

NEUTRALIZATION TESTS IN POLIOMYELITIS. SERA TAKEN
DURING THE ACUTE AND CONVALESCENT STAGES OF THE
DISEASE AND TESTED WITH A PASSAGE VIRUS AND A
STRAIN ISOLATED DURING THE 1935 NEW YORK CITY
OUTBREAK ¹

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(Received for publication January 9, 1937)

It has generally been accepted that recovery from paralytic poliomyelitis usually results in the appearance of antibodies in the blood serum (1 to 10). The neutralizing property of human convalescent sera from paralytic patients was first demonstrated in 1910 by Netter and Levaditi (1) and by Flexner and Lewis (2). Since then a number of investigators have confirmed this find-

recorded by several of the early workers. Recently, however, the absence of neutralizing power in convalescent sera has been noted more frequently (16 to 19). The majority of the neutralization tests were performed upon sera obtained from patients years after the onset of the disease; a smaller number were carried out on sera collected within one year; a few were secured

TABLE I
Summary of neutralization tests in paralytic cases compiled from the literature

Investigators	Within one week of onset			Within one year of onset			More than one year after onset		
	Number of cases tested	Number neutralized	Number failed to neutralize	Number of cases tested	Number neutralized	Number failed to neutralize	Number of cases tested	Number neutralized	Number failed to neutralize
Netter and Levaditi (1).....				4	4	0	2	1	1
Peabody, Draper and Dochez (4).....				1	1	0	3	2	1
Anderson and Frost (3).....				1	1	0			
Römer (5).....				2	2	0	1	1	0
Kling and Levaditi (6).....	2	2	0	2	2	0			
Flexner and Amoss (7).....	1	1	0						
Nuzum (8).....				1*	0	1			
Aycock and Kramer (9).....				1	0	1	16	15	1
Fairbrother and Brown (10).....				1	1	0			
Howitt (12).....							20	9	11
Stokes et al. (11).....							2	1	1
Harmon and Harkins (15).....	3	1	2	5	4	1			
Shaughnessy, Harmon and Gordon (16)				4	2	2	10	7	3
Schultz and Gebhardt (17).....							4	2	2
Jungeblut and Smith (18).....							26	15	11
Paul and Trask (19).....				7	1	6			
Total.....	6	4	2	29	18	11	84	53	31

* Recently recovered.

ing (Table I). This tabulation does not include tests performed on pooled convalescent sera or on sera of patients who had received serum therapy (11 to 15). An occasional failure of convalescent serum to neutralize poliomyelitis virus was

during the first week of illness. The proportion of patients showing neutralizing substances was approximately the same, regardless of the time after the onset of the disease that the sera were collected (Table I). This would suggest that the neutralizing substances either developed shortly after the onset of the disease or were present at the time of the infection. The former idea was

¹This work was made possible by grants from the Rockefeller Foundation and the President's Birthday Commission for Infantile Paralysis Research.

supported by Kling and Levaditi (6) who found neutralizing substances three and five days after the onset of the disease, and by Flexner and Amoss (7) who reported a similar finding at six days. The latter possibility was favored by Harmon and coworkers (14) who found neutralizing substances present in the serum of a patient before the onset of paralysis.

The higher incidence of neutralizing substances found in the convalescent sera by the early workers might have been due to chance in dealing with small numbers or to differences in technique of performing the tests. The age of the patient and the virulence of the virus must also be considered.

Very few studies have been carried out on sera from non-paralytic patients. Peabody, Draper and Dochez (4) found no neutralizing power in the serum of a child 7 days after the onset of the disease, while that of another child neutralized at 21 days. Paul and Trask (19) reported the case of a 6 year old non-paralytic child whose serum failed to neutralize both in the acute stage and 1 year later. Harmon and Harkins (15) demonstrated the presence of neutralizing substances in the serum from an 11 year old child obtained 5 days and again 5 months after the onset of the disease. Neutralization tests have also been recorded by several other investigators (1, 3, 5, 6) upon blood from patients about whom insufficient clinical data were given to determine whether they could be classified as non-paralytic or abortive cases of poliomyelitis.

Neutralizing substances have frequently been reported in the sera of so-called normal individuals (9, 16, 20). The incidence of these antibodies appears to increase with age (9, 16). Various workers (9, 16, 20) have found that the sera of at least 50 per cent of normal adults have neutralizing power. Paul and Trask (21) in a review of reports on neutralization tests carried out since 1929, pointed out that the neutralizing power of normal sera exceeded that of convalescent sera in all age groups.

All of the above recorded tests were performed with a monkey passage virus. In addition, Paul and Trask (19) tested 7 sera with a recently isolated human strain of virus, as well as with a passage strain. They found that neutralizing substances were present in 6 of the 7 sera tested

with the former strain as compared to one with the latter. Likewise, Howitt (22) found that 7 convalescent sera neutralized a recently isolated strain, while only 4 of them neutralized a passage virus.

In spite of so much contradictory evidence, the idea still prevails that recovery from an attack of poliomyelitis results in the development of antibodies. It, therefore, seemed advisable to investigate the matter further by tests on a sufficiently large number of sera taken at frequent intervals following the onset of the disease.

The purpose of this work was: (1) To test for the presence or absence of neutralizing substances in the sera of both paralytic and non-paralytic individuals of different ages in the acute and convalescent stages of the disease; (2) To determine whether poliomyelitis can develop in the presence of protective substances; (3) To determine whether paralytic or non-paralytic cases with no demonstrable protective substances in the acute stage of the disease developed them within 12 to 16 months after the onset; (4) To carry out the above tests with the F1 passage virus and with a strain of virus isolated during the summer of 1935.

METHODS

Sera were obtained from patients who were admitted to the Willard Parker Hospital for poliomyelitis during the summer and fall of 1935, when over 2,000 cases were reported in New York City. In each instance the diagnosis was established by clinical findings and examination of spinal fluids. Only those patients who had a definite loss of muscle function were classified as paralytic. Non-paralytic cases were those who had an acute onset with meningeal involvement and pleocytosis of the spinal fluid. Every case was seen by one of us and a careful follow-up of the non-paralytic and the paralytic patients was carried out. In this way, it was possible to eliminate from the non-paralytic group those patients who subsequently developed muscle weakness or paralysis. We also correlated, in the paralytics, the degree of muscle recovery with the results of the neutralization tests.

The work was begun in July 1935, and the patients were bled at short intervals during the following 9 months. A number were also bled 12 to 16 months after the onset. The majority of the sera were tested within a few weeks after collection. When comparative tests between early and later bleedings were made, the sera, which had been stored for 6 to 7 months, were again tested. All specimens were cultured before use and were kept at 2° to 4° C.

Passage virus and preparation of suspension. The passage virus (F1 strain) was ground without abrasive,

suspended in an equal amount of glycerine (Kahlbaum) by weight, and kept frozen. Small portions were removed, ground with sand, and to each gram was added 10 cc. of distilled water to make a 10 per cent suspension. The suspension was centrifuged lightly, and the resulting supernatant was referred to as a 5 per cent cord suspension because half of the gram of material mixed with the water was glycerine. Five different batches of infectious cords, each consisting of 4 or 5 spinal cords obtained from monkeys at the height of paralysis, were used. Four of the batches were used from July to November 1935, and the fifth from then until the completion of the work in October 1936. The virus, stored in this way, maintained its potency at a fairly uniform level of infectivity, as indicated by the results of repeated tests carried out with a batch over a period of 12 months.

The infectivity of each batch of virus was determined by making serial dilutions of the 5 per cent suspension with distilled water. This was accomplished by the addition of 0.5 to 1.0 cc. of virus suspension to the required amount of diluent, and then after being shaken for several minutes, 0.13 cc. was measured off with a 0.2 cc. pipette. Usually, 0.13 cc. of a 5 per cent suspension diluted 40 to 80 times and added to 0.6 cc. of saline or normal monkey serum resulted in infection.

Recently isolated strain. The virus was obtained from the nasal secretions of an acute bulbar case of poliomyelitis on the 9th day of illness. The washings were passed through a Seitz filter and concentrated *in vacuo* to approximately 5 cc. This was then inoculated intracerebrally and intraperitoneally into a monkey. For the tests described in this paper a single cord from the second passage was used, being kept in glycerine at freezing temperature. In making up the virus suspension, small pieces from 6 to 8 segments were used as previously described (23). The infectivity of the cord was sufficient to produce a rapid and complete paralysis in the usual incubation period when 0.13 cc. of a 5 per cent suspension diluted 20 to 40 times was injected. The virulence of this strain was, therefore, comparable to that of the passage virus.

Neutralization test. The neutralization or protection test was carried out as follows: 0.13 cc. of a 1:10 or 1:2 dilution of 5 per cent virus was added to 0.6 cc. of serum; the proportion was 0.1 cc. of diluted virus suspension to 0.45 cc. of serum. The mixtures were incubated for 2 hours at 37° C. and then kept in the ice box for approximately 2 hours. Of this mixture 0.5 to 0.6 cc. was inoculated into the frontal lobe of a monkey. A positive control, consisting of human serum known to have protective substances, was used in each experiment. Likewise, a negative control, consisting of normal monkey serum without protective properties, was included. When the serum failed to protect, the animal became paralyzed. With few exceptions, once paralysis set in, the animals became prostrate. When the paralysis was definite but failed to involve all extremities, it was referred to as "incomplete paralysis."

The usual minimal dose which was used to test the sera in this work was 0.13 cc. of a 1:10 dilution of 5 per

cent virus suspension. If the serum protected, a 1:2 dilution of the virus suspension was used in the next test. Protective substances were considered present only when the animal survived the greater dose. A more concentrated virus suspension was used to offset the irregularities which can occur in carrying out tests with small amounts of virus. When irregularities occurred, several retests were made and the average result recorded. Some of the sera were tested with as many as 3 different batches of virus with fairly consistent results.

Macacus rhesus monkeys weighing 2½ to 4 kilos were used. Those surviving were not used again within the month. After an animal was used 3 times, it was injected with 0.13 cc. of a 1:10 or 1:2 dilution of a 5 per cent virus suspension added to 0.6 cc. of saline or normal monkey serum to determine whether the protection tests carried out upon the monkey were valid. This was necessary because an occasional monkey is naturally resistant. Daily temperatures were taken for at least two weeks after inoculation.

EXPERIMENTAL

The sera of 82 paralytic, 32 non-paralytic and 3 encephalitic cases were tested in the acute stage of the disease, that is, within a week after the onset. Many of the sera were retested at intervals during convalescence. Thirty-three experiments were carried out. The following 9 were selected as typical protocols.

Neutralization tests with sera obtained in the acute stages of the disease

Experiment 1. The sera of 16 paralytic and 13 non-paralytic patients ranging from 5 months to 28 years of age were tested. It was found (Table II) that the sera of 6 of 7 paralytics 5 years and under and 2 non-paralytics in the same age group, failed to neutralize a 1:10 dilution of a 5 per cent virus suspension. In the group over 5 years the sera of 8 of 9 paralytics failed to protect; all these were tested with a 1:2 dilution of virus suspension. On the other hand, the sera of 9 of 11 non-paralytics over 5 years of age neutralized either a 1:10 or a 1:2 dilution; seven of them were tested against the 1:2 dilution of virus.

Experiment 2. Certain sera of the first and other experiments which had given protection with a given test dose of virus, were retested with larger doses. Sera that previously had neutralized 0.13 cc. of a 1:10 dilution were retested with a 1:2 dilution of a 5 per cent virus suspension; and those that had neutralized the latter amount

TABLE II
Summary of neutralization tests with sera taken in the first week of the disease

Patient's initials	Age	Type of case	Time after onset	Dilution of 5 per cent virus	Amount of virus	Amount of serum	Number of monkey	Results
Negative control	Normal monkey serum		days	1 : 10	cc. 0.13	cc. 0.60	J 127	Paralyzed 7 days
Negative control	Normal monkey serum			1 : 2	0.13	0.60	J 35	Paralyzed 7 days
Positive control	Normal adult human serum (M.B.)			*	0.20	0.45	J 1219	No paralysis
Positive control	Normal adult human serum (M.B.)			*	0.45	0.45	617	Paralyzed 11 days
S. S.	5 mos.	Paralytic	7	1 : 10	0.13	0.60	J 92	Paralyzed 10 days
D. O'B.	16 mos.	Paralytic	3	1 : 10	0.13	0.60	J 107	Paralyzed 6 days
	years							
H. D.	2	Paralytic	4	1 : 10	0.13	0.60	J 124	No paralysis
R. M.	4	Paralytic	5	1 : 10	0.13	0.60	J 108	Paralyzed 9 days
V. M.	4	Paralytic	3	1 : 10	0.13	0.60	J 102	Paralyzed 9 days
G. C.	5	Paralytic	5	1 : 10	0.13	0.60	J 112	Paralyzed 6 days
P. B.	5	Paralytic	3	1 : 10	0.13	0.60	J 111	Paralyzed 14 days
C. M.	6	Paralytic	5	1 : 2	0.13	0.60	J 122	Paralyzed 8 days
R. A.	7	Paralytic	2	1 : 10	0.13	0.60	J 120	No paralysis
R. A.	7	Paralytic	2	1 : 2	0.13	0.60	J 119	No paralysis
J. B.	8	Paralytic	1	1 : 2	0.13	0.60	J 105	Paralyzed 7 days
M. G.	8	Paralytic	3	1 : 10	0.13	0.60	J 121	Paralyzed 15 days
M. G.	8	Paralytic	3	1 : 2	0.13	0.60	J 53	Paralyzed 8 days
M. R.	9	Paralytic	4	1 : 2	0.13	0.60	J 103	Paralyzed 16 days
T. McT.	15	Paralytic	6	1 : 2	0.13	0.60	J 90	Paralyzed 6 days
J. M.	16	Paralytic	4	1 : 2	0.13	0.60	J 91	Paralyzed 6 days
E. P.	21	Paralytic	4	1 : 2	0.13	0.60	J 97	Paralyzed 6 days
V. C.	28	Paralytic	1	1 : 2	0.13	0.60	J 93	Paralyzed 6 days
P. M.	3	Non-paralytic	7	1 : 10	0.13	0.60	J 100	Paralyzed 7 days
H. B.	5	Non-paralytic	3	1 : 10	0.13	0.60	J 113	Paralyzed 4 days
C. E.	6	Non-paralytic	3	1 : 10	0.13	0.60	J 114	No paralysis
L. W.	8	Non-paralytic	2	1 : 10	0.13	0.60	J 123	No paralysis
L. W.	8	Non-paralytic	2	1 : 2	0.13	0.60	J 70	No paralysis
M. E.	9	Non-paralytic	3	1 : 2	0.13	0.60	J 96	No paralysis
H. E.	10	Non-paralytic	3	1 : 10	0.13	0.60	J 115	No paralysis
J. B.	13	Non-paralytic	1	1 : 2	0.13	0.60	J 98	No paralysis
C. S.	15	Non-paralytic	2	1 : 2	0.13	0.60	J 99	Paralyzed 9 days
A. P.	16	Non-paralytic	2	1 : 2	0.13	0.60	J 95	No paralysis
R. J.	16	Non-paralytic	4	1 : 2	0.13	0.60	J 106	No paralysis
A. C.	16	Non-paralytic	6	1 : 2	0.13	0.60	J 128	Paralyzed 16 days
B. M.	22	Non-paralytic	2	1 : 2	0.13	0.60	J 104	No paralysis
M. W.	27	Non-paralytic	4	1 : 2	0.13	0.60	J 94	No paralysis

* Undiluted.

were tested with 0.13 cc. or more of a 5 per cent suspension. The results (Table III) showed that the sera of 5 of 8 non-paralytically and two paralytically protected when the larger quantity of virus was used.

Summary of results of all tests on sera collected in the acute stage of the disease

The results of all of the first-week bleedings (Table IV) indicated that protective substances were present in 14 of 82 paralytically during the acute stage of the disease. In the non-paralytic group, on the other hand, they were present in

18 of 32 sera. When protective substances were present, they were detected as early as 1 and 2 days after the onset of the disease both in paralytic and non-paralytic cases. Four of the 82 patients were tested in the preparalytic stage. Two of them were found to have protective substances.

Protective substances in normal urban individuals

It was decided to test sera of a group of so-called normal residents of New York City between 11 and 25 years of age to determine whether the virus and technique used in the

TABLE III

Neutralization tests in the first week of the disease. (These sera had previously neutralized a smaller dose of virus)

Patient's initials	Age	Type of case	Time after onset	Dilution of 5 per cent virus	Amount of virus	Amount of serum	Number of monkey	Results
Negative control	Normal monkey serum			1 : 10	0.13	0.60	J 207	Paralyzed 7 days
Negative control	Normal monkey serum			1 : 2	0.13	0.60	J 126	Paralyzed 6 days
Positive control	Normal adult human serum (M.B.)			*	0.20	0.45	J 208	Partial paralysis
Positive control	Normal adult human serum (M.B.)			1 : 2	0.13	0.60	J 215	No paralysis
R. G.	3	Non-paralytic	4	*	0.20	0.45	J 239	No paralysis
C. E.	6	Non-paralytic	3	1 : 2	0.13	0.45	J 234	Paralyzed 15 days
L. W.	8	Non-paralytic	2	*	0.20	0.45	J	No paralysis
L. W.	8	Non-paralytic	2	*	0.45	0.45	J 229	No paralysis
J. B.	13	Non-paralytic	1	*	0.20	0.45	N 17	No paralysis
R. J.	16	Non-paralytic	4	*	0.20	0.45	N 19	Paralyzed 7 days
A. P.	16	Non-paralytic	2	*	0.20	0.45	J 119	No paralysis
A. P.	16	Non-paralytic	2	*	0.45	0.45	J 211	No paralysis
B. M.	22	Non-paralytic	2	*	0.20	0.45	J 237	No paralysis
M. W.	27	Non-paralytic	4	*	0.20	0.45	N 8	Paralyzed 9 days
H. D.	2	Paralytic	5	1 : 2	0.13	0.60	J 220	No paralysis
L. E.	8	Paralytic	3	1 : 2	0.13	0.60	J 216	No paralysis

* Undiluted.

experiments produced approximately the same proportion of neutralizations reported by others (9, 16, 21). Moreover, this would enable us also to compare the results of these tests with those of paralytic and non-paralytic patients in the acute stage of the disease.

TABLE IV

Summary of neutralization tests with sera taken in the first week of the disease

Age group	Paralytics			Non-paralytics			Encephalitic		
	Number tested	Sera neutralized	Sera failed to neutralize	Number tested	Sera neutralized	Sera failed to neutralize	Number tested	Sera neutralized	Sera failed to neutralize
years									
1-5	26	2	24	6	1	5			
6-10	25	4	21	14	10	4	3	1	2
11-17 . . .	19	4	15	10	5	5			
Adults . . .	12	4	8	2	2	0			
Totals . . .	82	14	68	32	18	14	3	1	2

Experiment 3. The sera of 18 individuals from 11 to 25 years of age were tested by the use of 0.6 cc. of serum mixed with 0.13 cc. of a 2½ per cent (a 1 : 2 dilution of 5 per cent) virus suspension. Protection occurred with 9 of the 18 sera. The proportion of positive neutralizations in the group approximates that found by

other workers for normal urban adolescents and adults. Protective substances were found in only 8 of the 31 sera from paralytic patients over 10 years of age during the acute stage (Table IV). In the non-paralytic patients of the same age group, 7 of the 12 sera protected. Thus, the incidence of protective substances during the acute stage of the disease in the paralytic group was lower than that in the normal urban residents, while in the non-paralytic group it was approximately the same.

Protection tests with sera secured during the first 9 months of convalescence

Repeated tests were carried out upon the sera of 44 paralytic, 13 non-paralytic and 2 encephalitic patients at various intervals from two weeks to nine months after the onset of the disease. The sera of a few of these individuals had protected in the first week of their illness, but the majority had failed to do so.

Protection tests with convalescent sera from individuals whose sera failed to protect in the first week of the disease

The following three experiments are typical of the results obtained.

Experiment 4. The sera of 2 paralytic cases obtained 6 weeks and 4 months, respectively, after the onset were tested together with those obtained in the acute stage of the disease. Similar tests were carried out with specimens of serum from a non-paralytic case and also with specimens obtained from an encephalitic case 3

child, G. C., specimens obtained in the 1st, 3d, and 12th weeks were tested against graded dilutions of virus (1:10 to 1:80). At the same time specimens from 15 paralytics and non-paralytics obtained 1 to 8 months after the onset were tested in the usual way against 1:10 and 1:2 dilutions of 5 per cent virus. From 3 of

TABLE V

Neutralization tests with convalescent sera of individuals whose sera failed to neutralize in the first week

Patient's initials	Age	Type of case	Time after onset	Dilution of 5 per cent virus	Amount of virus	Amount of serum	Number of monkey	Results
	<i>years</i>				<i>cc.</i>	<i>cc.</i>		
Negative control		Normal monkey serum		1 : 10	0.13	0.60	J 75	Paralyzed 8 days
Negative control		Normal monkey serum		1 : 5	0.13	0.60	J 63	Paralyzed 6 days
Negative control		Normal monkey serum		1 : 2	0.13	0.60	J 66	Paralyzed 8 days
Positive control		Normal adult human serum (M.B.)		1 : 10	0.13	0.60	J 64	No paralysis
Positive control		Normal adult human serum (M.B.)		1 : 5	0.13	0.60	J 74	No paralysis
Positive control		Normal adult human serum (M.B.)		1 : 2	0.13	0.60	145	No paralysis
G. M.	13	Paralytic	2 days	1 : 10	0.13	0.60	J 79	Paralyzed 6 days
G. M.	13	Paralytic	2 days	1 : 5	0.13	0.60	J 88	Prostrate 6 days
G. M.	13	Paralytic	2 days	1 : 2	0.13	0.60	226	Paralyzed 8 days
G. M.	13	Paralytic	4 months	1 : 10	0.13	0.60	J 87	Paralyzed 7 days
G. M.	13	Paralytic	4 months	1 : 5	0.13	0.60	J 84	Paralyzed 7 days
G. M.	13	Paralytic	4 months	1 : 2	0.13	0.60	J 56	Paralyzed 7 days
R. H.	6	Paralytic	2 days	1 : 10	0.13	0.60	J 83	Paralyzed 9 days
R. H.	6	Paralytic	2 days	1 : 5	0.13	0.60	J 77	Paralyzed 5 days
R. H.	6	Paralytic	2 days	1 : 2	0.13	0.60	J 28	Paralyzed 7 days
R. H.	6	Paralytic	6 weeks	1 : 10	0.13	0.60	J 85	Paralyzed 6 days
R. H.	6	Paralytic	6 weeks	1 : 5	0.13	0.60	J 81	Paralyzed 9 days
R. H.	6	Paralytic	6 weeks	1 : 2	0.13	0.60	J 1	Paralyzed 9 days
T. S.	6	Non-paralytic	3 days	1 : 10	0.13	0.60	J 80	Paralyzed 6 days
T. S.	6	Non-paralytic	3 days	1 : 5	0.13	0.60	J 76	Paralyzed 8 days
T. S.	6	Non-paralytic	3 days	1 : 2	0.13	0.60	J 57	Paralyzed 6 days
T. S.	6	Non-paralytic	3 weeks	1 : 10	0.13	0.60	J 69	Paralyzed 12 days
T. S.	6	Non-paralytic	3 weeks	1 : 5	0.13	0.60	J 78	Paralyzed 7 days
T. S.	6	Non-paralytic	3 weeks	1 : 2	0.13	0.60	211	Paralyzed 6 days
M. F.	7	Encephalitic	19 days	1 : 10	0.13	0.60	J 65	Paralyzed 7 days
M. F.	7	Encephalitic	19 days	1 : 5	0.13	0.60	J 67	Paralyzed 12 days
M. F.	7	Encephalitic	19 days	1 : 2	0.13	0.60	J 32	Paralyzed 7 days
M. F.	7	Encephalitic	5 months	1 : 10	0.13	0.60	J 89	Prostrate 6 days
M. F.	7	Encephalitic	5 months	1 : 5	0.13	0.60	J 86	Paralyzed 8 days
M. F.	7	Encephalitic	5 months	1 : 2	0.13	0.60	J 18	Paralyzed 6 days

weeks and 5 months, respectively, after the onset. Each serum was tested with three different dilutions of virus. The results of these tests (Table V) failed to show any evidence that protective substances had developed in these four convalescent sera.

Experiment 5. To determine whether any evidence whatsoever of protective power could be detected in convalescent sera from a paralytic

these individuals, the specimens collected during the first week and 3 subsequent specimens were tested. The results (Table VI) of this experiment agreed with those of the previous one, in that none of the 12 sera from paralytics had shown protective power when tested from 2 to 8 months after the onset. Likewise, the 3 sera from non-paralytics failed to neutralize the virus.

Unfortunately, the positive control serum failed

TABLE VI

Neutralization tests with convalescent sera from individuals whose sera failed to neutralize in the first week

Patient's initials	Age	Type of case	Time after onset	Dilution of 5 per cent virus	Amount of virus	Amount of serum	Number of monkey	Results
Negative control	years	Normal monkey serum		1 : 2	cc. 0.13	cc. 0.60	J 197	Paralyzed 6 days
Negative control		Normal monkey serum		1 : 10	0.13	0.60	J 199	Paralyzed 8 days
Negative control		Normal monkey serum		1 : 20	0.13	0.60	J 198	Paralyzed 6 days
Negative control		Normal monkey serum		1 : 40	0.13	0.60	J 196	Paralyzed 7 days
Negative control		Normal monkey serum		1 : 80	0.13	0.60	J 194	Paralyzed 6 days
Positive control		Normal adult human serum (M.B.)		*	0.20	0.45	J 178	Paralyzed 15 days
Positive control		Normal adult human serum (M.B.)		*	0.45	0.45	J 64	Partial paralysis 15 days
Positive control		Normal adult human serum (M.B.)		*	0.45	0.45†	J 74	Paralyzed 10 days
G. C.	5	Paralytic	5 days	1 : 10	0.13	0.60	J 186	Paralyzed 7 days
G. C.	5	Paralytic	5 days	1 : 20	0.13	0.60	J 187	Paralyzed 7 days
G. C.	5	Paralytic	5 days	1 : 80	0.13	0.60	J 185	Paralyzed 9 days
G. C.	5	Paralytic	19 days	1 : 10	0.13	0.60	J 191	Paralyzed 7 days
G. C.	5	Paralytic	19 days	1 : 20	0.13	0.60	J 190	Paralyzed 36 days
G. C.	5	Paralytic	19 days	1 : 40	0.13	0.60	J 189	Paralyzed 9 days
G. C.	5	Paralytic	19 days	1 : 80	0.13	0.60	J 188	Paralyzed 8 days
G. C.	5	Paralytic	3 months	1 : 10	0.13	0.60	J 192	Paralyzed 7 days
G. C.	5	Paralytic	3 months	1 : 20	0.13	0.60	J 193	Paralyzed 9 days
G. C.	5	Paralytic	3 months	1 : 40	0.13	0.60	J 175	Paralyzed 14 days
G. C.	5	Paralytic	3 months	1 : 80	0.13	0.60	B.U.	Died 8th day—Intercurrent infection
R. M.	4	Paralytic	15 days	1 : 10	0.13	0.60	J 170	Paralyzed 9 days
R. M.	4	Paralytic	26 days	1 : 10	0.13	0.60	J 164	Paralyzed 7 days
R. M.	4	Paralytic	3½ months	1 : 10	0.13	0.60	J 163	Paralyzed 4 days
V. M.†	4	Paralytic	2 days	1 : 10	0.13	0.60	J 173	Paralyzed 7 days
V. M.	4	Paralytic	19 days	1 : 10	0.13	0.60	J 174	Paralyzed 7 days
V. M.	4	Paralytic	4 months	1 : 10	0.13	0.60	J 168	Paralyzed 6 days
V. M.	4	Paralytic	7 months	1 : 10	0.13	0.60	J 167	Paralyzed 6 days
J. S.	3	Paralytic	8 months	1 : 10	0.13	0.60	J 171	Paralyzed 7 days
J. C.	8	Paralytic	8 months	1 : 10	0.13	0.60	J 172	Paralyzed 7 days
E. S.	5	Paralytic	2 months	1 : 2	0.13	0.60	N 12	Paralyzed 11 days
E. S.	5	Paralytic	6 months	1 : 2	0.13	0.60	J 158	Paralyzed 7 days
D. F.	17	Paralytic	7 weeks	1 : 2	0.13	0.60	J 182	Paralyzed 6 days
D. F.	17	Paralytic	7½ months	1 : 2	0.13	0.60	J 2	Paralyzed 9 days
K. G.	31	Paralytic	3 months	1 : 2	0.13	0.60	J 156	Paralyzed 8 days
K. G.	31	Paralytic	3 months	*	0.10	0.60	R 68	Paralyzed 6 days
K. G.	31	Paralytic	8 months	1 : 2	0.13	0.60	J 155	Paralyzed 7 days
K. G.	31	Paralytic	8 months	*	0.10	0.60	135	Paralyzed 11 days
S. H.	3	Paralytic	6 months	1 : 10	0.13	0.60	J 160	Paralyzed 8 days
S. H.	3	Paralytic	8½ months	1 : 2	0.13	0.60	R 49	Paralyzed 8 days
C. H.	6	Paralytic	1 month	1 : 2	0.13	0.60	49§	No paralysis
C. H.	6	Paralytic	7 months	1 : 2	0.13	0.60	J 159	Paralyzed 15 days
A. G.	29	Paralytic	3 weeks	1 : 2	0.13	0.60	R 62	Paralyzed 8 days
A. G.	29	Paralytic	7 months	1 : 2	0.13	0.60	J 157	Paralyzed 5 days
S. T.	8	Paralytic	7 months	1 : 2	0.13	0.60	J 176	Paralyzed 9 days
J. S.	5	Non-paralytic	6 months	1 : 10	0.13	0.60	J 165§	No paralysis
W. S.	11	Non-paralytic	6 months	1 : 10	0.13	0.60	J 166	Paralyzed 6 days
K. S.	11	Non-paralytic	7 months	1 : 10	0.13	0.60	J 177	Paralyzed 15 days
P. M.	3	Non-paralytic	3 weeks	1 : 10	0.13	0.60	J 180	Paralyzed 7 days
P. M.	3	Non-paralytic	3 months	1 : 10	0.13	0.60	J 181	Paralyzed 7 days
P. M.	3	Non-paralytic	6 months	1 : 10	0.13	0.60	J 183	Paralyzed 7 days

* Undiluted.

† Bulbar case.

‡ 1 : 6 dilution of serum.

§ These animals were subsequently proven to be resistant to poliomyelitis virus. The results have been omitted in the tabulations.

to neutralize 0.2 cc. of a 5 per cent virus suspension, an amount of the same batch of virus that this serum had neutralized in Experiment 1; the injected animal contracted poliomyelitis after a prolonged incubation period. In several other experiments, this serum had neutralized 0.13 cc. of a 1:2 dilution of a 5 per cent suspension, the usual test dose. In the next experiment, the same serum neutralized a similar test dose of the same batch of virus.

Experiment 6. Inasmuch as the positive control serum in Experiment 5 failed to neutralize the virus, the same positive and negative control

that the positive control serum in the previous experiment was tested against too large a dose of virus.

Summary of all tests within the first year with the convalescent sera of individuals whose sera failed to protect in the first week of the disease

A. Paralytic. Convalescent sera from 39 persons of various ages (Table VIII) were tested at frequent intervals after the onset of the disease.

At 2 or 3 weeks after the onset, only 1 of 24

TABLE VII
Recheck of part of Experiment 5

Patient's initials	Age	Type of case	Time after onset	Dilution of 5 per cent virus	Amount of virus	Amount of serum	Number of monkey	Results
Negative control	years	Normal monkey serum		1 : 20	cc. 0.13	cc. 0.60	J 260	Paralyzed 7 days
Negative control		Saline		1 : 20	0.13	0.60	J 274	Paralyzed 8 days
Positive control		Normal adult human serum (M.B.)		1 : 2	0.13	0.60	J 215	No paralysis
A. P.	16	Non-paralytic	2 days	*	0.45	0.45	J 211	No paralysis
V. M.†	4	Paralytic	3 days	1 : 20	0.13	0.60	J 273	Paralyzed 9 days
V. M.	4	Paralytic	7 months	1 : 20	0.13	0.60	J 277	Paralyzed 8 days
P. M.	3	Non-paralytic	3 months	1 : 20	0.13	0.60	J 272	Paralyzed 9 days

* Undiluted
† Bulbar case.

sera were retested. Whereas, in the previous experiment this positive control serum was tested against 0.2 cc. of a 5 per cent virus suspension, now 0.13 cc. of a 1:2 dilution of a 5 per cent suspension of the same batch of virus was used. As another positive control, the serum of a non-paralytic case (A. P.) which had previously protected was tested against a larger dose, 0.45 cc. of a 5 per cent virus suspension. In addition, 3 sera (V. M., 2 specimens, and P. M.) which had failed to protect were again tested with smaller amounts of virus than were used in the previous experiment. The results are shown in Table VII.

The positive control serum protected against the usual test dose of virus, while the negative control and the 3 other negative sera which were retested with smaller amounts of virus failed to do so. The serum from the non-paralytic case (A. P.) again protected. It appears, therefore,

sera protected; 7 of them were tested both at 2 and 3 weeks. Sera of 18 of the above individuals as well as of 12 others that failed to protect upon admission were retested once or twice at intervals of 1 to 6 months after the onset. Protection occurred in only 2 of these specimens, one, K. F., at 7 weeks, and the other, V. C., at 2½ and 3½ months. The latter showed protective substances for the first time on the 16th day.

At 7 to 9 months after the onset 20 of 21 sera tested failed to protect. The one serum that protected was obtained 9 months after the onset of the illness from K. F. who already had shown protective substance at 7 weeks. The other patient whose serum had developed protective substance previously was not available for retesting. The sera from 18 of the 21 individuals had been tested earlier during convalescence; 4 once, 12 two or three times and 2, four times always with

TABLE VIII

Summary of neutralization tests during first year of convalescence of individuals whose sera failed to neutralize in the first week

Age groups	Paralytic		Non-paralytic	
	Number tested	Number neutralized	Number tested	Number neutralized
<i>years</i>				
1-5.....	16	1	2	0
6-10.....	14	0	6	0
11-17.....	6	1	3	0
Adult.....	3	0	0	0
Total.....	39	2	11	0

negative results. The other 3 had been tested previously only in the acute stage of the disease.

B. Non-paralytically. Convalescent sera from 11 persons of various ages were tested at frequent intervals after the onset of the disease (Table VIII). Seven sera were tested 2 to 3 weeks after the onset and none neutralized the virus. Six sera, tested 1 to 3 months after the onset, showed no protective power; three of them had been tested previously during convalescence. Six sera tested 6 to 7 months after the onset also gave negative results; all but one of these had been tested earlier in convalescence.

C. Cases with encephalitic symptoms. The sera from two of these patients were tested and failed to show protective substances; one of them was tested 3 and 7 weeks, and 3 and 5 months after the onset, the other 6½ months after the onset.

Summary of tests with convalescent sera from individuals whose sera protected in the first week of the disease

The sera of 5 paralytically and 2 non-paralytically which showed protective power in the acute stage, when retested at later intervals still protected. The sera from two of the paralytically were retested in the second and third week respectively; two others at the end of 2 months; and the fifth after 6 months. One of the non-paralytically whose sera had protected on the 5th day, gave similar results in the 2d and 4th weeks, the other which had protected on the 4th day, also protected 6 months later. Thus, the ability of a serum to protect in the acute stage of the disease did not appear to be temporary, but was main-

tained for some time at least. However, no evidence of an increase in protective power was demonstrated either in two paralytically or in one non-paralytic. Serum from one of the paralytically had neutralized 0.1 cc. of a 5 per cent virus suspension in the acute stage, but failed to neutralize 0.2 cc. after 3 months. The other did not neutralize 0.1 cc. of a 5 per cent virus after 7 weeks, although a specimen obtained during the first week neutralized half that amount. A non-paralytic case also failed to show an increase of protective antibodies after 6½ months.

Tests with a strain of virus isolated in 1935

In order to compare the results with those obtained previously with the passage virus, two experiments were performed upon sera obtained in the acute and convalescent stages using the strain isolated from nasal washings.

Experiment 7. Ten specimens from 7 patients were tested; 3 were taken both in the acute and convalescent stages and 4 only in convalescence. None of these specimens had neutralized the F1 virus. The results, shown in Table IX, indicated that 2 of the 3 sera obtained soon after the onset and all of the 7 convalescent specimens failed to protect.

Experiment 8. Sixteen sera were tested. Six of these had previously protected against the passage virus. These included 3 specimens obtained from non-paralytically in the acute stage and 3 from paralytically during convalescence. Two of the latter (V. C. and K. F.) were from patients who showed protective substances both in the acute stage and in convalescence. Nine specimens obtained 5 to 9 months after the onset had failed to neutralize the F1 strain. Two of these were from non-paralytically and 7 from paralytically. The 16th serum was an acute-stage specimen of S. S. which appeared to protect against the recently isolated strain in Experiment 7 but had failed to neutralize the passage virus previously. The results are given in Table X. All 6 specimens which had protected against the passage virus also neutralized the recently isolated strain. The 9 convalescent sera which had failed to protect against the passage virus also failed to protect when the recently isolated strain was used. The specimen obtained from S. S. in the acute stage,

TABLE IX
Neutralization tests with a strain of virus isolated during the 1935 outbreak

Patient's initials	Age	Type of case	Time after onset	Dilution of 5 per cent virus	Amount of virus	Amount of serum	Number of monkey	Results
Negative control	<i>years</i>	Normal monkey serum		1 : 2	cc. 0.13	cc. 0.60	J 252	No paralysis
Negative control		Normal monkey serum		1 : 10	0.13	0.60	J 253	Paralyzed 14 days
Negative control		Normal monkey serum		1 : 40	0.13	0.60	J 248	Paralyzed 8 days
Positive control		Normal adult human serum (A.G.)		1 : 2	0.13	0.60	J 72	No paralysis
P. B.†	5	Paralytic	3 days	1 : 10	0.13	0.60	J 249	Paralyzed 6 days
P. B.	5	Paralytic	4 months	1 : 2	0.13	0.60	J 73	Paralyzed 15 days
P. B.	5	Paralytic	4 months	1 : 10	0.13	0.60	J 241	Paralyzed 8 days
S. S.	9	Paralytic	5 days	1 : 10	0.13	0.60	J 244	No paralysis
S. S.	9	Paralytic	7 months	1 : 10	0.13	0.60	J 243	Paralyzed 8 days
R. M.	4	Paralytic	5 days	1 : 2	0.13	0.60	J 258	Paralyzed 5 days
R. M.	4	Paralytic	3½ months	1 : 2	0.13	0.60	J 257	Paralyzed 6 days
G. C.	5	Paralytic	3 months	1 : 2	0.13	0.60	555	Paralyzed 13 days
G. C.	5	Paralytic	3 months	1 : 10	0.13	0.60	J 242	Paralyzed 8 days
G. M.	13	Paralytic	4 months	1 : 2	0.13	0.60	J 259	Paralyzed 6 days
J. C.	8	Paralytic	7 months	1 : 10	0.13	0.60	J 246	Paralyzed 8 days
J. S.	3	Paralytic	8 months	1 : 10	0.13	0.60	J 247	Paralyzed 12 days

† Bulbar case.

TABLE X
Neutralization tests with a strain of virus isolated during the 1935 outbreak

Patient's initials	Age	Type of case	Time after onset	Dilution of 5 per cent virus	Amount of virus	Amount of serum	Number of monkey	Results
Negative control	<i>years</i>	Normal monkey serum		1 : 10	cc. 0.13	cc. 0.60	J 250	Paralyzed 8 days
Negative control		Normal monkey serum		1 : 10	0.13	0.60	J 271	Paralyzed 12 days
Negative control		Normal monkey serum		1 : 80	0.13	0.60	J 289	No paralysis
Positive control		Normal adult human serum (A.G.)		1 : 2	0.13	0.60	J 267	No paralysis
R. G.	3	Non-paralytic	4 days	1 : 2	0.13	0.60	J 215	No paralysis
L. W.	8	Non-paralytic	2 days	1 : 2	0.13	0.60	J 256	No paralysis
R. J.	16	Non-paralytic	4 days	1 : 2	0.13	0.60	J 239	No paralysis
V. C.	2	Paralytic	3½ months	1 : 2	0.13	0.60	J 290	No paralysis
E. F.	4	Paralytic	3 months	1 : 2	0.13	0.60	J 266	No paralysis
K. F.	13	Paralytic	8½ months	1 : 2	0.13	0.60	J 263	No paralysis
D. D.	8	Non-paralytic	6 months	1 : 2	0.13	0.60	J 165†	No paralysis
E. B.	11	Non-paralytic	5½ months	1 : 2	0.13	0.60	J 291	Paralyzed 7 days
K. S.	11	Non-paralytic	6 months	1 : 10	0.13	0.60	J 278	Paralyzed 6 days
A. D.	2	Paralytic	8½ months	1 : 10	0.13	0.60	J 284	Paralyzed 10 days
S. T.	8	Paralytic	7 months	1 : 2	0.13	0.60	J 276	Paralyzed 7 days
H. P.	9	Paralytic	9 months	1 : 2	0.13	0.60	J 280	Paralyzed 9 days
D. F.	17	Paralytic	7½ months	1 : 2	0.13	0.60	J 264	Paralyzed 11 days
M. S.	26	Paralytic	7½ months	1 : 2	0.13	0.60	J 262	Paralyzed 11 days
A. G.	29	Paralytic	7 months	1 : 2	0.13	0.60	J 220	Paralyzed 9 days
K. G.	31	Paralytic	7½ months	1 : 2	0.13	0.60	J 275	Paralyzed 7 days
S. S.	9	Paralytic	5 months	1 : 10	0.13	0.60	J 286	Paralyzed 11 days

† This animal was resistant to poliomyelitis virus. The result was not included in the tabulations.

now failed to neutralize, a finding in keeping with that obtained with the passage virus.

These two experiments indicated that sera obtained in the acute and convalescent stages reacted similarly to both strains of virus. Three of 6 specimens obtained in the acute stage protected against both strains, and 3 failed to neutralize either strain. Three of 19 specimens obtained several months after the onset protected against both strains, whereas the other 16 neutralized neither strain of virus. The majority of convalescent sera thus failed to protect against a strain of virus obtained from the nasal washings of a patient in the same outbreak.

Neutralization tests in which passage virus and sera obtained from 12 to 16 months after the onset of the disease were used

Experiment 9. Thirty sera were tested, 24 from paralytic patients, 5 from non-paralytics and 1 from a patient who had the encephalitic form. Specimens taken from these patients during the acute stage and at intervals during the first year of convalescence had failed to neutralize the virus. The sera from two of the paralytic cases and one non-paralytic case taken 12 to 16 weeks after the onset neutralized the virus in duplicate tests. Thus, 3 of 30 patients apparently developed protective substances at the beginning of the second year following the illness. Two other persons whose sera protected during the acute stages of the disease were bled a year later in order to determine whether the neutralizing substances were still present. Both sera again protected.

DISCUSSION

Protective substances were found in 14 of 82 paralytic individuals in the first week of the disease; several of these patients had extensive paralysis. Kling and Levaditi (6), Flexner and Amoss (7), and Harmon and Harkins (15) also recorded instances in which protective substances were detected in the first week of illness. The first two authors (6, 7) thought that the detected neutralizing bodies had developed rapidly after the onset, while Harmon and Harkins (15) believed that they were present prior to infection. We concur with the latter opinion, for protective

substances were found very early in the disease and did not develop readily in convalescence. The presence of protective substances were demonstrated in the sera of two patients in the pre-paralytic stage, in one patient on the day paralysis developed, and in several others as early as one and two days after the onset. In one of the cases tested by Harmon and Harkins (15) neutralizing substance was also found in the pre-paralytic stage. On the other hand, when 39 of our 68 patients without protective substances in the acute stage were retested during the first year of convalescence, protective substances were found in the sera of only 2. Moreover, the sera of 2 individuals which had neutralizing power in the first week of the illness did not show a demonstrable increase during convalescence. Other observers also found that neutralizing substances did not develop readily during convalescence following paralysis (15, 19).

Non-paralytic cases showed protective substances in the acute stage of the disease more frequently (18 out of 32) than did paralytic cases. The neutralizing substances probably were present before the onset and did not develop rapidly after the disease set in. Their presence was detected as early as the first and second day of the illness. On the other hand, 11 of the 14 cases whose sera had no protective substances in the acute stage of the disease when retested several months later were still negative. This finding is in agreement with that of Paul and Trask (19) who found no neutralizing substance in a non-paralytic case several weeks and again one year after the onset.

The failure to develop protective substances in convalescent sera from paralytic cases also coincided with the findings of Paul and Trask (19) who showed that 6 of 7 sera taken 1½ to 10 weeks after the onset failed to neutralize the virus; 2 of them when retested a year later still failed to neutralize. On the other hand, Jensen (24) reported the presence of antibodies in paralytic and non-paralytic cases within a month after the onset. In a recent paper, however, Eagles and coworkers (25) stated that a pooled serum obtained from 67 paralytic patients approximately 4 weeks after the onset failed to neutralize the virus of poliomyelitis.

The failure of the majority of patients to de-

velop protective substances within 6 to 9 months after the onset is further confirmed by the negative results obtained with a strain of virus isolated from the same outbreak. Such findings differ from those of Howitt (22) who reported that convalescent serum more often neutralized a recently isolated strain than a passage virus. Likewise, Paul and Trask (19) found that convalescent sera from 6 of 7 paralytic cases neutralized a recently isolated strain, whereas only one serum neutralized the passage virus. It is possible that the strain used in the present work was more closely related to the passage virus than that used by the above workers (19, 22), for both strains of virus reacted similarly to 25 sera, 6 neutralizing and 19 failing to neutralize each strain. Moreover, 4 of 6 monkeys with residual paralysis that had recovered and were resistant to reinfection with F1 virus, were resistant also to the other strain. One of the 2 which failed to resist, developed only a mild attack with a rise in temperature and transitory weakness of the arms. On the other hand, the above mentioned investigators may have obtained neutralization more often with their recently isolated strains because they were less virulent than the passage strains. Howitt (22) indicated that the strain she used did not infect with regularity, while Paul and Trask (19) mentioned that their virus usually caused a less severe form of the disease than did the passage virus.

The absence of protective bodies during the first few months of convalescence in 48 of 50 paralytic and non-paralytic individuals whose sera failed to neutralize upon admission was demonstrated not only when 0.13 cc. of a 2.5 per cent (1:2 dilution of a 5 per cent virus suspension) was mixed with 0.6 cc. of serum, but also in many instances when tested against much smaller amounts of virus. Even though a number of sera were tested against 0.13 cc. of 5 per cent virus diluted 10 to 64 times, and in one instance 80 times, they still failed to neutralize. Smaller amounts of virus were used in testing these sera than most other investigators have employed. However, the intracerebral inoculation of a virus-serum mixture may not be suitable for the demonstration of neutralizing antibodies. Even

though diluted virus suspensions were used, the test still may be too severe.

A higher incidence of protective substances, 63 per cent, has been reported (Table I) in convalescent sera from paralytics obtained years after the disease, than was found in the present work in which the sera were collected within a year of onset. Our figure was approximately 22 per cent and included neutralizing substances found at the onset of the disease (17 per cent), and those which developed in convalescence (5 per cent). There have been no significantly large number of sera taken by any one group of investigators within one year of the onset of the disease for comparison with the results of the present work. The variations in technique would hardly explain the different results, for other workers frequently obtained neutralization although larger amounts of virus suspension were used. Howitt (20b) reported that pooled convalescent sera from long standing cases had a higher titer of neutralizing substances than those from recently recovered cases. In this investigation, the sera of 24 paralytics which had failed to neutralize during the first 9 months of convalescence, were retested 12 to 16 months after the onset. Two of these specimens obtained 14 months after the onset from children 7 and 9 years of age neutralized for the first time. It is also possible that more of these individuals may subsequently develop neutralizing substances. If this does occur, is it due to the slow development of neutralizing substances, is it produced by further exposure to the virus resulting in hyperimmunization, or can it appear as a result of nonspecific factors as Jungblut and Engle (26) have suggested? It is noteworthy that the sera which neutralized at the beginning of the second year after the onset were obtained in October 1936. However, there was practically no poliomyelitis in New York City during 1936. The children from whom the sera were obtained had not been in a locality where poliomyelitis was prevalent.

Until recently the presence of neutralizing substances in the serum was accepted as an index of immunity to poliomyelitis. Certain experimental and clinical observations, however, do not support this view. It has been shown (27, 28) that sera of monkeys injected subcutaneously with a

subinfective dose of virus may after a few weeks neutralize poliomyelitis virus *in vitro*, even though the animals are not able to resist an intracerebral or intranasal instillation of potent virus. Further, convalescent monkeys which are refractory to reinoculation may show no neutralizing substance in their sera (29, 30, 31). Evidence has been presented previously to support the contention that protective substances were present in 14 of 82 paralytic patients when the disease developed. There was no evidence to indicate that the paralysis was less extensive in the 14 persons whose serum neutralized in the acute stage than in the 68 whose serum failed to do so. Only 2 of the 14 had mild paralysis, that is a partial involvement of one limb, 2 had bulbar involvement, 2 others showed extensive paralysis of one leg. The remaining 8 had involvement of 2 extremities or more. Of the 68 whose serum failed to neutralize in the acute stage, 8 had a mild paralysis, 13 had involvement of one leg and 9 cases were bulbar. The remaining 38 had a muscular paralysis in at least two extremities. The majority of sera tested from the convalescent paralytic patients without protective substances in the first week of the disease did not develop any as late as 12 to 16 months after the onset of the disease. Nevertheless, second attacks of poliomyelitis are rare. Although one cannot always interpret human disease in the light of animal experimentation, one may assume an immunity in the majority of convalescent poliomyelitis patients in the absence of protective substances in the serum.

On the other hand, the incidence of protective substance in non-paralytic patients is greater than that in paralytics and approximates that in normal individuals of the same age. In the acute stage of the disease in non-paralytic patients its presence may help to limit the spread of the virus, while in the paralytics its frequent absence may be a factor accounting for the widespread involvement in some cases. Very noteworthy was the absence of neutralizing substances in the specimens of serum taken on admission and one year later from a paralytic boy who had had a previous attack of poliomyelitis in 1933 with residual paralysis.

Consequently, it appears that there must be factors other than the presence or absence of protective substances in the serum that determine

the resistance of the host to poliomyelitis. Indeed, in interpreting these results, we must consider the possibility that the neutralizing substances are not specific, that is, they may develop irrespective of exposure to the virus. The immunity present in the absence of neutralizing substances may be cellular rather than humoral.

The presence or absence of neutralizing substances in the acute and convalescent stages was not related to the degree of recovery from paralysis, a finding in agreement with that of Harmon and Harkins (15). Howitt (12), on the other hand, indicated that some correlation did exist between these two factors. We were unable to find that the 14 paralytic cases who had neutralizing substances in the acute stage improved more rapidly than the other 68 who had none. Indeed, in three of the 14, the paralysis was progressive. Of the two paralytic individuals (Table VIII) who developed demonstrable neutralizing substances within the first 9 months of convalescence, one showed considerable recovery of muscular power in several months, while the other improved slowly although a progressive increase in neutralizing substances was demonstrated after the 16th day. In 16 of 20 whose sera had failed to neutralize after 7 to 9 months, considerable clinical improvement occurred. One bulbar case and two spinal-paralytic cases made a complete recovery within a month after the onset.

The protection test offers no aid in the diagnosis of non-paralytic poliomyelitis. In the acute stage the proportion of sera from patients over the age of 5 that neutralized approximated that for normal individuals. Further, 11 of those whose sera showed no neutralizing power in the acute stage did not develop any within 6 to 7 months, thereby failing to give any serological evidence that the diagnosis was correct. Clinically, all the cases had the typical type of onset and symptoms, including pain and stiffness of the neck and back with pleocytosis of the cerebrospinal fluid. The lack of serological evidence does not invalidate the diagnosis, inasmuch as the majority of cases with frank paralysis also failed to develop protective substances.

SUMMARY

1. Neutralization tests carried out with the F1 strain (monkey passage) of virus on sera ob-

tained from paralytic and non-paralytic cases during an outbreak of epidemic proportions gave the following results: (a) Of 82 paralytic cases tested during the acute stage of the disease, the sera of 14 neutralized the virus; (b) 4 of these patients were tested in the preparalytic stage, and the sera of 2 possessed protective bodies; (c) of 32 non-paralytic cases tested in the acute stage of the disease, the sera of 18 neutralized; (d) of 3 encephalitic cases tested in the acute stage, the serum of one neutralized; (e) during convalescence only 2 of 39 paralytics who had no protective substances in the acute stage developed them within a few months after the onset. Two of 24 individuals whose sera had failed previously to neutralize in convalescence developed protective substances 12 to 16 months after the onset; (f) neutralizing substances which had not been present in the acute stage in 11 non-paralytics also failed to develop in the sera tested several months after the onset. One of 5 obtained 12 to 16 months after the onset neutralized; (g) the sera of 2 cases with encephalitic symptoms likewise failed to protect 5 to 6 months after the onset. One which was obtained 12 months after the onset was also negative; (h) two paralytics and one non-paralytic, whose sera neutralized in the acute stage of the disease, failed to show a demonstrable increase of neutralizing substances 2 months or longer after the onset.

2. Sera of 9 of 18 so-called normal individuals, over the age of 10, neutralized.

3. A strain of virus isolated from the outbreak from which the bloods were collected was tested against the sera and gave results comparable to those obtained with the passage (F1) virus.

4. No evidence of a definite relationship was found between the presence of protective substances in serum and (1) resistance to poliomyelitis, (2) the diagnosis of the non-paralytic form of poliomyelitis and (3) the degree of recovery from paralysis.

The authors wish to express their thanks to Dr. Thomas M. Rivers and to Dr. Morris Schaeffer for their valuable suggestions during the course of this work.

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