CLINICAL ASSESSMENT OF THE TREPONEMAL WASSERMANN REACTION*†

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The value of a new serological test for syphilis is best assessed by examining its effects in two groups of patients; the first group should be composed of patients with no evidence of treponemal disease as determined by clinical examination and testing by already established serological methods, and the second should consist of patients known to have syphilis. Preliminary assessments of the treponemal Wassermann reaction (TWR) were reported by Price and Whelan (1957) and by Price (1958) and, as a result of this work, the present clinical study was planned.

Material and Tests Employed

The TWR was performed as a routine, together with the Wassermann reaction (WR) and Price's precipitation reaction (PPR), on all patients attending the Whitechapel Clinic of The London Hospital between November 1, 1957, and October 31, 1958.

The serological technique employed during the year under review for the WR has been described by Price (1950) and that for the PPR by Price (1948).

Over 10,000 TWRs were carried out on 4,105 patients (Table I), all of whose case records were reviewed. There were 3,078 men (778 coloured and 2,300 white) and 1,027 women (152 coloured and 875 white).

TABLE I
TOTAL PATIENTS REVIEWED BY RACE AND SEX

Sex	Race		Total
	Coloured	White	
Women	152	875	1,027
Men	778	2,300	3,078
Total	930	3,175	4,105

Controls

Method of Study.—Requirements for the control group of patients without evidence of treponemal disease were as follows:

- (1) No history suggesting treponemal disease.

- (2) No clinical sign suggesting treponemal disease.
- (3) Negative initial standard serological tests for syphilis (STS).
- (4) Negative confirmatory standard STS (carried out after an interval of not less than 6 weeks).
- (5) No evidence of syphilis at the confirmatory clinical examination (carried out after an interval of not less than 6 weeks).
- (6) No antibiotic with antitreponemal action given between the initial and the confirmatory examination and STS.

To fulfil these requirements, all suitable patients, in whom there was no evidence of treponemal infection either clinically or by standard serological tests, were asked to return for confirmatory examination and blood tests 6 weeks or more after the initial examination.

Results.—The 647 patients in this control group were subdivided into those who gave a history of having received a treponemicidal antibiotic or drug before the review (107) and those without such history (540).

(A) The main control group, consisting of patients with no previous history of having been given an antibiotic or drug, is considered first. The results of this analysis are shown in Table II (overleaf).

White Patients.—This group included 130 white women (15 per cent. of the white women who attended during the year). The initial TWR was positive in eight cases, and a treponemal immobilization test (TPI) was performed in seven of these with negative results.

There were 352 white men (15 per cent. of the white men who attended during the year). The initial TWR was positive in thirteen cases, and a TPI was performed in four of these with negative results.

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TABLE II
RESULTS IN 647 CONTROLS

Controls		(A) With No Previous History of Antibiotic					(B) With Previous History of Antibiotic			Total Controls
Race	Sex	Number of Cases	Initial TWR Positive	TPI Negative	Initial TWR Negative Becoming Positive	TPI Negative	Number of Cases	Initial TWR Positive	TPI Negative	
White	Female	130 (15 per cent.)	8	7	3	1	17	2	1	147
	Male	352 (15 per cent.)	13	4	5	3	67	6	4	419
	Total	482	21 (4.4 per cent.)	11	8	4	84	8	5	566
Coloured	Female	11 (7 per cent.)	4	4	3	1	1	1	1	12
	Male	47 (6 per cent.)	20	11	6	4	22	9	2	69
	Total	58	24 (41.4 per cent.)	15	9	5	23	10	3	81
Total ..	..	540	45 (8.3 per cent.)	26	17	9	107	18 (16.8 per cent.)	8	647

Thus the initial TWR was positive in 21 cases (4.4 per cent.) of this group of white patients who had no evidence of syphilis either clinically or on testing by methods other than the TWR.

Coloured Patients.—There were only eleven coloured women who fulfilled the criteria for the control group and they comprised 7 per cent. of the coloured women who attended during the year. The initial TWR was positive in four cases, and the TPI was negative in all of them.

There were 47 coloured men (6 per cent. of those reviewed). The initial TWR was positive in twenty cases and the TPI was performed in eleven of them with negative results.

Thus, out of 58 coloured patients without evidence of treponemal disease, the initial TWR was positive in 24 (41 per cent.), and the TPI was performed in fifteen of these 24 cases with negative results.

Total.—Out of 540 patients in this main control group, who were selected as having no evidence of syphilis or yaws, no less than 45 (8.3 per cent.) had an initially positive TWR.

As already noted, for a patient to be admitted to the control series, the initially negative standard STS, performed in parallel with the TWR, had to be repeated with negative result after an interval of not less than 6 weeks. Thus all the 540 control cases had one or more TWRs after the original test.

The sixth column of Table II shows that in seventeen cases out of 540 (3.2 per cent.) the TWR,

although initially negative, became positive. This occurred in patients under supervision in whom there was no evidence of syphilis. In nine of these seventeen the TPI was performed, with negative results. This finding is relevant at the present stage, but it also gives information on the reproducibility of the TWR. If these seventeen patients are considered together with the 45 in which the TWR was initially positive, then a total of 62 (11.5 per cent.) of the 540 control cases gave positive results to the TWR.

In the 58 coloured patients the TWR was positive initially in 24 and subsequently in a further nine, to make a total of 33 cases (57 per cent.) in which the test was positive. The countries of origin of these patients were as follows:

Caribbean Zone	27	(TWR positive in 15)
West Africa	6	(TWR positive in 4)
India	11	(TWR positive in 4)
Pakistan	12	(TWR positive in 10)
Burma	1	(both TWR negative)
Arabia	1	

(B) There is further evidence from the second control group, comprising patients who fulfilled all the criteria originally laid down, but who had received a treponemicidal drug or antibiotic before the review. Table II shows that, of 107 such cases, the initial TWR was positive in eighteen (16.8 per cent.), and the TPI, performed in eight of these, was negative.

Other "Controls".—Further evidence is given in Table III. In 25 patients the TWR was positive, although there was no evidence of treponemal disease and the standard STS were negative on two or more occasions. They cannot be included in the control group proper either because they received a treponemical antibiotic after the initial blood test, or because they were under observation for less than 6 weeks. In these 25 patients, although the TWR was positive, the subsequent TPI was negative.

TABLE III
OTHER "CONTROLS"

All were TWR Positive and TPI Negative with No History and No Signs, but they had received Antibiotic Treatment after the Initial TWR, or were observed for less than 6 weeks

Sex ..	Male	Female	Total
Race .. Coloured ..	11	4	15
White ..	5	5	10
Total ..	16	9	25

The control group has been comprised selectively of patients with no evidence of treponemal disease either clinically, or by serological tests other than the TWR. In the 4,105 patients reviewed, however, thirteen were discovered without definite history or signs of treponemal disease, in whom the standard STS were negative but the TWR and subsequent TPI proved positive. These patients are shown in Table IV. There is no doubt that if the positive TWR had not led to the performance of a TPI test, these cases would have been overlooked. It is interesting that twelve out of the thirteen patients were coloured persons.

TABLE IV
PATIENTS WITH POSITIVE TWR AND TPI ONLY
(No History or Signs)

Sex ..	Male	Female	Total
Race .. Coloured ..	7	5	12
White ..	Nil	1	1
Total ..	7	6	13*

* TPI=100 per cent. in eight, "doubtful" in one, 58-82 per cent. in four.

Reproducibility.—A serological test for syphilis should be specific and sensitive, and the results it gives should be reproducible. Table V gives details of 22 cases in the control group in which the TWR, although initially negative, became positive.

Group A consisted of seventeen patients with no history of previous treatment with treponemical

drug or antibiotic. In eight white patients the TWR was initially negative and became positive; in four of them a TPI was performed and was negative. In the smaller group of coloured control patients there were nine such cases and in five of them the TPI was performed and was negative.

Group B consisted of five patients with a history of previous treatment with treponemical drug or antibiotic. There were four white patients; the TPI was performed in two and was negative. There was one coloured patient.

These patients were under clinical observation and, although the TWR became positive, none developed evidence of syphilis.

TABLE V
REPRODUCIBILITY OF TWR IN CONTROLS

Controls		(A) No Previous Antibiotic		(B) Previous Antibiotic	
Race	Sex	TWR Initially Negative Subsequently Positive	TPI Negative	TWR Initially Negative Subsequently Positive	TPI Negative
White	Female	3	1	1	—
	Male	5	3	3	2
	Total	8	4	4	2
Coloured	Female	3	1	—	—
	Male	6	4	1	—
	Total	9	5	1	—
Total ..	..	17	9	5	2

The TPI was negative in 35 members of the main control group in whom the TWR was positive in one or more tests. Reproducibility was studied in these cases by comparing the initial TWR with the subsequent TWR results (Table VI). In nineteen white patients, subsequent TWRs were "concordant" in fourteen and "discordant" in five. In sixteen coloured patients, subsequent TWRs were "concordant" in eleven and "discordant" in five. Thus,

TABLE VI
CONCORDANCE OF INITIAL TWR WITH SUBSEQUENT TWRs IN CONTROLS WITH NEGATIVE TPI

Race	Sex	"Concordant"	"Discordant"	Total
White	Female	5	2	7
	Male	9	3	12
	Total	14	5	19
Coloured	Female	2	2	4
	Male	9	3	12
	Total	11	5	16
Total ..	..	25	10	35

in these 35 patients, in all of whom the TPI was negative and none of whom had received a treponemidal antibiotic or drug during the period of observation, the results of the TWR were "concordant" in 25 and "discordant" in ten.

Another way of assessing reproducibility is to analyse the results of the TWR in all cases in the main control group (Table VII). There were 342 white men in whom two or more tests (a total of 841 TWRs) gave concordant results; in ten cases the results were discordant. Similarly 125 white women had 268 concordant TWRs, and in five cases the results were discordant. Thus, in 467 white patients, 1,109 concordant TWRs were performed, and in fifteen cases (3 per cent.) the results were discordant. This may be compared with the total of 43 coloured patients in whom 110 TWRs gave concordant results, and the fifteen (26 per cent.) in whom the results were discordant.

TABLE VII
REPRODUCIBILITY IN CONTROLS

Race	Sex	"Concordant"		"Discordant"
		Cases	Sera	Cases
White	Male	342	841	10
	Female	125	268	5
	Total	467	1,109	15 (3 per cent.)
Coloured	Male	37	98	10
	Female	6	12	5
	Total	43	110	15 (26 per cent.)
Total		510	1,219	30

It can be concluded that the reproducibility of the TWR is poor, especially in tests on sera from coloured patients.

Syphilis Group

Criteria for Selection.—All cases were studied in which a diagnosis of syphilis had been made at any time and in which the result of at least one TWR was available. From these a smaller group was selected which satisfied the strictest diagnostic criteria and, in this group, the results obtained with the TWR were compared with those obtained with the WR, PPR, and TPI.

Primary and Secondary Syphilis.—Only patients in whom treponemes had been found on darkfield examination were included.

Early Latent Syphilis.—This diagnosis was accepted only if one or more of the following criteria were satisfied:

- The patient was known to have had negative STS within the four years prior to diagnosis.
- The patient's consort had developed syphilis but was known to have had previous negative STS as above.
- The consort had darkfield positive lesions, or clinical secondary syphilis with positive STS.
- The patient had recently given birth to a child suffering from congenital syphilis.

Late Latent Syphilis.—If the historical evidence was sufficient this diagnosis was used, but the majority of cases of latent syphilis fell into the category of "latent syphilis of unknown duration".

"Syphilis or Yaws".—This category included coloured patients who had been born in areas where yaws occurs and were suffering from treponemal disease, other than early syphilis, congenital syphilis, or involvement of the neurological or cardiovascular systems.

Method of Study.—The results of the WR, PPR, and TWR at the time of diagnosis were recorded on an abstract sheet as "Positive", "Negative", or "No data", and the results of subsequent tests were entered in the same way under the appropriate time interval from the date of diagnosis. The TPI was similarly recorded with additional provision for the entering of "doubtful" results. Data so obtained were transferred to punched cards and mechanically sorted.

Results.—439 cases of syphilis proved suitable for this study and the results of the WR, PPR, and TWR in this group were analysed. It had been hoped that it would be possible to produce graphs to show the percentage of positive results for each test, at diagnosis, and at intervals after treatment. Owing to the small numbers in the different groups this has been possible in only some instances. Cases in which clinical or serological relapse occurred after treatment, or in which lesions of late syphilis developed, have been excluded, and the following charts show what happened after the successful treatment of syphilis. Follow-up tests were also excluded in those cases in which treatment had been inadequate.

In the construction of the charts which outline the results of individual tests before and after treatment in the different forms of syphilis, minor variations of

technique in the earlier tests have been ignored. All the tests were performed at the Laboratory of the Whitechapel Clinic.

Sero-Negative Primary Syphilis.—Fig. 1 shows the results in 44 cases. At diagnosis all forty cases in which the WR had been performed, and all 29 in which the PPR was carried out, gave negative results. This is so by definition. One out of the fourteen cases in which the TWR was performed, however, gave a positive result, and this was in a coloured patient.

The next set of columns shows the results of tests on all sera examined within 3 months after diagnosis. Here five out of eleven TWR results were positive and all the WR, PPR, and TPI results were negative.

Table VIII summarizes all the follow-up tests; only two proved positive out of 335 WRs and all the 299 PPRs were negative, but eight out of sixty TWRs were positive. This appears to be in keeping with the high percentage of positive TWRs in the control group.

TABLE VIII
SUMMARY OF FOLLOW-UP IN 44 CASES OF SERO-NEGATIVE PRIMARY SYPHILIS

Test	Result		Total No. of Sera
	Negative	Positive	
WR	333	2	335
PPR	299	0	299
TPI	1	0	1
TWR	52	8	60

In the seven coloured patients in this group the TWR at diagnosis was positive in one and negative in three. In follow-up tests the WR and PPR were negative in all 28 sera tested, the TWR being positive in one and negative in nine sera.

The number of coloured patients in the other diagnostic groups (except "Syphilis or Yaws") was small as in this first group, and no statistically valid difference between the white and coloured races was apparent in the results of serological tests. For this reason the two races are considered together in the following sections.

FIG. 1.—Sero-negative primary syphilis (44 cases).

Sero-Positive Primary Syphilis.—There were 33 cases in this group (Fig. 2). At diagnosis the WR was positive in 31 of 32 cases, the PPR in 26 of 27, and the TWR in all eight cases tested.

In the follow-up tests (Table IX) the WR was negative in 211 sera and positive in fifty (19 per cent.), the PPR was negative in 229 and positive in nineteen (8 per cent.), the TPI was negative in four cases and the TWR was negative in 33 sera and positive in nineteen (36 per cent.), a higher level of positivity than was obtained with the other tests.

TABLE IX
SUMMARY OF FOLLOW-UP IN 33 CASES OF
SERO-POSITIVE PRIMARY SYPHILIS

Test	Result			Total No. of Sera Tested
	Negative	Positive		
		No.	Per cent.	
WR	211	50	19	261
PPR	229	19	8	248
TPI	4	0	0	4
TWR	33	19	36	52

FIG. 2.—Sero-positive primary syphilis (33 cases).

Secondary Syphilis.—There were forty cases in this group (Fig. 3). All tests gave positive results at diagnosis, and after treatment the WR and PPR showed a rapid reversal to negative. In the follow-up tests (Table X) the WR was negative in 272 sera and positive in 53 (16 per cent.), the PPR was negative in 243 and positive in 46 (16 per cent.), and the TWR was negative in thirty and positive in 40 (16 per cent.). Again there was a relatively large number of positive results with the TWR.

TABLE X
SUMMARY OF FOLLOW-UP IN 40 CASES OF
SECONDARY SYPHILIS

Test	Result		Total No. of Sera Tested
	Negative	Positive	
		No. Per cent.	
WR	272	53	325
PPR	243	46	289
TPI	—	—	—
TWR	30	20	50

FIG. 3.—Secondary syphilis (40 cases).

FIG. 4.—Early latent syphilis (12 cases).

Early Latent Syphilis.—There were only twelve cases in this selected group (Fig. 4). At diagnosis the WR was positive in all twelve, the PPR in eight cases out of nine, the TPI in four out of four, and the TWR in four out of five. In the follow-up tests (Table XI), the WR was negative in fifteen sera and positive in fifteen (50 per cent.), the PPR was negative in sixteen and positive in fourteen (47 per cent.), and the TWR was negative in four and positive in seven.

Latent Syphilis of Unknown Duration.—There were 66 cases in this group (Fig. 5, opposite). At diagnosis the WR was positive in 61 of 62 cases, the PPR in 46 of 48, the TPI in all of twelve, and the TWR in ten of eleven. In the follow-up tests (Table XII), the WR was positive in 310 sera (73 per cent.) and negative in 112, the PPR was positive in 248 (65 per cent.) and negative in 134, the TPI was positive in eighteen cases (the result in a further one being reported as “doubtful”), and the TWR was positive in 48 sera (69 per cent.) and negative in 22.

TABLE XI
SUMMARY OF FOLLOW-UP IN 12 CASES OF
EARLY LATENT SYPHILIS

Test	Result			Total No. of Sera Tested
	Negative	Positive		
		No.	Per cent.	
WR	15	15	50	30
PPR	16	14	47	30
TPI	0	1	—	1
TWR	4	7	—	11

TABLE XII
SUMMARY OF FOLLOW-UP IN 66 CASES OF
LATENT SYPHILIS OF UNKNOWN DURATION

Test	Result			Total No. of Sera Tested
	Negative	Positive		
		No.	Per cent.	
WR	112	310	73	422
PPR	134	248	65	382
TPI	0	18*	100	18
TWR	22	48	69	70

* Plus one “doubtful”

FIG. 5.—Latent syphilis of unknown duration (66 cases).

Late Syphilis.—There were 96 cases in this group. In Fig. 6, the positive results are expressed as a percentage of tests performed.

At diagnosis the WR was positive in 85 out of 91 cases (93 per cent.), the PPR in 61 out of 69 (88 per cent.), the TWR in twenty out of 23 (87 per cent.), and the TPI in all of the nine cases tested.

In the follow-up tests (Table XIII), the WR was positive in 491 sera (71 per cent.) and negative in 197, the PPR was positive in 440 (69 per cent.) and negative in 198, the TPI was positive in fifteen cases and negative in one (the result in a further one being reported as "doubtful"), and the TWR was positive in 61 sera (61 per cent.) and negative in 38.

In view of the small number of sera tested by the TWR, the results of the various tests may be

regarded as similar.

The increase in the percentage of positive tests fifteen years after diagnosis and subsequently may

TABLE XIII
SUMMARY OF FOLLOW-UP IN 96 CASES OF
LATE SYPHILIS

Test	Result			Total No. of Sera Tested
	Negative	Positive		
		No.	Per cent.	
WR	197	491	71	688
PPR	198	440	69	638
TPI	1	15*	94	16
TWR	38	61	61	99

* Plus one "doubtful"

FIG. 6.—Late syphilis (96 cases).

be due to the fact that more of the patients with positive STS tended to return for further observation.

Late Congenital Syphilis.—There were 45 cases in this group (Fig. 7). At diagnosis the WR was positive in all of 42 cases, the PPR in 22 out of 23, and the TWR in six out of seven. Approximately 70 per cent. of sera tested between 10 and 15 years after diagnosis were positive.

In the total follow-up tests (Table XIV), the WR was positive in 266 sera (82 per cent.) and negative in 59, the PPR was positive in 258 (84 per cent.) and negative in fifty, the TPI was positive in nine cases

and negative in two (the result in one further case being reported as "doubtful"), and the TWR was positive in 31 sera (74 per cent.) and negative in eleven.

TABLE XIV
SUMMARY OF FOLLOW-UP IN 45 CASES OF
LATE CONGENITAL SYPHILIS

Test	Result			Total No. of Sera Tested
	Negative	Positive		
		No.	Per cent.	
WR	59	266	82	325
PPR	50	258	84	308
TPI	2	9*	—	11
TWR	11	31	74	42

* Plus one "doubtful"

FIG. 7.—Late congenital syphilis (45 cases).

"*Syphilis or Yaws*".—There were 103 coloured patients in this group (Fig. 8). At diagnosis the TPI was positive in 64 cases and "doubtful" in two more, the TWR was positive in 77 out of 86 (90 per cent.), the WR in 77 out of 103 (75 per cent.), and the PPR in seventy out of one hundred. In tests carried out in the 3 months after diagnosis, the TPI was positive in all of thirteen cases, the TWR in 85 per cent. of sera tested, the WR in 79 per cent., and the PPR in 73 per cent. In the total follow-up tests (Table XV), the WR was positive in 118 sera (76 per cent.) and negative in 37, the PPR was positive in 103 (67 per cent.) and negative in 52, the TPI was positive in all of 20 cases, and the TWR was positive in 101 sera (83 per cent.) and negative in 21.

TABLE XV
SUMMARY OF FOLLOW-UP IN 103 CASES OF
"SYPHILIS OR YAWS"

Test	Result			Total No. of Sera Tested
	Negative	Positive		
		No.	Per cent.	
WR	37	118	76	155
PPR	52	103	67	155
TPI	0	20	100	20
TWR	21	101	83	122

In this group the TWR comes closer to the TPI in performance than do the standard STS. But this could be related to the fact that the TWR gave positive results in 41 per cent. of coloured controls.

FIG. 8.—"Syphilis or Yaws" (103 cases).

Non-Treponemal Reactors.—One or more TWRs were performed in 24 patients (nine white and fifteen coloured) diagnosed as non-treponemal reactors, in whom there was no clinical evidence of treponemal disease and the TPI was negative (Table XVIA and B). The TWR was positive in fifteen and negative in nine; the WR positive in thirteen and negative in eleven, and the PPR positive in only five and negative in nineteen.

TABLE XVIA
NON-TREPONEMAL REACTORS
(TPI NEGATIVE)

Sex	Male	Female	Total
Race			
White ..	4	5	9
Coloured ..	4	11	15
Total	8	16	24

TABLE XVIB
RESULTS OF TESTS IN NON-TREPONEMAL REACTORS

Tests			TWR	WR	PPR
Results	No. of Cases	Positive ..	15	13	5
		Negative ..	9	11	19
	No. of Sera	Positive ..	39	40	7
		Negative ..	30	29	62

With regard to sera which were tested in parallel by the three methods, the TWR was positive in 39 sera and negative in thirty, the WR positive in forty and negative in 29, and the PPR positive in only seven and negative in 62.

In this group, therefore, results of the TWR did not approximate to those of the TPI.

Summary

(1) The results of standard serological tests for syphilis (STS) were compared with those of the TWR in 4,105 patients attending the Whitechapel Clinic of The London Hospital between November 1, 1957, and October 31, 1958.

(2) In a selected group of 482 white patients with no evidence of treponemal infection, the initial TWR gave positive results in 21 cases (4.4 per cent.), whereas in a similar group of 58 coloured patients it was positive in 24 (41 per cent.).

(3) Reproducibility of the TWR was unsatisfactory.

(4) The results obtained in cases of syphilis which conformed to strict diagnostic criteria were analysed.

(5) In 24 non-treponemal reactors the TWR was positive in fifteen cases and negative in nine.

(6) There was considerable discrepancy between the results of the TWR and those of the WR, PPR, and TPI. This was apparently due to a high incidence of false positive reactions with the TWR.

(7) Because of the non-specificity and poor reproducibility of the TWR, this test in its present form cannot be regarded as a substitute for the TPI.

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