Quality control in bacteriology: preliminary trials¹

E. JOAN STOKES AND J. L. WHITBY²

From University College Hospital, London, and Queen Elizabeth Hospital, Birmingham

SYNOPSIS Six trials of quality control material sent in the form of artificial specimens are reported. A method of assessment using a computer for complex results, including antibiotic sensitivity tests, was employed. The trials were successful in showing the need for a comprehensive service and in bringing to light by this method a wide variety of errors.

If diagnostic laboratories are to maintain a high standard, quality control is essential. In chemical pathology and haematology it is already routine practice but in bacteriology test material is not readily available. Ampoules of dried cultures to check tests of identification can be purchased but are seldom employed because identification is only one of the many problems which face the diagnostic bacteriologist. Indeed when a potential pathogen has been isolated in pure culture half the battle is won.

If positive test material as close as possible to the specimens undergoing investigation were available for repeated tests throughout the year, it would enable bacteriologists to check their procedures and faulty methods of isolation as well as of identification would be disclosed.

The possibility of obtaining commercially prepared material in this form was investigated but was found to be impracticable and the alternatives are either to set up a separate quality control laboratory to manufacture 'specimens' or for the task to be undertaken by one or more practising laboratories; tests using the latter alternative are reported here. It has the advantage that the manufacturing laboratory has problems common to users of the service and can test its own materials under field conditions. Moreover suitable material for specimens is to hand and the strains sent out can be freshly isolated and are therefore likely to be nearer in their growth requirements and behaviour to wild strains.

These trials were made by a group of laboratories in various parts of England, whose directors were interested in establishing a regular service, in order to test the proposed methods of manufacture of 'specimens' and assessment of results and to estimate the cost. They were financed by the Department of Health. Participants were 13 public health laboratories, eight laboratories linked with teaching hospitals, and four non-teaching hospital laboratories (five in the last three trials). One of the teaching hospital laboratories which had previous experience in the ACP Sensitivity Test Trial (Association of Clinical Pathologists, 1965) sent the specimens and took part in the trial by posting a sample box to itself; another teaching hospital laboratory, experienced in computer techniques, received the results from all laboratories.

Regular testing is essential in quality control and participating laboratories gain more from the tests if they can compare their results with others examining similar material sent out at the same time. For this reason and because preparation in batches at regular intervals is less disturbing to working routine than test material on demand, 'specimens' were sent at approximately monthly intervals on three occasions, followed by an interval for assessment and then a further three trials. In order to give laboratories the same kind of information they should normally receive the 'specimens' were said to have come from appropriate sites, but no attempt was made to disguise them as anything other than quality control, nor do we regard this as necessary.

Methods

MANUFACTURE OF SPECIMENS

Six specimens were sent for each trial (three faeces, two swabs in transport medium, and one urine).

¹Other participants: Blowers, R., Bradley, J. M., Brumfitt, W., Crone, P. B., Cunliffe, A. C., Darrell, J. H., Dowsett, L. M., Furniss, A. L., Greenbury, C. L., Gruneberg, R. N., Hughes, M. H., Hurley, R., Hutchison, J. G. P., Jebb, W. H. H., Kingsley Smith, A. J., McCoy, M. H., Meers, P. D., Mohun, A. F., Mortimer, P. R., Speller, D. C. E., Taylor, C. E. D., Turner, G. C., Williams, R. E. O., and Willis, A. T.

²Present address: Department of Bacteriology and Immunology, University of Western Ontario, London, Canada.

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Faeces

Faeces sent for occult blood estimation from patients not receiving antibiotics was emulsified in about five times its volume of peptone water in an electric homogeniser. It was then distributed in 2 ml volumes, refrigerated overnight, and seeded on the day of dispatch with 2 drops, 0.04 ml, of diluted overnight culture of an intestinal pathogen. The culture was counted at the time of seeding by the Miles and Misra method so that the approximate number of viable pathogens per millilitre faeces suspension could be calculated. The aim was to grade the specimens so that one sample should yield a positive result on primary plates and one only after enrichment, the third being intermediate between them. The numbers of pathogens seeded per ml faeces suspension were 10^6 - 10^7 for the strong samples, 10^4 - 10^5 for the intermediate samples, and 10^2 - 10^3 for the weak samples.

Tests on extra samples of the faecal specimens sent in trials 4 and 6 were also made by three of the participating laboratories knowing with what they had been seeded. Good survival of *Salmonella* was demonstrated. It could be isolated from specimens kept at room temperature for at least two weeks. There was growth of *Salmonella* initially at room temperature (approximately 22°C) in samples heavily seeded at the expense of the normal commensals.

The possibility of using a suspending medium resembling faeces instead of the natural material has been considered, and in the fifth trial a sterile yeast suspension kindly supplied by Oxoid Limited as employed for the Chick-Martin test was used. This was seeded with *Sh. sonnei* in the same manner as the previous samples and to simulate natural conditions 3 drops of a peptone water culture of *E. coli* and 2 drops of diluted culture of *Proteus mirabilis* were added to the suspension before seeding with the pathogen, all bottles receiving the same amount of commensals. The results were disappointing since few laboratories were able to isolate the pathogen. This was probably because too much commensal was added which overwhelmed the pathogen in many cultures. Even if it were proved that this *Shigella* was abnormal and that most pathogens would survive in the yeast suspension, there are disadvantages in the method. If a natural specimen is to be simulated commensal organisms must be added and it would be necessary either to use stock tested strains or to do preliminary tests with freshly isolated commensals before preparing material for each trial. Moreover, one could not hope to cover the variety of strains of different commensals in natural faeces and if selective media are to be improved in the light of results from many

laboratories testing these 'specimens' over a long period one would be in danger of suiting media to an artificial set of circumstances.

Swabs

Swabs were dipped in a 1/10 dilution of overnight broth cultures of the pathogens and then placed in transport medium. When a mixed culture was sent the mixture was prepared and well mixed immediately before dipping.

Tests made at the end of the trials showed that the extra staphylococci occasionally reported may have been due to contamination from the hands of the person introducing the swab into the transport medium while breaking off the swab stick.

Normal urine was brought to the boil to kill contaminating organisms. After distribution in 2 ml volumes it was seeded with 1 drop overnight culture per bottle. In the last three trials sterile peptone water was used instead of urine.

In the first five trials the pathogens inoculated into the 'specimens' were sent in pure culture to the Queen Elizabeth Hospital, Birmingham, and to Hammersmith Hospital, London, where their identity and antibiotic sensitivity was checked and agreed with the distributing laboratory.

METHODS OF ASSESSMENT

The results of the first three trials were assessed without computer aid. At the end of the third trial it was considered desirable that an extension be considered, unfortunately funds, for instance, for assessing laboratory performance in all bacteriology laboratories in a region were not available, so it was proposed to repeat the trial with the same participants, to try to discover something about methods employed and to experiment with the use of computer facilities for a more rapid analysis and print-out of the results.

The computer aspects were dealt with by sending each participating laboratory punch cards in which their laboratory number had been punched and with allotted spaces for entry of results similar to those described (Whitby and Blair, 1970) for routine laboratory working (Fig. 1). On receipt of completed cards the recorded findings were punched into the same cards and fed to the computer, which produced in tabulated form a complete printout of all laboratories findings and a summary (Fig. 2). The tables were photocopied and then sent to all participants. Little difficulty was experienced with using the cards save that they were unsuitable for one of the samples submitted, where more sophisticated tests for antibiotic sensitivity required recording. Faecal specimens were not included in the computer process.

Top Card (Before Completion):

LAB	DATE	ORG	ANTIBIOTIC SENSITIVITY	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32
			S = SENSITIVE R = RESISTANT M = MOD. SENSITIVE (IF USED)	PEN	AMP	METH/CLOX	CARB	CEPHALOR	CEPHALEX	TETR	ERYTH	STREP	NEO/KANA	GENT	LINC/CLIND	NOVO		
			ORGANISM ISOLATED	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47
				BACI	RIFAMP	FUS.A	CHLOR	COL/POLY	NITROF	NAL.A.	SULPH	TRIMETH	TRI+SULPH	MYSTAT	AMPHO			

Bottom Card (After Completion):

LAB	DATE	ORG	ANTIBIOTIC SENSITIVITY	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32
	29.4.71		S = SENSITIVE R = RESISTANT M = MOD. SENSITIVE (IF USED)		S				S		R				S			
			ORGANISM ISOLATED SPECIMEN No. 6 PROTEUS MIRABILIS (mode positive)					R	S	S	S	S						
				BACI	RIFAMP	FUS.A	CHLOR	COL/POLY	NITROF	NAL.A.	SULPH	TRIMETH	TRI+SULPH	MYSTAT	AMPHO			

Fig. 1 Punch cards sent to participants before and after completion.

Results

FAECAL SPECIMENS

Three specimens of faeces were included in each trial. These consisted of graded numbers of a *Salmonella* species as outlined in the Methods section. In trial 2 an unseeded sample was included and the weakest seeding omitted and in trial 5 *Shigella sonnei* was submitted in yeast suspension. Excluding the results for trial 5, where the suspension proved unsuitable, the performance of laboratories showed considerable improvement in the later trials. Thus, in trials 1 and 3, 54 negative and 93 positive reports were received.

Thirty-three of the 54 negative reports were from the samples containing the least numbers of salmonella and 21 were from samples where *Salmonella* was present in considerable numbers. In trials 4 and 6, 22 negative and 122 positive reports were received, 18 of the 22 negative reports being from the samples that contained the least numbers of salmonella. Table I sets out the relative performance in trials 4 and 6 compared with trials 1-3, and also the performance of public health laboratories and hospital laboratories. Although public health laboratories had performed better in the first trials this no longer appeared to be the case and did

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QUALITY CONTROL TRIAL																					
DATE 28.04.71																					
SPECIMEN SIX																					
LABORATORY	ORGANISM ISOLATED	P	AM	M/CCAR	CP	CX	T	ER	ST	N/KGEN	ANTIBIOTIC SENSITIVITY				NF	NA	SU	TR	T+S	NY	AM
											L/CNV	BA	RI	FU	CH	PMX					
1	512	0		S			R										S	S	R		
2	510	0		S			R		S	S					R		S	S	S		
3	510	0		S			R		S	S							R	S	S		
4	512	0		S			R		S								R	S	S	S	
5	510	0		R	S		R										R	S	S	S	
6	510	0		S			R			S							S	S	S	S	
7	512	0		P	S		R		S									S	S		
8	512	0		S			R										R	S	S	S	
9	510	0		S													R	S	S		
10	510	0		S			R										S	S	R	S	
11	512	0		S		S	R		S					R			R	S	S		
12	512	0		S		S	R		S								R	S	S	S	
13	512	0		S			R		S								R		S	S	
15	512	0		S		S	R			S				R			S	S	S	S	
18	512	0		S													M		S	S	
20	512	0		S													M		S	M	
21	512	0		S			R							R			M	S	S	S	
22	510	0		S			R							S			R	S	R	S	S
23	512	0		R			R										R	S	S	S	
26	510	0		S			R										R		S	S	S

ANTIBIOTIC	TOTAL NO. TESTED	NO. RECORDED SENSITIVE	PCENT. RECORDED SENSITIVE
PENICILLIN	2	0	0.0
AMPICILLIN	20	19	95.0
CEPHALORDN	2	2	100.0
CEPHALEXIN	1	1	100.0
TETRACYCLN	17	0	0.0
STREPTOMCN	4	4	100.0
NEO. KANA.	5	5	100.0
GENTAMYCIN	2	2	100.0
COL. POLYMX	5	1	20.0
NITROFURNT	19	8	42.1
NALADIXIC	16	16	100.0
SULPHONAMD	20	17	85.0
TRIMETH.	5	5	100.0
TRIMETH+SU	11	11	100.0

ORGANISM CODE	ORGANISM	NO. OF ISOLATIONS
510	PROTEUS SP	8
512	P. MIRABIL.	12

NUMBER OF TIMES ALTERNATIVE NAME USED	0
NUMBER OF TIMES MODERATLEY SENSITIVE IS USED	
NITROFURNT	3
TRIMETH.	1
NCARD	21

***

Fig. 2 Computer printout (detail and summary).

Trials 1—3 ¹			Trials 4 and 6 ²		
Number of Failures (8 positive samples per laboratory)	Public Health Service Laboratories	Hospital Laboratories	Number of Failures (6 positive samples per laboratory)	Public Health Service Laboratories	Hospital Laboratories
0	5	0	0	7	4
1	3	1	1	3	5
2	0	2	2	0	4
3	0	2	3	2	1
4	3	5	4	0	0
5	1	3	5	0	0

Table I Number of times laboratories failed to isolate *Salmonella* spp from positive samples in the two halves of the trial¹One public health and two hospital laboratories failed to report.²One hospital laboratory failed to report.

perhaps represent improved performance in hospital laboratories.

ISOLATION AND IDENTIFICATION OF BACTERIA FROM SWABS AND 'URINE'

No difficulty in the isolation and identification of *Staph. aureus*, *E. coli*, *Proteus*, or *Pseudomonas* was encountered. In trial 2, two strains of *Staph. aureus* were included in one sample and this sample was not well handled. Sixteen laboratories found a single strain, eight two strains, and one three strains. In trial 3 *Cl. welchii* was included with *Staph. aureus* in a swab said to have come from a compound fracture caused by a road traffic accident. Two laboratories of 24 failed to isolate the anaerobe and one other failed to identify it correctly. In trial 6 *Cl. welchii* was sent alone on a swab said to have come from the uterus of a patient with septic abortion. Four laboratories failed to isolate it although subsequent tests with this strain showed it to be viable in transport medium for at least three days. On this occasion all those isolating it identified it correctly.

Klebsiella aerogenes was sent in urine in trial 6. Correct identification was reported by nine laboratories, *Klebsiella* species by seven, *Klebsiella pneumoniae* by one, *Enterobacter* species by one, and five falsely reported *Escherichia coli*.

Out of 409 isolations of seven species there were only nine incorrect identifications. Nomenclature varied and some laboratories identified as far as the genus only, especially for urinary pathogens.

ANTIBIOTIC SENSITIVITY TEST RESULTS

It was to be expected that more difficulty would be encountered in testing some antibiotics than others. Most of the errors reported in the A.C.P. Antibiotic Sensitivity Test Trial (1965) occurred in tests of penicillinase-producing staphylococci with penicillin which were falsely reported sensitive and false reports of resistance to sulphonamide. Also *Proteus*

mirabilis was sometimes falsely reported sensitive to nitrofurantoin. These errors were again most commonly made and in addition some laboratories had difficulty in recognizing methicillin resistance of *Staph. aureus* and carbenicillin sensitivity of *Pseudomonas aeruginosa* which were not previously tested.

In these trials penicillinase-producing *Staph. aureus* was reported penicillin sensitive eight times in 184 tests. Resistance to sulphonamide was wrongly reported 25 times in 128 tests of urinary pathogens. In 13 of 39 tests *Proteus mirabilis* was reported sensitive to nitrofurantoin although the minimal inhibitory concentration of the two strains sent out was greater than 128 µg, that of the sensitive *E. coli* control being 16 µg per ml. The results for methicillin sensitivity of *Staph. aureus* are seen in Table II.

Trial	No. of Tests Reported	No. Reported Sensitive	MIC ¹ (µg/ml)
1	21	4	64
3	19	4	> 64
4	22	0	> 64
6	23	2	> 64
			Control 1

Table II Individual laboratory reports of sensitivity of methicillin-resistant *Staph. aureus*¹Plate dilution technique with heavy inoculum, Oxford staphylococcus control.

Carbenicillin sensitivity of *Pseudomonas* is not easy to determine because the species is never very sensitive although it is treatable by large doses of this penicillin. It is, however, essential that laboratories should be able to distinguish between strains which are treatable and those which are too highly resistant to respond. The same sensitive strain was sent on two occasions in trials 3 and 5. There were nine reports of resistance out of 44 tests, eight of them occurring in trial 3. The improvement was probably the result of discussion among the parti-

cipants at the end of the first three trials about the meaning of resistance in this species and the recommendation that a sensitive *Pseudomonas* should be tested in parallel (Waterworth, 1969). No difficulty was encountered with gentamicin tests.

Streptococcus faecalis is notoriously difficult to test to penicillins (College of Pathologists of Australia, 1968). A strain recently isolated from a blood culture happened to be available and was sent in trial 5. The results are seen in Table III.

	Sensitive	Moderately Sensitive	Resistant	MIC ($\mu\text{g/ml}$) ¹	
				Strept. faecalis	Control
Penicillin	2	3	19	1.0	0.03
Ampicillin	19	4	1	1.0	0.06

Table III Reporting of sensitivity of *Strept. faecalis* to penicillins by 24 laboratories

¹Plate dilution technique Oxford staphylococcus control

In trial 6 the strain of *Klebsiella* was moderately resistant to a number of antibiotics but deemed treatable with higher dosage. Scoring errors would be difficult but Table IV sets out the findings of the various laboratories for six antibiotics; there are marked differences but the findings would not indicate overreporting of ampicillin sensitivity (cf Table III).

No attempt has been made to score the total number of errors in examination of swabs and urine because in some of the samples scoring becomes too arbitrary, nor has any attempt been made to compare one laboratory with another, and the results have been returned to laboratories for their own comments and action.

LABORATORY METHODS

While with faeces there does seem to have been a significant improvement in performance in the later three trials, it would be hard to conclude that there had been much change with antibiotic sensitivity

testing since the number of laboratories reporting discrepant results remained much the same. In an attempt to examine some of the reasons for discrepant findings in sensitivity tests a short questionnaire was circulated to participating laboratories; unfortunately, because of a desire not to ask a leading question, the answer to the degree to which testing was controlled did not emerge clearly for all laboratories.

The use of control organisms was mentioned by seven laboratories, queried by one, and not mentioned by the other 16. Individual discs were the most popular method for applying antibiotics to the plate, being used exclusively in eight laboratories and for part of the routine by a further 12. Twelve laboratories used Multodiscs and four impregnated rings. Oxoid DST with or without the addition of lysed blood was the most popular medium for testing antibiotic sensitivity routinely. Special conditions to determine methicillin sensitivity (Hewitt, Coe, and Parker, 1969) were used in 21 out of 24 laboratories. Table V summarizes the amount of antibiotic in micrograms incorporated in the discs or rings for sensitivity testing.

Table VI shows the number of laboratories performing direct sensitivity tests on swabs. The point of the question was to relate it to the findings in trial 2, question 5, where, when two strains of *Staph. aureus* were submitted on a single swab, only nine out of 25 laboratories submitting answers found both organisms; seven of these performed direct tests routinely and two sometimes did so.

Table VII outlines the methods employed in participating laboratories for faeces. It can be seen that the only discrepancies are in the number of laboratories using additional methods. It is notable how many laboratories do not check the efficiency of fresh batches of culture medium but there was no correlation between this fact and laboratory performances. Three laboratories which successfully isolated salmonella from all specimens did not use any control procedure. Nor again did laboratories picking a single example of each colonial type fare

Antibiotic	Report				MIC ($\mu\text{g/ml}$)	
	Sensitive	Moderately Sensitive	Resistant	Not Tested	Kleb. aerogenes	Sensitive Control <i>E. coli</i>
Ampicillin	4	7	14	0	16	4
Cephalosporins	7	1	0	17	8 ¹	2 ¹
Sulphonamide	17	2	6	0	32	8
Tetracycline	18	0	2	5	2	0.25
Nitrofurantoin	21	2	2	0	64	16
Naladixic acid	19	0	1	5	4	1

Table IV Reporting of antibiotic sensitivity of *K. aerogenes* in trial 6 by 25 laboratories

¹Cephaloridine

Anti- biotic in Disc (µg)	Pen ¹	Amp	Met	Clo	Car	Cep	Cepx	Str	Kan	Neo	Gen	Tet	Chl	Pol ¹	Col	Ery	Lin	Clin	Fuc	Nov	Bac	Nit	Nal
1	15			1				2			4					1	12	12		9	1		
2	3	8																					
5	5	4		5		13 (1)	1		8			2	1			10							
10		2	20	1		1		13 (1)		13	17 (10)	14	7			11	4		21		9		
25		9 (15)			5	7 (12)	10 (5)	6 (5)				6	7										
30		1							14 (13)	3	(1)									2			
50								2		1		2	8 (4)	(1)	11 (1)	2				1		1	24
100					17 (23)										4								
200															2	7 (10)						22	
300															3 (5)								
3,000																							1
Not used	1	0	4	17	2	3	13	1	2	7	3	0	1	15	6	0	8	12	3	12	14	1	0

Table V Number of laboratories employing paper discs of various potencies for antibiotic sensitivity testing

¹Units²Where figures are in brackets laboratory has indicated a different strength for use with organisms isolated from urine

Direct Sensitivity Testing			No. of Colonies Picked for Sensitivity Tests				
Routine	Sometimes	No	1	2-4	>5	Sweep	Direct Tests Only
14	8	2	10	4	4	4	2

Table VI Swab routine in 24 laboratories

Primary Culture Media			Enrichment Broths		Number of Colonies of Each Type Picked for Identification					Control of Performance of Culture Media		
Desoxy- cholate Citrate	Mac- Conkey	Wilson and Blair	Selenite	Tetra- thionate	Rappaport	1	2-5	10	Serology Positive	Regular	Sometimes	None
24	16	5	24	5	4	15	6	1	2	8	7	9

Table VII Laboratory routine for faeces in 24 laboratories employing various techniques

Quantitative Procedure Employed					Nature of Specimens Accepted			
None	Dip Slide	Blotting Paper	Standard Loop	Modified Miles and Misra Viable Count	Fresh or Refrigerated	Insist on a Fresh Specimen	Same Day	All Processed
1	6	7	12	6	8	12 ¹	2	2

Table VIII Urine routine in 24 laboratories

¹Two laboratories insisted but not rigidly

any worse than those picking more, in this small sample.

Table VIII shows the quantitative routine used for urines. Despite much criticism of 'standard loops' in the past the use of such a loop was the most popular quantitative method, most laboratories made some attempt to obtain a fresh specimen of urine on which to perform their quantitative routine.

Discussion

If quality control is to have any impact or meaning in laboratories it is important that it should be treated in a routine manner, yet the time taken for some reports to be received was as long as two months and some reports have never been received. This is clearly shown by the variations in the number

of reports for analysis in the different trials. It is of course equally important to put work through a laboratory when the director is away as when he is present, and most routine work is dealt with within 48 hours of receipt. Some delay in our trial arose because laboratories performed 'phage typing on staphylococci and serology on salmonellae but in those circumstances as in clinical practice an interim report should have been sent. Specimens were usually posted on Tuesdays, and, even allowing for the postal services must have been received by Friday in all but the most exceptional circumstances, we would therefore have expected to receive all the results by the end of the following week. Using the computer for the tables would have then meant returning results to participants within 14 days of the initial receipt of the specimens. In the last three trials when detailed records were kept there were signs of improvement as in trial 4, seven laboratories, trial 5, five laboratories, and trial 6 only two laboratories submitted results later than the twelfth day after specimens had been posted to them.

The object of quality control is to improve performance but these trials are not sufficiently extensive to be able to assess improvement satisfactorily. It appears from Table I that there was some improvement in the isolation of salmonella from faeces, especially in hospital laboratories, but with the exception of improvement in testing *Pseudomonas* sensitivity to carbenicillin and perhaps a little improvement in the recognition of methicillin-resistant *Staph. aureus* there was no appreciable difference in performance in the later trials.

It might have been expected that the false reports of sensitivity of *Strept. faecalis* to ampicillin (Table III) was due to the use of high content discs but this cannot have been the only factor since seven laboratories reporting sensitivity were using 2 µg discs for all tests. Ten laboratories reported further tests of sensitivity to antibiotic combinations, notably penicillin with streptomycin. In these circumstances even a preliminary report of sensitivity to ampicillin may be dangerous as treatment with this drug in normal dosage will not succeed in *Strept. faecalis* endocarditis.

There was evidence of lack of knowledge of the normal resistance of species to antibiotics: for example, one laboratory identified *Proteus mirabilis* correctly and reported it sensitive to colistin. Another reported *Klebsiella* in urine sensitive to fusidic acid. A further correct report of an unnecessary test was *Pseudomonas* in urine resistant to methicillin.

These trials had the advantage that there were few participants who could meet and discuss results and this probably played a large part in bringing particular faults to the notice of directors who could then discuss remedies for them. In a comprehensive service this would not be possible and some substitute for it, perhaps a quarterly bulletin from manufacturers of specimens and assessors is desirable. This need not interfere with anonymity which should be maintained.

The results show that quality control by sending known positive specimens brings to light a wide variety of errors. In all tests, except sensitivity of *Strept. faecalis* to ampicillin, most of the results were correct. Extra tests of validity undertaken by the manufacturer of specimens and other participants in these trials would probably not be needed in a comprehensive service.

We thank Mrs J. N. Blair for writing the computer programme and handling computer data and Miss P. M. Waterworth for estimating the MICs of antimicrobials for the bacteria sent.

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