarlet fever	i	2		1	2 1	1	1	3
Trachoma Tuberculosis Typhoid fever			16		32 1		2	5

<sup>Reports received weekly.
Exclusive of Oklahoma City and Tulsa.</sup>

GENERAL CURRENT SUMMARY AND WEEKLY REPORTS FROM CITIES

The 97 cities reporting cases used in the following table are situated in all parts of the country and have an estimated aggregate population of more than 33,-240,000. The estimated population of the 90 cities reporting deaths is more than 31,695,000. The estimated expectancy is based on the experience of the last nine years, excluding epidemics.

Weeks ended September 19, 1931, and September 20, 1930

	1931	1930	Esti- mated ex- pectancy
Diphtheria: Cases reported			
46 States	1, 149	804	
97 cities	217	291	468
Measles:			
45 States	490	463	
97 cities	142	101	
Meningococcus meningitis:			
43 States	68	€7	
97 cities	41	28	
Poliomyelitis:	i		
43 States	1, 268	503	
Scarlet fever:	1		_
46 States	1, 226	1, 050	
97 cities	366	381	332
Smallpox:	1		
45 States	69	128	
97 cities	4	28	7
Typhoid fever:	i i		
46 States	1,037	940	
97 cities	267	137	157
Influenza and pneumonia: Deaths reported	1		
90 cities	377	356	
Smallpox:	311	999	
EO cities	0	0	
EU CILIES.	<u> </u>		

City reports for week ended September 19, 1931

The "estimated expectancy" given for diphtheria, poliomyelitis, scarlet fever, smallpox, and typhoid fever is the result of an attempt to ascertain from previous occurrence the number of cases of the disease under consideration that may be expected to occur during a certain week in the absence of epidemics. It is based on reports to the Public Health Service during the past nine years. It is in most instances the median number of cases reported in the corresponding weeks of the preceding years. When the reports include several epidemics, or when for other reasons the median is unsatisfactory, the epidemic periods are excluded, and the estimated expectancy is the mean number of cases reported for the week during nonepidemic years.

If the reports have not been received for the full nine years, data are used for as many years as possible, but no year earlier than 1922 is included. In obtaining the estimated expectancy, the figures are smoothed when necessary to avoid abrupt deviation from the usual trend. For some of the diseases given in the table the available data were not sufficient to make it practicable to compute the estimated expectancy.

1		-	theria	Influ	lenza	I	1	Pneu-
Division, State, and city	Chicken pox, cases reported		Cases reported	Cases reported	Deaths reported	Measles, cases re- ported	Mumps, cases re- ported	monia, deaths reported
NEW ENGLAND					l			
Maine:				i				
Portland	0	0	0		0	0	0	1
New Hampshire:	_	ا ا	•			_	_	
Concord	0	0	0		0	0	0	0
Manchester	0	0	0		0	0	0	0
Nashua	0	0	0		0	0	0	0
Vermont:		_	_			_		_
Barre	0	0	0		0	0	0	0
Burlington	0	0	0		0	0	0	0
Massachusetts:								
Boston	10	14	10	4	0	3	0	8
Fall River	1	2	0	1	0	1	0	1
Springfield	0	1	0		0	0	1	3
Worcester	1	3	1	1	0	0	17	2
Rhode Island:	_					-		_
Pawtucket	0	1	2	0	0	1	0	0
Providence	ŏ	3	1		ŏl	7	i	i
Connecticut:	Ĭ	-	- 1			1	-	_
Bridgeport	1	2	1	1 1	1	0	o l	8
Hartford	ñ	ĩ	ñ		á	ŏl	ĭI	2
New Haven	ŏ	أة	ŏ		ŏl	ĭI	اة	ñ
11011 TIGA (TT)	0,	0 1	٠,		0,	4 1	٠,	v

	i	Diph	theria	Infl	uenza	1		Pneu-
Division, State, and city	Chicken pox, cases reported	Cases, estimated expect- ancy	Cases reported	Cases reported	Deaths reported	Measles, cases re- ported	Mumps, cases re- ported	monia, deaths reported
MIDDLE ATLANTIC								1
New York: Buffalo		7	2	1		1	1	7
New York	12	74	33	8	3	16	13	86
Rochester Syracuse	2	2 1	0		0	3 1	8	1
New Jersey:							U	1
Camden Newark	0	2 8	0 3		0	0	0	7
Trenton	ō	i	ŏ		i	2	0	1
Pennsylvania: Philadelphia	11	27	3	2	1	5		1
Pittsburgh	4	11	8	6	î	9	3	24 19
Reading	0	1	0		0	0	Ö	0
EAST NORTH CENTRAL								
Ohio:								
Cincinnati Cleveland	2 12	5	5		0	.0	1	4
Columbus	12	22 2	0	2 1	0 1	10 0	28 1	12
Toledo	2	4	Ō		Ō	ĭ	Ô	4
Indiana: Fort Wayne	0	1	1		0	0	0	,
Indianapolis	1	4	ī		0	2	8	2 1 2
South Bend Terre Haute	0	0	.0		0	0	0	2 1
Illinois:		1	- 1			1		1
Chicago Springfield	22	56 0	27 0	2	3	8	7	26
Michigan:	1	1	1		· ·	٧	3	1
DetroitFlint	8	29	7		1	1	2	14
Grand Rapids	1	2	1 0		0	0 3	6	2 0
Wisconsin: Kenosha							-	
Madison	0	0	0		0	0	13 3	0
Milwaukee	12	5	2		0	4	12	5
Racine Superior	3	0	0		0	0	9	0 5 0
WEST NORTH	1	-	-		•	Ĭ	Ů	•
CENTRAL Minnesota:	1	1		į		1		
Duluth	0	0	0		0	0	1	2
Minneapolis	7	14	2		2	2	23	4
St. PaulIowa:	5	7	1 .		0	2	2	0
Davenport	3	0	0 .			0	0	
Des Moines	0	0	1 0			0	0	
Waterloo	ŏ	ō	i .			ŏ	ō	
Missouri: Kansas City	0	2	2		0	1	o	2
St. Joseph	1	0	1 .		ŏ	0	Ö	3
St. Louis North Dakota:	2	17	8 -			1	1	3
Fargo	1	0	0 -		0	0	1	0
South Dakota:	9	0	0 -			0	0	
Aberdeen	12	0	0			0	0	
Sioux Falls Nebraska:	0	0	0 -	-		0	0	
Omaha	0	5	2 _		0	0	0	1
Kansas: Topeka	0		,	1			,	•
Wichita	2	î	4		ŏ	ĭ	1	Ö
SOUTH ATLANTIC			1			i	1	
Delaware: Wilmington	o	1	0 -	- 1	اه	0		^
Maryland:	1	ı	1		0	į.	0	0
Baltimore Cumberland	12	15 1	8	3	0	3	4	10
Frederick	ŏ	ŏ	0 -		0	0	0	1 0
District of Columbia: Washington			1		1	i	- 1	
Virginia:	0	9	9	1	1	0	0	6
Lynchburg	0	2	1		0	2	0	0
Norfolk Richmond	0	1 12	1		0	0	0	2 2
Roanoke	ŏ	3	6		δj	ŏ	ŏ	í
							•	

	I	Diph	theria	Infl	uenza	1		Draw
Division, State, and city	Chicken pox, cases reported	Cases, estimated expect- ancy	Cases reported	Cases reported	Deaths reported	Measles, cases re- ported	Mumps, cases re- ported	Pneu- monia, deaths reported
SOUTH ATLANTIC—COD								
West Virginia: Charleston		o	0	İ		1	o	م ا
Wheeling	ŏ	ì	ŏ		ŏ	Ô	ŏ	ŏ
North Carolina: Raleigh	1	2	1		0	0	0	0
Wilmington Winston-Salem	0	0 2	0		0	0	0	9
South Carolina:					1		_	
Charleston Columbia	8	1 1	0 1	3	0	0	0	2 1
Greenville Ceorgia:	0	1	0		0	0	0	0
Atlanta	1 0	5 0	3		0	1 0	1	4
Savannah	ŏ	i	0	4	0	ŏ	0	0
Florida: Miami	0	2	1		0	1	. 0	1
Tampa	0	ī	Ō		Ŏ	Ö	Ō	Õ
EAST SOUTH CENTRAL								
Kentucky: Covington	o	0	0		0	0	0	0
Tennessee: Memphis	1	3	7		0	0	0	1
Nashville	Ô	2	4		ŏ	ŏ	ŏ	6
Alabama: Birmingham	0	3	0		0	0	0	2
Mobile	0	1 2	3 2		0	0	0	0
WEST SOUTH CENTRAL		- 1	-			Ĭ	Ĭ	
Arkansas:								
Fort Smith Little Rock	0	0	2 1		0	1 1	0	
Louisiana:		8	1		0	0	0	8
New Orleans Shreveport	1 0	î	. 0	1	ŏ	3	ŏ	õ
Oklahoma: Muskogee	ol	0	3		0	o	0	0
Texas:	0	6	2		0	0	0	
Dallas Fort Worth	1	1	1		Ŏ	0	Ö	1 1 2
Galveston Houston	0	0 5	0		0	0	0	4
San Antonio		2						
MOUNTAIN	Ī	1				l		
Montana: Billings	0	0	o l		o l	7	o	Q
Great Falls Helens	1 0	0	0		0	0 5	0	1 0 1
MissoulaIdaho:	Ŏ	Ō	Ō		Ō	Ö	Ō	Ĭ
Boise	1	0	0		0	0	1	1
Colorado: Denver	2	8	1		0	2	2	5
Pueblo	1	0	0		0	0	4	0
Albuquerque	0	0	0	i	0	0	0	0
Arizona: Phoenix	0	o	o		0	0	0	1
Utah: Salt Lake City	0	2	1		0	0	2	1
Nevada:	1	_	0			0	0	0
Reno	0	0	•		١	١	١	U
Washington:		į						
Seattle Spokane	7	3	0			0	1 0	
Tacoma	i	2	4		0	ŏ	ŏ	2
Oregon: Portland	1	5	2		0	4	5	2
SalemCalifornia:	Ō	0	0		Ō	Ō	1	0
Los Angeles	2	13	7	19	1	4	4	24
Sacramento San Francisco	1 9	8	0	1	0	20	1 3	3 6
				1				

	Scarlet fever			Smallpo)X	Tuber-	T	phoid f	ever	Whoop	
Division, State, and city	Cases, esti- mated expect- ancy	Cases re- ported	Cases, esti- mated expect- ancy	Cases re- ported	Deaths re- ported	re-	mated	Cases re- ported	Deaths re- ported	ing cough,	Deaths, all causes
NEW ENGLAND											
Maine: Portland	1	o	0	0	0	o	0	2	0	5	22
New Hampshire: Concord	0	0	0	0	0	0	0	0	0	0	7
Manchester Nashua	1 0	0	0	0	0	0	0	0	0	0	16
Vermont: Barre	0	1	0	0	0	1	0	0	0	0	1
Burlington Massachusetts:	ŏ	ō	ŏ	š	Ŏ	Ō	ŏ	ŏ	ŏ	ĭ	7
Boston Fall River	16 1	16 4	0	0	0	9	3 0	3 0	1 0	24 1	202 21
Springfield Worcester	1 3	0 6	ŏ	ŏ	ŏ	0 2	Ŏ 1	0	Ŏ	1 13	19 42
Rhode Island: Pawtucket	0	0	0	0	0	1	0	0	0	0	12
Providence Connecticut:	2	7	ŏ	ŏ	ŏ	î	2	ŏ	ŏ	5	57
Bridgeport Hartford	2	1	0	0	0	1	0	0	0	1	34 36
New Haven	í	6	ŏ	ŏ	ŏ	2	2	2	ŏ	5 2	31
MIDDLE ATLANTIC											
New York: Buffalo New York	6 27	6 22	0	0	0	9 83	1 37	1 25	0	21 166	125 1, 343
Rochester Syracuse	2 2	13 7	0	0	0	0	0	0	0	6 24	76 46
New Jersey: Camden	0	0	0	0	0	0	1	3	0	3	25
, Newark Trenton	3	6 2	0	0	0	7 3	1	0	0	81	100 37
Pennsylvania: Philadelphia	19	28	o	0	0	35	10	1	0	109	442
Pittsburgh Reading	11 0	12	0	0	0	8 0	3 1	5	0	24 0	160 19
EAST NORTH CENTRAL								İ			
Ghio: Cincinnati	اء	17	o	,	0	4	2	- ,	o	10	147
Cleveland	13	10 2	0	0	0	12 5	4	130	7	102	198 79
Toledo Indiana:	4	5	ĭ	ĭ	ŏ	6	1 2	ŏ	ŏ	19	56
Fort Wayne	1 3	o l	0	0	0	9 7	1 1	1 0	0	5 8	23
Indianapolis South Bend Terre Haute	1	5 1 0	ö	ŏ	0	i	1 0	ö	ö	ő	15 18
Illinois:	33	- 1	0	0	0	46	6	6		182	632
Chicago	30	40 2	ŏ	ŏ	ŏ	0	î	õ	ŏ	102	15
Detroit	27	9	Q	o l	8	25. 0	4 0	8	1 0	147	229 21
Grand Rapids. Wisconsin:	5	3	8	0	ŏ	2	ŏ	ő	ŏ	8	28
Kenosha Madison	ņ	1 0	0	0	0	0	0	2	0	5	4
Milwaukee	1 8 2	4 3	ö	8 -	0	6	0 1 0	0 -	0	49	97 9
Racine Superior Superior	1	2	81	0 1	8	ĭ	8	8	ŏ	õ	6

	Scarle	Scarlet fever		Smallpo)X	Tuber-				Whoop-	
Division, State, and city	Cases, esti- mated expect- ancy	Cases re- ported	Cases, esti- mated expect- ancy	Cases re- ported	Deaths re- ported	re-	Cases, esti- mated expect- ancy		Deaths re- ported	ing cough, cases re- ported	Deaths, all causes
WEST NORTH CENTRAL											
Minnesota:											1
Duluth Minneapolis	16	6	0	0	0	2	0 1	8 2	0	8	36 105
St. Paul	8	ŏ	ŏ	ŏ	ŏ	ĭ	î	õ	ŏ	. 7	49
Iowa: Davenport	ا ا	0	0	0			0	0		3	
Des Moines	2	1	0	2			0	1		0	24
Sioux City Waterloo	0	1 0	0	0			0	1 0		2	
Missouri:	1			_						1	
Kansas City	4	0	0	0	0	6	2	0	0	3	97
St. Joseph St. Louis	12	13	ŏ	ŏ	0	13	0 6	0 7	0	0 51	200
North Dakota:		0	0	0	0	0		0	-		
Fargo Grand Forks	2 0	ő	ŏ	ö	U	0	0	ŏ	0	1	
South Dakota:										-	
Aberdeen Sioux Falls	0	0	0	0			0	0		7	11
Nebraska:		- 1		- 1	_						
Omaha Kansas:	1	0	0	0	0	2	1	0	0	0	52
Topeka	1	1	0	0	0	0	2	0	0	0	10
Wichita	2	4	0	0	0	1	0	2	0	, 0	34
SOUTH ATLANTIC											
Delaware:			_	_ 1				_			
Wilmington Maryland:	0	1	0	0	0	0	0	0	0	2	36
Baltimore	6	10	0	0	0	11	8	5	2	125	206
Cumberland Frederick	0	1 0	0	0	0	1 0	1 0	0	1 0	0	13
District of Colum-	•	١	١	١	١	١	١	١	"		
bia: Washington	6	4	0	0	0	11	3	2	2	25	
Virginia:	١٥	*	١	١	١	ł	°		2	25	141
Lynchburg Norfolk	0	2	0	0	0	2 4	1	0 3	0	.0	15
Richmond	1 4	3 8	0	ŏ	ŏ	õ	1 2	ő	0	11 2	45
Roanoke	1	0	0	0	0	1	1	0	Ō	2	18
West Virginia: Charleston	1	o	o	o	0	0	2	o	o	3	26
Wheeling	1	Ö	0	Ó	Ō	Ō	2	Ŏ	ŏ	ŏ	15
North Carolina: Raleigh	0	2	o	0	0	1	o	0	0	12	9
Wilmington	Ō	0	0	0	0	1	0	0	Ō	3	18
Winston-Salem South Carolina:	2	1	1	0	0	1	1	0	0	12	15
Charleston	0	4	0	0	0	3	2	1	1	0	22
Columbia Greenville	0	0	0	0	0	3 0	0	2	8	0	22
Georgia:	- 1	i	i	i	1	- 1	į	1		1	
Atlanta Brunswick	5	1	0	0	0	3 0	3 0	2	0	1 0	60
Savannah	ŏ	ŏ	ŏ	ŏ	ŏ	ĭ	ŏ	ō	ŏ	ĭ	5 26
Florida: Miami	0	0	0	o	0	o	1	2	0	0	20
Tampa	i	ŏ	ŏ	ŏ	ŏ l	2	δl	ő	ĭ	8	20 26

	Scarle	t fever		Smallp	ox	Tuber-				Whoop	
Division, State, and city	Cases, esti- mated expect- ancy	Cases re- ported	Cases, esti- mated expect- ancy	Cases re- ported	Deaths re- ported	re-	mated	Cases re- ported	Deaths re- ported	ing cough,	Deaths, all causes
EAST SOUTH CENTRAL											
Kentucky: Covington	0	1	- 0	0	0	0	1	0	0	0	17
Tennessee: Memphis Nashville	2 2	3	0	0	0	11 5	5 5	3 0	0	11 7	102 57
Alabama: Birmingham Mobile Montgomery	4 0 0	4 2 3	0 0 0	0 0 0	0	5 1	4 0 0	4 0 1	0 0	4 0 6	53 19
WEST SOUTH CENTRAL											
Arkansas: Fort Smith Little Rock Louisiana:	1 1	0	0	0	<u>-</u>	1	0 1	0	-	0	6
New Orleans Shreveport Oklahoma:	2 1	3 1	0	0	0	10 1	4 0	9 2	0 1	0 6	130 26
Muskogee Texas:	0	0	0	0	0	0	0	1	0	0	
Dallas Fort Worth Galveston Houston San Antonio	2 2 0 1 1	6 4 0 3	1 0 0 0	0 0 0	0 0 0	4 2 0 5	2 1 0 1 0	0 1 0 1	0 0 0 1	10 0 0 0	41 31 11 83
MOUNTAIN	_										
Montana: Billings	0 0 0	0 0 0	0 1 0 0	0 0 0 0	0 0 0	0 0 0 2	0 0 0 0	0 0 0	0 0 0 1	1 0 1 0	12 7 7 8
Boise Colorado:	0	0	0	0	0	0	0	0	0	1	, 5
Denver Pueblo New Mexico:	4 0	9	0	0	0	8	1	0 2	0	7 0	64 12
Albuquerque Arizona:	0	0	0	0	0	3	1	1	0	0	7
Phoenix Utah: Salt Lake City_	1	0	0	0	0	1 2	0 2	0	0	0	35
Nevada: Reno	0	0	0	o	0	0	0	0	0	0	6
PACIFIC		1						l		ĺ	
Washington: Seattle Spokane Tacoma	5 2 1	8 0 1	1 1 1	0	0	0	1 1 0	1 0 0	0	12 0 3	26
Oregon: Portland Salem	3	0	2	1 0	0	4 0	2 0	0	0	0	62
California: Los Angeles Sacramento San Francisco.	9 1 6	16 0 3	1 0 0	0 0 2	0	18 2 9	2 0 0	3 2 12	1 0 1	10 3 5	252 30

	000	ningo- cus ingitis	Letha ceph	rgic en- alitis	Pel	lagra	Poliomyelitis (infan- tile paralysis)		
Division, State, and city	Cases	Deaths	Cases	Deaths	Cases	Deaths	Cases, esti- mated expect- ancy	Cases	Deaths
NEW ENGLAND									
Maine: Portland	0	o	0	0	0	o	0	2	1
New Hampshire: Concord	0	0	0	0	0	0	0	1	. 0
Manchester Nashua	0	Ŏ	0	Ŏ	Ŏ	0	Ŏ	1	Ŏ
Massachusetts: Boston	0	0	0	0	0	0	4	34	ŀ
Fall River Springfield	6 0	1 0	0	0	0	0	0	6 10	1 0
Worcester Rhode Island:	0	0	0	0	0	0	1	3	1
Pawtucket Providence	0	0	0	0	0	0	0 1	2 2	0 1
Connecticut: Bridgeport	0	Q	Q	0	0	0	0	8	0
Hartford New Haven	0	0	0	0	0	0	0 1	8 8	2 0
MIDDLE ATLANTIC									,
New York: New York	13	1	4	1	0	0	14	226	38
Rochester Syracuse	0	0	Ô	Õ	Ŏ	ŏ	1 2	200	0
New Jersey: Newark	0	0	1	0	0	0	0	6	1
TrentonPennsylvania:	ŏ	ŏ	Ō	0	ŏ	Ŏ	ŏ	ŏ	î
Philadelphia Pittsburgh	0	1 2	0 2	0	0	0	1	9	1
EAST NORTH CENTRAL									
Ohio:			0			1			•
Cleveland Toledo	1	0	ŏ	ŏ	0	Ŏ	3 0	5 1	0
Indiana: IndianapolisIllinois:	0	1	0	0	0	0	1	0	0
Chicago Springfield	3	1 0	1 0	0	1 0	1 0	4 0	6	1
Michigan: Detroit	3	0	2	1		0	3	44	2
Grand Rapids	ŏ	ŏ	ō	õ	ŏ	ŏ	ŏ	ī	i
Kenosha Madison	0	0	0	0	0	0	0	1 4	0
Milwaukee	0	0	0	0	0	0	1 0	5	0 0 0
Superior	0	0	Ò	0	Ō	0	0	1	Ō
WEST NORTH CENTRAL	İ		l				l	ĺ	
Minnesota: Duluth	o l	o l	o l	o l	o l	o	o	.9	
Minneapolis St. Paul	0	0	0	0	0	0	1 1	15 34	1 2
Iowa: Des Moines Missouri:	0	0	0	0	0	0	1	2	0
Kansas City St. Louis	0 3	0	0	0	0	0	1 0	1	1
North Dakota: Grand Forks	0	0	0	0	0	0	ol	1	0
Nebraska: Omaha	2	0	0	0	o	ol	o l	1	0

	coc	ingo- cus ngitis	Letha: ceph	rgic en- alitis	Pell	lagra		myelitis e paraly			
Division, State, and city	Cases	Deaths	Cases	Deaths	Cases	Deaths	Cases, esti- mated expect- ancy	Cases	Deaths		
SOUTH ATLANTIC 1											
Maryland: Baltimore Cumberland Virginia:	0 1	0	0	8	0	0	1 0	2 0	0		
Lynchburg Richmond North Carolina	0	0	0	0	0	1 0	0	0	0		
Raleigh	0	0	0	0	0 0		0	1	0		
Charleston	0	0	0	0	6 0	0	0	0	0		
Georgia: Atlanta ¹ Brunswick Savannah ¹	0 0 1	0	ŏ	Ŏ	0 0 0	0 0 0	0 0 0 1 0 0		0 0 0	1 0 0	1 0 0
EAST SOUTH CENTRAL											
Tennessee: Memphis Nashville	1	0	0	0	0 1 0		0	5 1	1 0		
WEST SOUTH CENTRAL											
Louisiana: New Orleans Texas: 1	0	0	1	1	1	0	0	2	0		
Dallas Houston	0	0	0	0	1 0	1 1	0	0	0		
MOUNTAIN											
Montana: Great Falls Missoula	0	0	0	0	0	0	0	1 1	0		
New Mexico: Albuquerque Utah:	0	0	0	0	0	0	0	1	1		
Salt Lake	1	0	0	0	0	0	1	0	0		
Reno	0	0	0	0	0	0	1	1	0		
Washington:											
Seattle Tacoma Oregon:	0	0	0	0	0	0	0	2 2	0		
California:	0	0	0	0	0	0	1	1	1		
Los Angeles San Francisco	0 1	1	0	0	1 1	1 0	0	1 1	0		

¹ Typhus fever, 6 cases: 1 case at Atlanta, Ga.; 2 cases at Savannah, Ga.; 2 cases at Miami, Fla.; and 1 case at Fort Worth, Tex.

The following tables give the rates per 100,000 population for 98 cities for the 5-week period ended September 19, 1931, compared with those for a like period ended September 20, 1930. The population figures used in computing the rates are estimated mid-year populations for 1930 and 1931, respectively, derived from the 1930 census. The 98 cities reporting cases have an estimated aggregate population of more than 33,000,000. The 91 cities reporting deaths have more than 31,500,000 estimated population.

Summary of weekly reports from cities, August 16 to September 19, 1931.—Annual rates per 100,000 population compared with rates for the corresponding period of 1930 i

DIPHTHERIA CASE RATES

	<u> </u>			<u> </u>	Week	ended-				
	Aug. 22, 1931	Aug. 23, 1930	Aug. 29, 1931	Aug. 30, 1930	Sept. 5, 1931	Sept. 6, 1930	Sept. 12, 1931	Sept. 13, 1930	Sept. 19, 1931	Sept. 20, 1930
98 cities	2 30	.33	2 31	38	3 37	40	35	- 44	4 34	46
New England Middle Atlantie East North Central West North Central South Atlantie East South Central West South Central Mountain Pacific	67 19 228 31 24 35 68 44 35	44 27 40 25 40 12 63 44 22	41 18 233 36 63 52 34 17 24	53 20 45 27 64 12 66 70	55 24 38 5 26 34 81 6 107 52 27	39 29 48 35 66 48 56 44 32	58 26 32 34 45 99 41 26 29	60 26 63 56 68 24 45 35 22	36 22 29 42 73 93 4 52 17 29	34 36 74 48 46 24 63 26
		MEA	SLES (CASE	RATES					
98 cities	2 29	28	2 22	20	³ 19	24	14	16	4 22	16
New England Middle Atlantic. East North Central West North Central South Atlantic East South Central West South Central West South Central Mountain Pacific	63 25 237 13 20 23 7 70 22	65 31 21 19 20 6 0 26 40	63 13 223 8 4 6 24 52 53	22 22 7 27 32 12 10 35 30	58 14 11 * 9 8 6 . • 0 52 67	36 27 12 31 28 24 0 53 34	29 8 13 11 6 6 10 35 45	41 19 9 15 6 6 3 35 16	31 18 17 13 14 0 420 122 53	19 16 14 19 22 0 9 44 18
	SC.	ARLET	r FEVI	ER CA	SE RA	TES				
98 cities	2 43	32	3 41	41	3 48	42	49	50	4 57	61
New England Middle Atlantic East North Central West North Central South Atlantic East South Central West South Central Mountain Pacific	99 38 2 57 19 36 17 27 44 31	51 25 35 35 30 30 35 88 28	46 30 243 31 30 70 64 165 39	56 26 47 43 72 102 14 88 26	87 37 56 50 51 87 655 26 43	60 24 47 58 72 60 63 35 28	106 30 64 36 55 64 41 61 39	56 26 84 35 56 36 24 79 63	87 43 62 50 71 81 4 52 87 55	77 45 90 45 44 36 52 76
		SMALI	JPOX (CASE	RATES					
98 cities	21	2	21	2	31	3	1	3	41	4
New England Middle Atlantic East North Central West North Central South Atlantic East South Central West South Central Mountain Pacific	0 0 2 0 6 4 0 0 0 4	0 0 0 8 2 0 7 0	0 0 20 4 4 0 0 0 0	0 0 0 8 0 0 3 0	0 0 4 5 4 0 0 0 6 0	0 0 2 14 4 0 0 0	2 0 2 6 0 6 0 0	0 0 2 27 0 0 0 0 8	0 0 1 0 0 0 0 0 40	0 9 21 0 0 0

¹ The figures given in this table are rates per 100,000 population, annual basis, and not the number of cases reported. Populations used are estimated as of July 1, 1931 and 1930, respectively.

² Terre Haute, Ind., not included.

³ St. Paul, Minn., and Fort Smith, Ark., not included.

⁴ San Antonio, Tex., not included.

⁵ St. Paul, Minn., not included.

⁵ St. Paul, Minn., not included.

⁶ Fort Smith, Ark., not included.

Summary of weekly reports from cities, August 16 to September 19, 1931—Annual rates per 100,000 population compared with rates for the corresponding period of 1930—Continued

TYPHOID	FEVER	CARE	DATES
	PEVER	CADE	RAILO

				Week	ended—				
Aug. 22, 1931	Aug. 23, 1930	Aug. 29, 1931	Aug. 30, 1930	Sept. 5, 1931	Sept. 6, 1930	Sept. 12, 1931	Sept. 13, 1930	Sept. 19, 1931	Sept. 20, 1930
2 21	19	1 22	24	3 20	21	23	26	4 42	22
5 14 2 11 19 55 70 91 9	17 13 9 21 60 78 24 26 6	22 20 20 10 13 38 47 98 9	12 20 10 19 88 42 66 44 8	7 13 16 5 6 49 41 6 78 44 10	12 20 12 14 58 48 45 9	7 13 10 13 79 35 91 35 27	22 24 17 21 70 48 52 62 4	22 16 91 38 26 47 448 28 35	12 15 11 29 68 48 63 0
I	NFLUI	ENZA 1	DEATI	H RAT	ES				
2 2	3	12	4	8 2	3	4	3	4 3	3
2 2 2 2 3 6 0 0 0 7	0 3 1 0 8 0 4 9	0 2 11 3 6 13 0 0	0 3 4 3 8 6 7 0 2	2 1 1 4 3 2 6 10 0 2	0 3 2 6 8 0 11 9	2 4 3 9 2 0 17 0 2	0 4 3 0 2 19 0	2 3 3 6 4 0 40 0 2	2 2 2 0 0 26 7 18 0
PI	NEUM	ONIA I	DEATI	RAT	ES				
1 48	45	2 48	52	å 50	53	55	54	4 59	57
36 56 32 44 63 57 59 44 53	56 53 27 36 52 65 57 53 40	46 60 26 50 69 57 59 61 29	51 57 50 39 60 45 36 53 45	24 62 33 73 61 38 83 96	56 65 36 51 68 91 50 53 27	58 65 36 44 63 82 73 70 46	68 63 43 45 58 26 57 123 25	50 66 45 44 57 57 482 78 84	56 65 42 75 58 71 46 115 40
	22, 1931 2 21 5 14 2 11 1 19 5 5 70 91 9 8 II 2 2 2 2 2 2 2 2 2 2 2 2 3 6 0 0 0 7 P1 2 48 36 56 56 56 32 44 63 57 59 44	22, 1931 1930 221 1930 221 19 5 17 14 13 211 9 19 21 55 60 70 78 91 26 8 6 INFLUI 22 3 2 0 3 22 1 3 0 6 6 8 0 0 0 0 4 0 9 7 7 PNEUM 248 45 36 56 53 32 27 44 36 65 57 59 57 44 55	22, 23, 29, 1931 2 21	22, 23, 29, 30, 1931 1930 2 21 19 22 24 5 17 22 12 14 13 20 20 211 9 10 10 19 21 13 19 55 60 38 88 70 78 47 42 91 24 98 66 9 26 9 44 8 6 12 8 INFLUENZA DEATH 2 2 0 0 0 0 2 3 2 2 1 2 1 2 1 4 2 0 0 0 0 2 3 3 2 2 3 2 2 1 3 6 6 8 6 8 0 0 0 13 6 6 8 0 0 0 13 6 6 8 0 0 0 13 6 6 8 0 0 0 13 6 6 8 0 0 0 13 6 6 8 0 0 0 0 7 7 2 2 2 PNEUMONIA DEATH 2 48 45 2 48 52 36 56 46 51 56 53 60 67 32 27 26 50 44 36 50 39 63 52 69 69 63 55 69 69 65 57 45 59 57 45 59 57 45 59 57 45 36 153 66 57 59 57 45 59 57 45 59 57 45 59 57 45 59 57 45 59 57 59 36	Aug. 22, 23, 29, 30, 1931 221 19 22 24 20 5 17 22 12 7 14 13 20 20 13 111 9 10 10 16 19 21 13 19 46 55 60 38 88 49 70 78 47 42 41 91 24 98 66 76 9 26 9 44 44 8 6 12 8 10 INFLUENZA DEATH RAT 2 2 0 0 0 0 2 2 3 1 2 1 1 1 4 1 3 0 3 3 3 2 2 3 1 2 1 1 3 1 3 0 6 6 6 0 4 0 7 0 0 9 7 7 7 2 2 2 2 2 2 2 PNEUMONIA DEATH RAT 2 48 45 2 48 52 2 PNEUMONIA DEATH RAT 2 48 45 2 48 52 2 2 3 3 2 3 4 4 36 50 39 67 3 32 27 26 50 33 4 4 36 50 39 67 3 59 57 59 36 88 3 19 44 3 8 6 50 39 67 3 8 56 45 57 59 36 3 8 36 3 8 36 3 9 63 52 69 60 39 47 36 57 55 55 57 45 38 59 57 59 36 88 8 3 49 70 78 47 42 2 10 0 0 0 0 2 2 2 3 1 2 4 1 3 1 4 1 3 0 3 3 3 3 3 4 5 6 6 6 6 6 7 7 10 7 7 7 2 2 2 2 2 3 2 3 4 4 36 50 39 67 5 50 57 55 59 36 8 8 6 8 8 8 3 49 7 10 7 7 7 2 2 2 2 3 50 3 3 50 3 3 50 3 3 63 56 50 39 67 5 57 55 57 45 38 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88 5 9 57 59 36 88	Aug. Aug. Aug. 30, 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1930 1930 1930 1930 1930 1930 1930 1930 1930 1930 1930 1930 1930 1930 193	22	Aug. Aug. 23, 29, 30, 5, 6, 12, 13, 1930 1931 1930	Aug. Aug. Aug. Sept. Sept. Sept. Sept. 12, 13, 19, 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1930 1931 1931 1930 1931 1931 1931 1930 1931 1931 1931 1930 1931 193

<sup>Terre Haute, Ind., not included.
St. Paul, Minn., and Fort Smith, Ark., not included.
San Antonio, Tex., not included.</sup>

<sup>St. Paul, Minn., not included.
Fort Smith, Ark., not included.</sup>

FOREIGN AND INSULAR

CANADA

Provinces—Communicable diseases—Week ended September 12, 1931.—The Department of Pensions and National Health of Canada reports cases of certain communicable diseases for the week ended September 12, 1931, as follows:

Frovince	Cerebro- spinal fever	Influ- enza	Poliomye- litis	Small- pox	Typhoid fever
Prince Edward Island 1 Nova Scotia 2 New Brunswick Quebec Ontario Manitoba Saskatchewan Alberta British Columbia	1 1	3	75 18 2	12	1 32 32 13 2 2 2 2
Total	3	- 5	100	14	86

¹ No case of any disease included in the table was reported during the week.
2 Two cases of undulant fever.

Quebec Province—Communicable diseases—Week ended September 12, 1931.—The Bureau of Health of the Province of Quebec, Canada, reports cases of certain communicable diseases for the week ended September 12, 1931, as follows:

Disease	Cases	Disease	Cases
Chicken pox Diphtheria Erysipelas German measles Itch Measles	3 34 5 1 1 15	Mumps Poliomyelitis Scarlet fever Tuberculosis Typhoid fever Whooping cough	2 75 26 36 32 35

CUBA

Habana—Communicable diseases—Four weeks ended September 12, 1931.—During the four weeks ended September 12, 1931, certain communicable diseases were reported in Habana, Cuba, as follows:

Disease	Cases	Deaths	Disease	Cases	Deaths
Chicken pox	1 6 2 1 5	1 3	Measles Scarlet fever Tuberculosis Typhoid fever	38 1 26 1 8	3 4 2

¹ Many of these cases are from the island of Cuba, outside of Habana.

CZECHOSLOVAKIA

Communicable diseases—June, 1931.—During the month of June, 1931, certain communicable diseases were reported in the Republic of Czechoslovakia, as follows:

Disease	Cases	Deaths	Disease	Cases	Deaths
Anthrax Cerebrospinal meningitis Diphtheria Dysentery Malaria Paratyphoid fever	16 10 1, 158 19 125 16	3 64 2	Puerperal fever	27 1, 134 209 326 2	11 21 25

JAMAICA

Communicable diseases—Four weeks ended September 12, 1931.—During the four weeks ended September 12, 1931, cases of certain communicable diseases were reported in Kingston, Jamaica, and in the island of Jamaica, outside of Kingston, as follows:

Discase	Kings- ton	Other locali- ties	Discase	Kings- ton	Other locali- ties
Cerebrospinal meningitis Chicken pox Diphtheria Dysentry Leprosy Lethargic encephalitis	1	2 1 1 3 1	Paratyphoid fever Poliomyelitis Scarlet fever Tuberculosis Typhoid fever	39 25	1 1 3 65 62

TRINIDAD

Port of Spain—Vital statistics—August, 1930, 1931.—The following statistics for the month of August, 1930 and 1931, are taken from a report issued by the public health department of Port of Spain, Trinidad:

	1930	1931		1930	1931
Number of births	123 21. 5 104	144 24. 7 123	Death rate per 7,000 population Deaths under 1 year Deaths under 1 year per 1,000 births	18. 2 28 269. 2	21. 1 15 104. 2

CHOLERA, PLAGUE, SMALLPOX, TYPHUS FEVER, AND YELLOW FEVER

From medical officers of the Public Health Service, American consuls, International Office of Public Hygiene. Pan American Sanitary Burseu, health section of the League of Nations, and other sources. The reports contained in the following tables must not be considered as complete or final as regards either the list of countries included or the figures for the particular countries for which reports are given.

CHOLERA

[C indicates cases; D, deaths; P, present]

										Week ended-	popu						
Place	Mar. 8- Apr. 4, 1931	8- Apr. 5- 1, May 2, 1931	May 30, 1931	May 3- May 31- 30, June 1931 27, 1931	 	July, 1931	1931			Aug	August, 1931			æ	September, 1931	er, 193	
					-	ıı	81	83	1	00 -	15	ឌ	8	5	13	91	8
	00 00		2-						HH		HH 10			- 6	4	64	
Swatow Crientsin Crientsin I I I Bombay Crientsin Crient	8,4,	8 11,462 0 5,767	13,604 7,270	10, 18, 001 10, 337	2,677	2, 848 1	5, 707 3, 064 11	6, 628 3,504 11	7,357	9, 848 5, 584	9	=	72	9	100		
		256 310 256 176 12 19 12 20 20 26 10 13	265 149 129 17 17 17	168	82	88.	සැපි කි		4 4 0	27.	-8	220 6	œ54∞	- 533	250 I		
	00000			4.62	 		8-						-				
India (French): Chandernagor Pondicherry	0000	7 0 0 8 18	6 4 4 17 17					2		44-	88-14		8-				

CHOLERA, PLAGUE, SMALLPOX, TYPHUS FEVER, AND YELLOW FEVER-Continued

CHOLERA—Continued [O indicates cases; D, deaths; P, present]

		2	lo morestes cases; D, nestus; F, present	Casses; L	, desco	, F, DE	жены										
										Week ended-	-pepu	_					
Place	Mar. 8- Apr. 4, 1931	Apr. 5- May 2, 1931	May 3- May 31- 30, June 1931 27, 1931	May 31- June 27, 1931		July, 1931	1831			Aug	August, 193	#		8	September, 1931	r, 1931	
					4	Ħ	18	8	1	œ	15	ឌ	83	2		61	8
India (Portuguese)				1		61-											
Indo-China (see also table balow): Cochina—Rachgia		6		-			•		д	1 -		-			-		
Saigon and Cholon.		778	~ <u>%</u>	178:	000	1-00				-				$\overrightarrow{\Pi}$			
Iraq: Abulkhasib		3	9	4	es .	.9	23						9				
												C	~ 덩	88	80	+	
												7	20	8=	106	-12	32
Basra									3	6	88	272	148	4	\$2	<u></u>	18
									2	œ	140	137	64	9	82	& 83	38
Dinwaniyah Province C Dinwaniyah Province C												23	2	ro	2	2	3.1.6
Iwaniyah													Ħ		H		= 12
												Ħ	01	22	8	83	288
Nasiriyah													0	38:	389	0 28	840
Suqelshuyukh													67	2	3	3	
Persia: Rafsanjan 1			36										7				
	_		14							_				-	-	-	:

Columbia Columbia	Columbia Columbia	Philippine Islands, Provinces—Capiz.	.	-8	12	7							_		-	_
1	Colored Colo		4	*	12	•					H	-		a 10		
September Columbia	Colored Colo	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	-			នគ	28	800			-	-				
Shorthan Columbia	Sheepers Co Co Co Co Co Co Co Co Co Co Co Co Co		; ; ;;		$\frac{1}{11}$	i			<u> </u>							
Colored Colo	Sharita. C C 2 3 3 2 3 3 1 1 1 1 1 1 2 2 2 2 3 3 1 1 1 1			<u> </u>			H				$\frac{1}{11}$	#				
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¹ From May 3 to 25, 1931, 152 cases of cholers with 75 deaths were reported in Rafsanjan and vicinity, Karman district, Persia.

² Figures for cholers in the Philippine Islands are subject to correction.

⁸ Reports incomplete.

CHOLERA, PLAGUE, SMALLPOX, TYPHUS FEVER, AND YELLOW FEVER-Continued

PLAGUE
O indicates cases; D, deaths; P, present]

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Peru (see table below). Senegal (see table below).		•			+	-	+	+		T	İ	-	┼		-	:
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1 On July 27, 1931, 1,250 cases of plague were reported in Chiobe and Changchow, China, sinc April.

CHOLERA, PLAGUE, SMALLPOX, TYPHUS FEVER, AND YELLOW FEVER-Continued

PLAGUE—Continued [C indicates cases; D, deaths; P, present]

			≟	Indicat	es cases	C indicates cases; D. deaths; P. present	ns; F, I	resent							′			
												Week	Week ended-	Ţ				
Place		<u> </u>	Mar. 8- Apr. 4, 1931	Apr. 5- May 2, 1931	May 1931,	May31- June <i>27</i> , 1931		July, 1931	188			August, 1931	1931			Septo	September, 1931	123
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Syria: Belrut Tripolitania Tunisia: Tunis		0000	-2	16	16	=		7		-			-					
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Plague-infected rats. Orange Free State.		A OA	91	m mm	(36)								-					
Place	Mar., Apr., 1931 1931	., May,	y, June, 31 1931	16, July,	, Aug. 1 1931	.:-			Place			A	Mar., 1931	Apr., 1931	May, 1931	June, 1931	July, 1931	Aug.,
British East Africa (see also table above): Kenya. Indo-China (see also table above)C	F-41	345	245	154 484		Poru	Poru.					OA C	ω ₆₁	80	8	1011	6 6	<u> </u>
ove):		88	618		<u> </u>		Dakar					900 1111		64-	£ &	2,2	282	8 4 8
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		$\left \cdot \right $																

¹ Reports incomplete.

SMALLPOX
[C indicates cases; D, deaths; P, present]

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Nova Scotia. Ontario. Ottawa. Ottawa. Sault Sto. Marie.	1 9	-0°2 44	17	32	14	69	0	13	7		61	69	4		· -
Quebec Saskatchewan. Regina. C Canary Islands: Las Palmas. C Chile:	886-	. 8 1 6	± 24 €2	20		13	10	19		9	6	9	oo i	σο .	
Chanaral			1		•			,	<u> </u>		H	H	H	H	

1 An epidemic of smallpox was reported on May 18 with 716 cases and 314 deaths since the middle of April, 1931, in Mendez Province, Bolivis.

CHOLERA, PLAGUE, SMALLPOX, TYPHUS FEVER, AND YELLOW FEVER—Continued

SMALLPOX-Continued

[C indicates cases; D, deaths; P, present]

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									Week o	Week ended-					
Place	Mar. 8- Apr. 4, 1931	Apr. 5- May 2, 1931	May 3-30, 1931	May 31- June 27, 1931		July, 1931	1831			Aug	August, 1931		<u> </u>	September, 1931	. per
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China: Amoy. Amoy. Canton. Foochow. Hankow. How Kong. Manchuria— Harbin (see also table below). Charking. Foreigners only. Tocluding natives. Swatow. Choesen (see table below).	1117 620 101 008 1 2 2 1 2 2 0 1 2 2 1 1 2 1 1 1 1 1 1 1	911476861 97 1286 1 409188897	\$28080 \$28080 \$28080 \$28080 \$28080 \$28080	28 28 28 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 227	2 111	27.2	1	8 92	8 28	91 19	40 TO # 01	- B 8 8 8		
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CHOLERA, PLAGUE, SMALLPOX, TYPHUS FEVER, AND YELLOW FEVER-Continued

SMALLPOX—Continued

[O indicates cases; D, deaths; P, present]

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									Week	Week ended-	,				
Place	Mar. 8- Apr. 4, 1931	Apr. 5- May 2, 1931	May 3-30, 1931	May31- June 27, 1931		July, 1931	1831			γng	August, 1931	15 15		September, 1931	1 Der
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Turkey (see table below). Union of Socialist Soviet Republics (see table below). Union of South Africa: Cape Province.	Д	. А	A												
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S. S. Rotterdam at Naples from Venice. S. S. Clan McTavib at Mandle from Chittagong. C. S. S. Tall (pilgrim ship) at Suakin from Jeddah. C. S. Talodi at Suakin.	-	ÀA										-			
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	1931 Feb.,	Mar. 1831	Apr.,	May, 1931	June, 1831	July, 1931		!	Place			Jan., 1931	Feb., 1931	Mar., 1931	Apr., 1931	May, 1931	June, 1931	July, 1931
China: Harbin (see also table above)		117		13	014		Rumar Turkey	nia / M Rocial	Rumania. C Turkey C Union of Socialist Societ Reports.	at Range	OOA	282	7.8			10	1	
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CHOLERA, PLAGUE, SMALLPOX, TYPHUS FEVER, AND YELLOW FEVER-Continued

TYPHUS FEVER

C indicates cases: D. deaths: P. presentl

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1 On Feb. 27, 1931, the Director General of Public Health of Guatemals reported an unusual outbreak of typhus fever in a small village in Guatemala.

CHOLERA, PLAGUE, SMALLPOX, TYPHUS FEVER, AND YELLOW FEVER-Continued

TYPHUS FEVER-Continued

[C indicates cases; D, deaths; P, present]

YELLOW FEVER

[C indicates cases; D, deaths; P, present]

2478

							}			Week	Week ended-	1					
Place	Mar. 8 Apr. 4, 1931	Mar. 8- Apr. 5- May 3- May 31- Apr. 4, May 2, 30, June 27, 1931 1931 1931 1931	May 3- 30, 1931	May31- June 27, 1931		July, 1931	1931	-		Augn	August, 1931	31		88	September, 1931	3t, 198	_
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