STUDY OF AN OUTBREAK OF DIARRHEA IN A CONVICT CAMP NEAR SAN JUAN

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In the latter part of November 1927 we were requested by the Assistant Commissioner of Health and the Director of the Biological Laboratory of the Department of Health of Porto Rico to investigate an outbreak of acute diarrhea in a camp for convicts on the grounds of the Insular Tuberculosis Sanatorium, about five miles from San Juan. The study was undertaken as a part of an investigation of dysentery in Porto Rico which was already in progress at the School of Tropical Medicine.

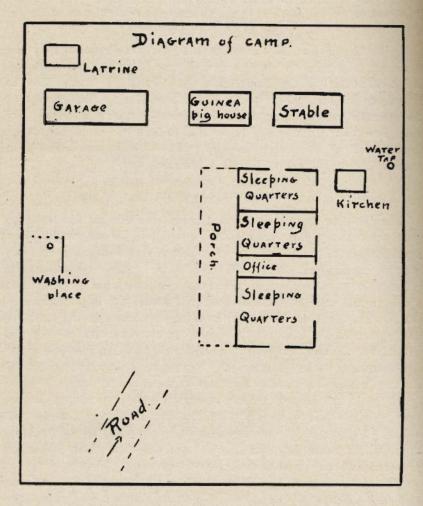
PRELIMINARY OBSERVATIONS

The camp in question contained forty convicts, all from the insular penitentiary, together with one guard and his assistant. These convicts live in three large rooms, arranged as shown in the accompanying diagram. They are supplied the same foodstuffs as the patients in the Sanatorium with the sole exception of the milk, which in the case of the convicts is condensed milk, "Milkmaid" brand. The food of the latter, however, is cooked at the camp itself by two of the convicts. Some of the convicts take care of and milk sixty-five cows that provide milk for the two hundred and eighteen inmates of the Sanatorium. The nearest living quarters of the institution is about five hundred yards from the camp.

At the time of our arrival (November 22), thirty-two of the forty convicts had been sick with diarrhea in the two weeks preceding. Neither of the two guards had been ill. The guards are supplied the same foodstuffs as the convicts but their meals are prepared and served at their homes.

The outbreak started on November 8th when twelve convicts were suddenly taken sick almost simultaneously. From that time until November 22nd, twenty more became similarly ill, and new attacks or relapses were noted in several of those first taken ill.

The following is a typical history of practically all of the cases: The patient would go to bed apparently well and during the night, generally in the early morning hours, would awake with an acute pain over the lower abdomen and a violent desire to evacuate. He would have to hurry to the latrine, and sometimes before reaching it he would defecate in his clothing. Defecation was accompanied by tenesmus and pain. The stools were liquid and contained abundant mucus. In many cases there was also blood. Stools were very



frequent, four to ten or more in twenty-four hours. The patients would at once take a large dose of Glauber's salts (sodium sulphate) and in two or three days the symptoms would pass. In practically all the cases, however, a second or third attack occurred.

The constitutional symptoms were very mild. Most of the patients kept at work except when evacuations were so frequent that

they had to remain in quarters. No fever was noticed except in one case (No. 32). All patients complained of pain on pressure on the lower abdomen.

Physical examination revealed nothing except marked tenderness over the hypogastric region and slight tenderness in both iliac regions on deep pressure. It is interesting to note that every one of the men blamed the red beans for their illness, although many of them had a second attack after omitting the beans from the diet.

With respect to the sanitary conditions of the camp we noted that there were great numbers of flies all around, that there was only one latrine (and to this flies and other insects had free access), that the kitchen was freely accessible to rats and vermin, that a stable for horses with manure and waste stood close to the camp, that there was free interchange in eating and drinking utensils among the men, and that all of the clothing of the camp was washed on the premises by one of the convicts.

It is noteworthy that during the whole month of November there were heavy rains.

METHODS OF STUDY

The general plan followed by us in the investigation of this outbreak was as follows:

 Bacteriological study of all the foods and of the fresh stools of all the occupants of the camp.

The bacteriological technique employed was as follows: The stool was cultured immediately after passage, one hour at most after collection. A portion of the stool (about 0.5 gram) was placed in plain nutrient broth pH. 7.6, and shreds of mucus removed, washed in saline and seeded on one-half of each of four plates of Endo's agar pH. 7.6, and two of Levine's cosine methylene blue agar (pH. 7.6. The other half of the plate was planted with the supernatant broth after letting it set for thirty minutes. The plates were incubated twenty-four hours at 37°C and colonies, both lactose and non-lactose fermenters, were picked from all six plates, about 15–25 colonies in each case. The cultures were studied, other media inoculated, and agglutination reactions made until the organism recovered, was classified.

In the case of the foodstuffs, samples of each were placed in tubes of nutrient bullion, placed in the incubator for a half hour, and seeding made in the way described above with the supernatant broth.

2. Study of the gross and microscopic picture of the fresh stool and of stained smears of the same.

The technique followed in our fecal examinations was as follows: In our search for *Endamoeba histolytica* only fresh stools were examined. A bit of the fecal material, preferably that containing mucus, was diluted with normal salt solution and examined under the microscope without a warm stage. If cysts were found they were identified in a second smear made with iodin as the diluting fluid.

In cases where amoebae were found smears were fixed in Schaudinn's fixing fluid and stained with iron-hematoxylin. The same procedure was followed in the final examination for cysts except that in this case permanent preparations were made from all the stools examined.

3. Hemoglobin estimation (Dare), total white and red blood cell counts and differential counts were made in eighteen of the affected men and five of the unaffected.

FINDINGS

Bacteriological findings in case of the foodstuffs and feces are given in Tables I and IV.

At the first examination made by us most of the stools were very mushy and several were diarrhoeic. Mucus was present in them only in slightly abnormal amounts. Blood was visible microscopically in a large number of them, but only in four cases was blood discernible by the naked eye. Endamoeba histolytica were found in three cases. Actively motile amoebae with ingested red blood corpuscles were never observed. In three more cases small amoebae were seen but it was impossible to identify them because cysts were not present and stained preparations could not be made from the material. Vegative forms of End. coli were found in another case.

Stools passed after the taking of a saline purge were obtainable in seven cases out of the eleven to whom the purge was administered. The three cases previously found to be infected with *End. histolytica* again showed the amoebae in their stools, but the remaining four cases proved negative.

The results obtained in our final examination made fifty-four days after the cessation of symptoms are given in Table III.

All parasitological findings, actually made in one or another of the examinations are given in Tables II and IV.

The results of the study of the blood are given in Table IV.

DISCUSSION

In our bacteriological study we failed to make anaërobic cultures of the foodstuffs and feces.

The inability to find dysentery or other allied bacilli in any of the feces, although fresh samples of these were studied, leads us to discard this group of organisms as a possible factor in the outbreak. The absence of definite constitutional disturbances, including fever, in all of the cases is also against such an etiology although many cases of bacillary dysentery do present very mild manifestations at times.

The blood picture of the cases studied did not throw any light on the problem. The high percentage of eosinophiles and the anemia found in some can be explained by the presence of helminths.

Our protozoological findings clearly show that about one-half of the diarrhea patients must have been infected with End. histolytica. This estimated percentage incidence, though high, is not unduly high for Porto Rico. Hegner (1921) found End. histolytica in twelve per cent of eighty-three stools from healthy individuals from different parts of the Island. According to his figures the estimated incidence would have been thirty-six per cent. Hill and Hill (1927) examined 269 persons from the general population and found 10.4 per cent infected. From their figures the estimated incidence would have been 31.2 per cent for the general population. This figure for the whole population necessarily means a much higher percentage for the lower, poorer classes to which our convicts belong. If in addition it is recalled that they form part of an institution, an incidence of End. histolytica above the average would be expected.

These findings, together with some theoretical consideration, we believe rule out End. histolytica as the cause of the diarrhea. From the work of Walker and Sellards (1913) it is clear that only a minority of the individuals infected with End. histolytica showsymptoms at one time or another and it also becomes apparent that there is not a definable incubation period in amoebiasis. With these two facts in mind and from what we know about the transmission of End. histolytica it is impossible to understand how the forty men could have become infected simultaneously or how eighty per cent of them should have come down with symptoms at nearly the same time.

The presence of Charcot-Leyden crystals in about fifteen per cent of the stools examined has no significance because they are known to be commonly found associated with hookworm and other helminths (Ashford, 1911).

SUMMARY AND CONCLUSIONS

The results of a bacteriological and parasitological study of an outbreak of diarrhea affecting thirty-two out of forty persons living under the same conditions in a convict camp were essentially negative, in that no bacterium or protozoa was found that could be

considered responsible for the disturbance. No organisms of the typhoid-dysentery group were demonstrated and End. histolytica was ruled out on the basis of the comparative incidence.

Since anaërobic cultures were not made, the possibility of an anaërobe as the cause must be considered.

Another possibility is a non-living chemical irritant derived from or associated with one of the foodstuffs in the common dietary.

These negative findings are considered worthy of publication largely because the frequency of intestinal disorders in Porto Rico makes it advisable to place on record all systematic studies, whether conclusive or not.

We wish to thank Dr. A. Fernós Isern, Assistant Commissioner of Health, and Dr. P. Morales Otero, Director of the Public Health Laboratory, for affording us the opportunity to study this outbreak. We are greatly indebted also to Dr. Roure, acting Director of the Tuberculosis Sanatorium, who cooperated with us in every way, placing all the facilities of the Sanatorium at our service.

LITERATURE CITED

ASHFORD, B. K. Unicinariasis in Porto Rico. Senate Document No. 808. Government Printing Office, Washington, D. C., 1911. Hegner, R. W. The Prevalence of Intestinal Protozoa and Related Organisms in Porto Rico. Jour. Amer. Med. Ass., 1921, 77: 1439-1440.

Hegner. R. Host-Parasite Relations Between Man and His intestinal Protozoa. Century Co., New York, 1927.

HILL, C. McDowell, and Hill, R. B. Infection with Protozoa and the Incidence of Diarrhea and Dysentry in Porto Rican Children of the Pre-School Age. Amer. Jour. of Hyg., 1927, 7: 134-146.

TABLE I
BACTERIOLOGIC STUDY OF FOODSTUFFS

No.	Food	Findings
14	Rice (uncooked)	Flavobacterium racemosum
15	Lard	Negative
16	Chick peas	Escherichia coli (B. coli communis) Staphylococcus albus
17	Condensed milk "Milk maid"	Negative
18	Table salt	Escherichia coli
19	Water	Negative
20	Beans (white)	Escherichia coli
21	Brown sugar	Staphylococcus albus
22	Ground coffee	Negative
23	Annato	Escherichia coli Flavobacterium racemosum
24	Beans (red)	Negative
25	Bacon	Negative
26	Cod fish	

TABLE II
RESULTS OF EXAMINATIONS OF STOOLS FOR HELMINTHS

•			Total positive	helminths		Hookworm	Trichnris	trletiura	Stronovioldos	stereolaris	Sehistosoma	mansoni	Ascaris lum- bricoldes		
Cases studied	Number	Per cent	Number	Per cent	Number	Per cent	Number	Per cent	Number	Per cent	Number	Per cent	Number	Per cent	
With diarrhea Not affected		76.1 25.3	27 7	84.3 70	22 6	68.7 60	15 2	46.8 20	3	9.3 10	1	3.1	1 0	3.1	
Total	42		34	80.9	28	66.6	17	40.4	4	9.5	2	4.7	1	2.3	

TABLE III

RESULTS OF ONE EXAMINATION FOR INTESTINAL PROTOZOA
FIFTY-FOUR DAYS AFTER CESSATION OF SYMPTOMS

	ases		Endamoeba			Endamoeba			Endollmax	папа			Iodamoeba, williamsi		Dientamoeba	ILERIIIS	Trichomonas	Bominis	Giardia	lamblia
Cases	Number of cases	Number	Per cent	Estimated Per cent	Number	Per cent	Estimated Per cent	Number		Per cent	Estimated Per cent	Number	Per cent	Estimated Per cent	Number	Per cent	Number	Per cent	Number	Per cent
With diarrhea. Not affected	27 6	5	18.	55.5	4	14.8 16.6	44. 49.	1 8	3 11	1.1	33.8	1 0	3.6	10.8	1 0	3.6	1 0	3.6	1 0	3.6
Total	33	5	15.1	45.3	5	15.1	15.	3	3	9	27	1	3	9	1	3	1	3	1	8

^{*} This is the first case of infection with *Dientamoeba jragilis* reported for Porto Rico. Dr. W. H. Taliaferro, professor of protozoology in the University of Chicago, at present visiting lecturer to the School of Tropical Medicine, has examined the slides and corroborated our diagnosis of this rare intestinal amoeba of man.

TABLE IV
CLINICAL AND LABORATORY FINDINGS IN ALL CASES

	Gros	s Ex	am.							e at			Ble	boo		1			
				cells			en		attacks	of residence		uko. mm.	b c	Dir	Tere	ntial	col	int	Organisms isolated from
Case number	Diarrhea	Blood	Mueus	Red blood ce	Leukocytes	Epithelium	Charcot-Leyden crystals			Months of res	% hemoglobin (Dare)	Number of lenko- cytes per c. mm	Millions of r	Lymph \$	Polys &	Eosino 9	Transg	Other %	the feces
1e	+	_	+		+	+		Amoeba (?)	2	14	82	7520	3.720	33	64	1	1	1	Escherichia coli *
2	+	_	+	+	+	+	_	Endamoeba histolytica.	3	7	94	7800	5.624	29	60	9	2		Escherichia coli
3	+	_	+	+	+	+	_	Trichuris trichiura	3	5	81	9230	3.910	30	58	6	2	4	Escherichia coli
4	+	-	+	+	1	+	-	{ Trichuris trichinra }	2	11	58	8320	2.480	30	55	8	4	3	Escherichia coli
5	+	+	+	+	+	-	-	Hookworm	2	11	72	6940	4.610	25	58	16	0.2	0.7	Escherichia coli
6c	+	+	+	+	-	+		Schistosoma mansoni	3	6	79	8040	3.800	34	57	5.6	1.5	1.5	Escherichia coli
7	+	_	+	_	-	+	-	Ascaris lumbricoides	2	1	62	7890	2.810	40	48	11	1		Escherichia communio
8	+	+	+	+	+	+	-	(Endolimax nana)	2	19									Escherichia coli
9	+	_	+	+	_	1+	-	(Trichuris trichiura) Endamoeba histolytica	3	4	85	7170	4.736	37	50	9	2	2	Escherichia coli
0	+	_	+	+	_	-	-	Amoeba (?)	3	20	72	8740	4.240	32	55	10	2	1	Escherichia communio
11	+		+					Trichuris trichiura	2	2	88	7500	4.060	194	70	2	9	2	(Escherichia communi Aerobacter aerogenes

12	+	-	+	-	-	-	1+1	Hookworm 1 20		.1	1	١	1	Escherichia coli *
						1		(Endamoeba histolytica.)				-37	1311	
13	+	-	+	+	-	+	-							Escherichia coli
27	+	-	+	+	+	+		(Trichuris trichiura)						{ Escherichia coli } Aerobacter aerogenes
28	+	-	+	+	+	+	+	12-1	00 3.760 2	0 69	9	1	1	Serichia coli Aerobacter aerogenes
· 29	+	-	+	+	+	+	-	Trichomonas hominis \ 4 1 83 9.90	00 2.140 4	5 47	6	2		Escherichia coli Aerobacter aerogenes
30	+	-	+	+	+	+	-	Hookworm	40 4.810 2	6 60	13	1		Escherichia coli
31	+	-	-	+	+	+	-	(Hookworm)						Escherichia communior Aerobacter cloacae
32w	+	-	+	+	+	+	+	Hookworm	50 2.46 2	1 56	22	0.6	0.7	Escherichia coli Aerobacter aerogenes Staphylococcus albus
33	-	-	-	-	-	-	-	Dientamoeba fragilis 0 18 86 5.80 Trichuris trichiura	00 4.404 2	8 66	1	3	2	Escherichia coli
34	+	-	_	-	-	+		Endamoeba histolytica. Endolimax nana } 2 21 81 5.81 Trichuris trichiura	10 4.312 2	8 69	1	1	1	Escherichia coli
35	-	-		-	-	+	_	Endamoeba coli 2 7						Escherichia coli
36	-	-	_	_	-	-	-	Trichuris trichiura 0 9 85 11.4 Hookworm 0 9 85 11.4 Strongyloides stercolaris	00 4.386 2	2 53	23	2		Escherichia coli
37	-	-	_	-	-	-	_	Hookworm			.,			Escherichia coli Proteus valeriae
39	+	+	+	+	+	+	-	$ \left\{ \begin{array}{ll} Endamoeba\ coli & \dots \\ Giardia\ lamblia & \dots \\ Hookworm, & \dots \end{array} \right\} 2 20 95 4.30 $	00 4.120 3	0 55	12	2	1	Aerobacter aerogenes Escherichia coli
40	-	I —	-	-	-	-	-	Hookworm 0 19			١	١		Escherichia communior

^{*} Escherichia coli has been recently adopted as the name for B. coli communis.

CLINICAL AND LABORATORY FINDINGS IN ALL CASES-Continued

	Gross Exam								Stools microscopic	Jo	e at			Blo	ood					
					cells			den.		attack	sidenc	_	uko- mm.	rbe	Di	fere	ntial	col	unt	
Case number	Diarrhae	Dinimen	Blood	Muens	Red blood ce	Leucocytes	Epithelium	Charcot-Leyden, cristals	Parasites or ova	Number of a	Months of residence	% hemoglobin (Dare)	Number of leuko- cytes per c mm.	Millions of r	Lymph #	Polys \$	Eosino 3	Trans \$	Other %	Organisms isolated from the feces
41 42	-	-	-	_	-	_	_	+	Strongyloides stercolaris	1	11			,.						Escherichia coli Escherichia coli
44		-	_	+	+	+	+		Hookworm	1	15	65	8.200	2.040	23	56	19	2	::	Escherichia communior
46	4		_	(8)	+	+	_	4	Giardia lamblia	1	7	92	9,400	3.240	24	66	5	2	2) Escherichia coli
47		-	_	_	_	_	_	-	Hookworm.	1	32									Aerobacter aerogenes Secherichia coli Aerobacter aerogenes
48	-	-	-		-	-	-	-	Strongyloides stereolaris	1	10									Escherichia coli
49	-	-	-	_	-		-	_	Endamoeba histolytica. Hookworm	2	6									Escherichia coli
50	-	-	-	-	-	1	-	-	\ Hookworm\ Schistosoma mansoni\	0	48	74	8.000	4.030	26	67	7			Escherichia coli
51 52		-	-	-	-	_	-	-	Iodamoeba williamsi (Endamoeba coli	0	10									Escherichia communior Staphylococcus albus
53	-	-	-	_	_	-	_		Hookworm	2	10			· · · · · ·						Escherichia coli Aerobacter nerogenes Escherichia coli
54	-	-	-	-	-	_		-		0	20	91	8.100	4.290	27	66	1	3	3	Escherichia coli
55 56		-	-	-	_	_	-	_	Hookworm	0	9									Escherichia communior Escherichia coli
58	-	-		_	_	_	_	-	Hookworm,	0	36	1::								Escherichia coli
59	-		-	-	-	+	+	+-	Hookworm	0	4	89	6.930	4.110	37	57	9	13		Escherichia coli