An automated method for serum bilirubin determination

N. A. SIMMONS

From the Department of Clinical Pathology, Guy's Hospital, London

SYNOPSIS A method for the determination of both direct and indirect serum bilirubin on the AutoAnalyzer is described which has a number of advantages over the automated method that is currently recommended. It gives total bilirubin results which are similar to those obtained with the equivalent manual method and direct bilirubin results which are usually about 10% lower.

The effect of haemolysis on estimations was investigated. It has little influence on the total, but significantly lowers the direct bilirubin results.

Simple methods for the preparation of true bilirubin standards and a stable control serum are described.

In a very careful study Nosslin (1960) showed that the method of Jendrassik and Grof (1938) for the estimation of total serum bilirubin had many advantages over other methods and that a modification of it gave a direct reaction which was the best available index of conjugated bilirubin in the serum.

In the total bilirubin method the serum was mixed with a buffered caffeine solution which acted as an accelerator and then with diazo reagent. The azo pigment formed was converted from red to blue by the addition of alkali. In the direct reaction the accelerator solution was replaced by water and the diazo reaction terminated by ascorbic acid before treatment with alkali. Serum blanks were prepared with diazo I instead of mixed diazo reagent. The methods currently recommended for the determination of bilirubin on the Technicon AutoAnalyzer (Gambino and Schreiber, 1964) are based on these procedures.

Michaëllsson (1961) demonstrated a number of disadvantages in Nosslin's method. First, caffeine was found to reduce the optical density of alkaline azobilirubin solutions. Since it was used in the total bilirubin, but not direct bilirubin estimations, and since the amount of direct-reacting bilirubin was calculated from data obtained by putting standards through the total bilirubin procedure this could cause misleading results. Secondly, ascorbic acid and diazo reagent react to form a coloured product in the direct reaction. They were not present in the blanks for which values were consequently low. Thirdly, haemolysis reduced the values for total bilirubin.

Nosslin (1960) did not use ascorbic acid in the total bilirubin procedure. Michaëllsson (1961) showed that its inclusion minimized the effect of haemolysis. He went on to show that the other disadvantages could be overcome if all three reagents, ascorbic acid, diazo reagent, and accelerator, were all included in all three estimations, total bilirubin, direct bilirubin, and blanks. By varying the order in which the reagents were used he devised a method in which this was done and the automated method described in this paper is based on it. In addition, simple and convenient methods for the preparation of bilirubin standards and control serum are described here and the effect of haemolysis upon direct and total bilirubin determinations is investigated.

METHOD

PRINCIPLE In the determination of total bilirubin the serum is added to a caffeine reagent, which acts as an accelerator, and then mixed with combined diazo reagent. The diazo reaction is terminated by the addition of ascorbic acid which destroys the excess diazo reagent, the azobilirubin is made alkaline by the addition of a tartrate buffer, and the intensity of the colour read at 600 m\mu.

To determine the direct-reacting bilirubin the serum is mixed with water and diazo reagent. The reaction is terminated by ascorbic acid and then caffeine is added. Again the azobilirubin is made alkaline with tartrate buffer.

When blanks are run the serum is mixed with water and ascorbic acid. Diazo reagent, caffeine, and tartrate reagent are added subsequently.

1Present address: Department of Pathology, Chase Farm Hospital, Enfield, Middlesex.

Received for publication 21 August 1967.
An automated method for serum bilirubin determination

REAGENTS The following are required:

Ascorbic acid A 0·8% solution is made up each day by dissolving 400 mg in 50 ml distilled water. (This was found to be sufficient for total and direct bilirubin determinations on at least 20 specimens.)

Caffeine mixture Hydrated sodium acetate, 84 g, 50 g sodium benzoate, and 33 g caffeine are dissolved in approximately 800 ml distilled water at 50° to 60°C. This is made up to 1,000 ml when cool.

Diazo I solution Sulphanilic acid, 5·0 g, is dissolved in distilled water containing 15 ml concentrated hydrochloric acid (specific gravity 1·19) and the volume made up to 1,000 ml.

Diazo II solution Sodium nitrite, 0·5% in distilled water, is stored in a dark bottle and renewed every 14 days.

Diazo reagent Diazo II, 1·25 ml, is added to 50 ml diazo I. This is stable for 24 hours (Gambino and Schreiber, 1964) and is made up each day.

Alkaline reagent (§ strength Fehling’s II solution) Sodium hydroxide, 66 g, and 217 g sodium potassium tartrate are dissolved in distilled water and the volume made up to 1,000 ml.

Distilled water BRIJ-35 (Technicon), 0·5 ml, may be added to each litre if it is to be run through the manifold.

STANDARDS The method for their preparation is a modification of that described by Gadd (1966). A 5% solution of bovine albumin in 0·1 M phosphate buffer, pH 7·4, is prepared. It is stored in the refrigerator. The buffer is made by dissolving 13·97 g anhydrous dipotassium hydrogen phosphate and 2·69 g anhydrous potassium dihydrogen phosphate in distilled water and making the volume up to 1,000 ml.

Bilirubin (molar absorbivity 60,700 ± 800, British Drug Houses Ltd), 50 mg is dissolved in 50 ml dimethylsulphoxide. Without delay 7·5 ml of this solution is made up to 50 ml and 10 ml made up to 100 ml with the albumin solution. These standards contain 15 mg and 10 mg bilirubin per 100 ml respectively. Standards containing 5·0, 2·5, 1·25, and 0·6 mg per 100 ml of albumin solution are prepared from the 10 mg per 100 ml standard.

At the same time as the standards are prepared, the concentration of bilirubin is checked spectrophotometrically as follows: 1·0 ml of the solution of 50 mg bilirubin in 50 ml dimethylsulphoxide is added to 9·0 ml dimethylsulphoxide and 0·2 ml of this solution added to a further 3·8 ml dimethylsulphoxide (total dilution 1:200). The optical density of this solution at 453 m$log$ is read in a 1 cm cuvette against a blank of dimethylsulphoxide. The optical density is multiplied by 19·26 (based on a molar absorbivity of 60,700) to give the concentration of bilirubin in the standard expected to contain 10 mg per 100 ml. The molar absorbivity was confirmed by observation.

This method differs slightly from that of Gadd (1966). To check the bilirubin concentration spectrophotometrically he used a dilution in dimethylsulphoxide of the 10 mg per 100 ml standard which contains albumin, whereas the solution used to check the concentration in this method contains none. His method was found to be unsatisfactory, for the small amount of albumin in the 10 mg standard increased the optical density of the bilirubin.
rubin solutions and changed the peak of the bilirubin absorption curve from 453 m\(\mu\) to 458 m\(\mu\).

Standards are dispensed into small quantities and stored in the dark at \(-20^\circ C\). They are stable for at least 30 days.

**CONTROL SERUM** This is prepared from discarded blood removed at exchange transfusion from babies with haemolytic disease of the newborn who have a pre-exchange serum bilirubin greater than 10 mg per 100 ml. Blood diluted with heparinized saline during the exchange transfusion is suitable. Plasma is separated by centrifugation from approximately 500 ml. To render it clear and free from clots it is first stored at 4°C for three days. Large clots are removed and it is transferred to a deep freeze at \(-20^\circ C\). After 24 hours it is thawed, centrifuged, and the supernatant cleared by passing it through a sintered glass filter ("grade 3"). The value for the bilirubin concentration is determined by repeated estimations using commercial serum as a control. The material is dispensed into small quantities and stored in the dark at \(-20^\circ C\). It is stable for at least 10 months.

**OPERATING PROCEDURE** Samples are analysed at a rate of up to 70 per hour. The sample line aspirates 0-16 ml per min, therefore about 0.1 ml of sample is aspirated for each of the three runs, total and direct bilirubin and blank. For specimens containing more than 15 mg bilirubin per 100 ml the serum is diluted 1:3 with distilled water. It is usually easy to see with the naked eye specimens containing more than 2 mg per 100 ml, and to avoid carry-over these can be placed at the end of the run or followed by a sample cup containing distilled water. A 15 mm tubular flowcell and 600 m\(\mu\) filters are used. A recorder chart speed of 18 inches per hour is suitable and chart paper calibrated to show optical density is used. Reagents are aspirated in a different order for the estimation of total and direct bilirubin and blanks (Fig. 1). Changing the order is simplified by mounting the manifold on a platter and numbering the lines. Between each different type of estimation water is washed through the manifold rapidly for about a minute. Although blanks usually give low values, this is not so if the samples are cloudy or haemolysed and for accurate results they should be run routinely.

To read the results the optical density of the standards' blanks is subtracted from the optical density when their total bilirubin is estimated and a horizontal line is drawn near the top of their peaks to show the corrected value (Fig. 2). These points are used to draw the standard curve on the chart reader and this is used to read the total and direct bilirubin of the samples and their blanks. The figures obtained for the total and direct bilirubins of the samples are corrected by subtracting from them the values obtained for the blanks.

**RESULTS**

Unless stated otherwise tests were performed on plasma from heparinized blood.

**ACCURACY AND REPRODUCIBILITY** Twenty-five total bilirubin determinations were performed on different days on a control serum prepared in the laboratory by the method described above. The mean concentration was 7.3 mg per 100 ml, the standard deviation 0.27 mg per 100 ml, and the coefficient of variation 3.7%.

Estimations of the total bilirubin content of four commercial control sera were made. Each serum was put through six consecutive times in a single run. Individual values obtained with each serum varied from the mean by less than 2%. The mean values were close to those given by the manufacturer (Table I).

**TABLE I**

<table>
<thead>
<tr>
<th>Serum No.</th>
<th>Manufacturer's Method</th>
<th>Total Bilirubin (mg/100 ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Manufacturer's Value</td>
</tr>
<tr>
<td>I</td>
<td>Weight</td>
<td>0.65</td>
</tr>
<tr>
<td>II</td>
<td>King and Wootton (1956)</td>
<td>2.8</td>
</tr>
<tr>
<td>III</td>
<td>Weight</td>
<td>5.0</td>
</tr>
<tr>
<td>IV</td>
<td>Malloy and Evelyn (1937)</td>
<td>7.1</td>
</tr>
</tbody>
</table>

Six consecutive determinations of direct bilirubin were performed in one run on a specimen known to have a high value. Individual values varied from the mean of 7.7 mg per 100 ml by less than 1%.
Determinations of total bilirubin were made on pooled human serum whose initial bilirubin concentration was known and to which bilirubin in 5% bovine albumin had been added. Recoveries varied between 95 and 102% (Table II).

### Table II

<table>
<thead>
<tr>
<th>Bilirubin Added (mg/100 ml)</th>
<th>Theoretical Value (mg/100 ml)</th>
<th>Determined Value (mg/100 ml)</th>
<th>Bilirubin Recovered (mg/100 ml)</th>
<th>Percentage Recovery</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.0</td>
<td>1.15</td>
<td>1.1</td>
<td>0.95</td>
<td>95</td>
</tr>
<tr>
<td>2.5</td>
<td>2.65</td>
<td>2.7</td>
<td>2.55</td>
<td>102</td>
</tr>
<tr>
<td>5.0</td>
<td>5.15</td>
<td>5.0</td>
<td>4.85</td>
<td>97</td>
</tr>
</tbody>
</table>

Comparison with a Manual Method

The manual method employed was in routine use at Guy's Hospital and differed from the method recommended by Michaelsson (1961) only in that a caffeine solution was used as the accelerator instead of a diphylline mixture. Total bilirubin determinations were performed on 103 specimens and direct bilirubin determinations on 100 specimens by the manual and automated methods.

Fifty-six specimens were found to have a total bilirubin concentration of more than 0.5 mg per 100 ml by both methods and these results are plotted on a scattergram (Fig. 3) which shows that the two methods gave similar results. With values between 1-0 and 2-0 mg per 100 ml, the two methods occasionally differed by up to 0.4 mg per 100 ml, but the results of manual estimations in this range performed by different workers on the same specimen occasionally differed from each other by the same amount; repeated automated determinations in this range varied by no more than 0.1 mg per 100 ml. Forty-seven specimens were found to have a total bilirubin concentration of 0.5 mg or less per 100 ml. In 37 of these the results obtained with the two methods differed by no more than 0.1 mg per 100 ml. In nine they differed by 0.2 mg and in one by 0.3 mg per 100 ml.

Twenty-eight specimens were found to have a direct reacting bilirubin of more than 0.5 mg per 100 ml by both methods and these results are shown in Figure 4. In this range the automated method tended to give values approximately 10% lower than the manual method.

Seventy-two specimens were found to have a direct reacting bilirubin of 0.5 mg or less per 100 ml. In 54 of these results obtained with the two methods differed by no more than 0.1 mg per 100 ml. In 14 they differed by 0.2 mg and in four by 0.3 mg per 100 ml.

Normal Range

The total and direct reacting bilirubin concentrations were determined in 100 specimens from blood donors. Figure 5 shows the distribution of the total bilirubin results. In 95 specimens the concentration was between 0.2 mg and 0.8 mg per 100 ml. No direct reacting bilirubin was
detected in 44 of the specimens. In 55 of the others the concentration was found to be not higher than 0-1 mg per 100 ml. In one specimen it was 0-2 mg per 100 ml.

THE EFFECT OF ADDED HAEMOLYSED BLOOD

A solution containing 10 g haemoglobin per 100 ml, prepared from pooled human erythrocytes lysed with distilled water, was added to sera containing bilirubin to give final haemoglobin concentrations of 1·0 g, 0·5 g, and 0·1 g per 100 ml. Controls were prepared by the addition of distilled water to the sera instead of haemoglobin solution and determinations of total and direct bilirubin carried out.

Some typical results are shown in Table III.

<table>
<thead>
<tr>
<th>Haemoglobin Concentration (g/100 ml)</th>
<th>Total Bilirubin mg/100 ml</th>
<th>Percentage of Control</th>
<th>Direct Bilirubin mg/100 ml</th>
<th>Percentage of Control</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1-0</td>
<td>8·3</td>
<td>91</td>
<td>1·9</td>
<td>27</td>
</tr>
<tr>
<td>0·5</td>
<td>8·7</td>
<td>96</td>
<td>3·8</td>
<td>54</td>
</tr>
<tr>
<td>0·1</td>
<td>9·0</td>
<td>99</td>
<td>6·3</td>
<td>90</td>
</tr>
</tbody>
</table>

Addition of lysed red cells had little effect on the total bilirubin results. However, the direct-reacting bilirubin was significantly reduced and when haemoglobin was present even in the lowest concentration the results obtained were approximately 10% lower than in the control. Similar results were obtained when determinations were performed by the manual method.

When haemoglobin was present in plasma, values for the blanks were significantly increased and their routine inclusion was essential since small amounts of haemoglobin were difficult to detect when the bilirubin level was raised.

DISCUSSION

The manual method of Jendrassik and Grof (1938) for the determination of total bilirubin is usually regarded as the best available (With, 1954). Nosslin (1960) devised an excellent method based on it for the quantitative determination of direct-reacting bilirubin and the methods currently recommended for the determination of total and direct bilirubin on the AutoAnalyzer are based on these procedures (Gambino and Schreiber, 1964). Michaëlsson (1961) pointed out a number of disadvantages in the Nosslin-Jendrassik-Grof procedures and introduced modifications which overcame them. The automated method described in this paper is based on Michaëlsson's procedure. It gave total bilirubin results similar to those obtained with a manual method, but the direct bilirubins were usually lower. This was not unexpected for colour development in the direct diazo reaction may be rapid for an initial period of about one minute, but then continue at a slower rate for at least 45 minutes (Nosslin, 1960), and the time allowed for diazotization in the automated method is only about three minutes compared with 10 minutes in the manual method.

The amount of direct-reacting bilirubin found in normal serum by this method was small and in 99 of the 100 specimens from blood donors it was no higher than 0-1 mg per 100 ml. However, it must be emphasized that levels slightly above this are not necessarily due to an increase in conjugated bilirubin and may simply reflect a much larger increase in total bilirubin (Nosslin, 1960).

The instability of bilirubin in strong alkali has always been a source of difficulty in the preparation of standards. Gadd (1966) found that it was more stable in dimethylsulphoxide and, slightly modified, his method of preparation was found to be satisfactory.

In automated procedures the regular use of a control serum is desirable. Commercial preparations may be used, but if they are included in each run they may prove to be expensive. The bilirubin control serum described in this paper was found to be very easy to prepare and stable for at least 10 months.

The present investigation showed that haemolysis has little effect upon the total bilirubin measured by the automated method and the results are in agreement with those of Michaëlsson (1961). However, it
was found that the amount of direct-reacting bilirubin was significantly reduced in the presence of even small amounts of haemoglobin. Failure to recognize this fact could lead to a serious underestimation of the proportion of conjugated bilirubin in the serum.

I wish to thank Dr. J. Liddell and Mrs. C. Michael for their help and advice.

The January 1968 Issue

THE JANUARY 1968 ISSUE CONTAINS THE FOLLOWING PAPERS

Consistency in the histological diagnosis of epithelial abnormalities of the cervix uteri J. COCKER, H. FOX and F. A. LANGLEY

Morphogenesis of testicular tumours J. C. CROOK

Sensitivity of pencillase-forming strains of Staphylococcus aureus and of their pencillase-negative variants to cephaloridine, cephalothin, methicillin and benzylpenicillin J. H. HEWITT and M. T. PARKER

Effect of light on the Lactobacillus casei microbial assay BARBARA B. ANDERSON and JEAN D. COWAN

Absorption of antibiotics during peritoneal dialysis in patients with renal failure A. C. BUCK and SIMON L. COHEN

Classification of Staphylococcus albus strains isolated from the urinary tract R. G. MITCHELL

Epidemiological aspects of Proteus infections with particular reference to phage typing DIANA R. FRANCE and N. P. MARKHAM

Drug sensitivity of Proteus species C. T. HUANG and GRACE CHOU

Technical methods

Demonstration of Paneth cell granules using Naphthalene Black DIANA BOWER and C. G. CHADWIN

A fluorescent technique for demonstrating treponemes in films made from suspected chancrens M. F. GARNER and J. H. ROBSON

Recovery of human foetal liver cells after storage in liquid nitrogen A. J. ZUCKERMAN, H. E. M. KAY, and A. B. HOCKLEY

Rapid method for the estimation of plasma haemoglobin levels C. M. CRIFFS

Specific determination of plasma and urinary lactose P. A. TOSELAND

A simple device for microdialysis A. A. CODD

Letter to the Editor

Journal of Medical Microbiology

Postgraduate education

Book reviews

Copies are still available and may be obtained from the PUBLISHING MANAGER

BRITISH MEDICAL ASSOCIATION, TAVISTOCK SQUARE, W.C.1, price 18s. 6d.