Micromethod for the determination of uric acid in biological fluids

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SYNOPSIS  A micro procedure for estimating uric acid in body fluids, based on the carbonate method, is described. Interferences are elucidated. The stoichiometry of the reaction is carefully worked out. Reliability of the method is established by comparison with two other methods, by recovery experiments, and by replicate analyses.

Most methods for the quantification of uric acid in biological fluids are based on the reducing power of this hexavalent body. Uric acid has the ability to reduce the hexavalent tungsten of phosphotungstic acid to lower valences to form a blue colour. In principle, the reaction is similar to the one used for the estimation of glucose by the copper reduction technique and to the estimation of inorganic phosphorus. Since the resulting system is basically unstable, many workers have attempted to stabilize the final colour and prevent turbidity (Kern and Stransky, 1937; Caraway, 1955; Archibald, 1957; Henry, Sobel, and Kim, 1957; Eichhorn, Zelmanowksi, Lew, Rutenberg, and Fanias, 1961).

This article is devoted to an elucidation of a modification of the carbonate method of Caraway (1955) resulting in smaller sample requirements, greater adherence to Beer's law, and much more rapid development of maximal colour with no loss in colour stability, thus making it more attractive for routine use.

REAGENTS

Chemicals of A.R. quality should be used where possible.

1 WORKING DEPROTEINIZING REAGENT  Dissolve 1:19 g. of sodium pyrophosphate (Na₃P₂O₇·10H₂O) and 15 mg. of metaphosphoric acid in about 50 ml. of water. Slowly add 1-45 ml. of concentrated sulphuric acid. Cool in a water bath to 40°C. Slowly add 5-55 g. of sodium tungstate (Na₂WO₄·SH₂O) dissolved in about 400 ml. of water. Make up with water to 1 litre. This solution is stable for at least five years at room temperature.

2 10% (w/v) SODIUM CARBONATE  Dissolve 100 g. of anhydrous sodium carbonate in and dilute to 1 litre with water. Store in a polyethylene bottle. This solution is stable for at least one year at room temperature.

3 PHOSPHOTUNGSTIC ACID REAGENT  Dissolve 10 g. of sodium tungstate (Na₂WO₄·2H₂O) in 80 ml. of water. Add 8 ml. of 85% phosphoric acid. Boil gently with the aid of a reflux condenser for two hours. Allow to cool to room temperature and transfer to a 1 litre volumetric flask with the aid of water. Add about 400 ml. of water and dissolve 16 g. of lithium sulphate in the mixture. Dilute to 1 litre with water. Store in an amber glass bottle. This solution is stable for at least five years at room temperature.

4 URIC ACID STANDARD  For the stock standard (10 mg. per 100 ml.) dissolve 100 mg. of anhydrous uric acid and 50 mg. of lithium carbonate in about 150 ml. of water (warm if necessary). When completely dissolved, add about 500 ml. of water, 2-5 ml. of 40% formalin, and 0·3 ml. of glacial acetic acid. Make up to 1 litre with water. Store in an amber glass bottle. This solution is stable for at least one year at room temperature.

METHOD

To 2·7 ml. of working deproteinizing reagent in a test tube (13 x 100 mm.) add 0·3 ml. of unhaemolysed serum, cerebrospinal fluid, or diluted urine (1 : 10 with water). Mix well by lateral shaking. Centrifuge at moderate speed for five minutes. Pipette 2·0 ml. of supernatant fluid and 1·0 ml. of 10% (w/v) sodium carbonate into a round cuvette and mix. Let it stand for 10 minutes. Add 10 ml. of phosphotungstic acid reagent and immediately mix by lateral shaking. Within the next five to 23 minutes, read the absorbance at 700 mµ against a partial reagent blank prepared by treating 2·0 ml. of water exactly as the supernatant fluid. Determine the concentration with reference to a 5 mg. per 100 ml. standard carried through
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the procedure at the same time, or by referring to a
calibration curve prepared as described below. The colour
is linear, up to 14 mg. per 100 ml. In the case of urine
multiply by 10 to correct for the dilution.

For calibration, measure 2.0, 4.0, 6.0, 8.0, 10.0, 12.0,
14.0, 16.0, and 18.0 ml. of uric acid standard in
100-ml. volumetric flasks. Dilute to the 100-ml. mark in
each case with water. These standards are equivalent to
protein-free filtrates (1 : 10) of biological fluids containing
2.4, 6.8, 10, 12, 14, 16, and 18 mg. % uric acid. Use 2.0
ml. of each standard in place of the supernatant fluid in
the routine procedure above. This should be checked at
regular intervals. A typical calibration curve is shown
(Fig. 1). It can be seen that the reaction follows Beer's law
to about 14 mg. % uric acid. In the Caraway (1955)
procedure the colour is linear to only 8 to 10 mg. %,
depending on the instrument used.

All the figures quoted in this paper were obtained with
a Colemanootnote{Coleman Instruments, Inc., Maywood, Ill., U.S.A.} Junior spectrophotometer, model 6D using
19-mm. round cuvettes. To make readings on as little as
4 ml. of solution in 19-mm. round cuvettes, elevate them
by placing a slice of solid rubber stopper at the bottom
of the cuvette adaptor.

EXPERIMENTAL

OPTIMAL CONCENTRATIONS OF REAGENTS. The sodium
carbonate solution was varied in strength between 2 %
(w/v) and 14 % (w/v). These concentrations were tested
in an actual serum assay. Similarly, the phosphotungstic
acid concentration of that reagent was varied from 0.8
of the present concentration to 10 times the present
concentration while keeping the sodium carbonate
constant at 10 % (w/v). The results (Tables I and II)
indicate that both reagents are well within the optimal
limits.

TABLE I

<table>
<thead>
<tr>
<th>Sodium Carbonate (%) (w/v)</th>
<th>Final Colour (Absorbance)</th>
</tr>
</thead>
<tbody>
<tr>
<td>2</td>
<td>0.167</td>
</tr>
<tr>
<td>4</td>
<td>0.191</td>
</tr>
<tr>
<td>5</td>
<td>0.199</td>
</tr>
<tr>
<td>6</td>
<td>0.209</td>
</tr>
<tr>
<td>8</td>
<td>0.215</td>
</tr>
<tr>
<td>9</td>
<td>0.222</td>
</tr>
<tr>
<td>10</td>
<td>0.222</td>
</tr>
<tr>
<td>12</td>
<td>0.222</td>
</tr>
<tr>
<td>14</td>
<td>0.222</td>
</tr>
</tbody>
</table>

TABLE II

<table>
<thead>
<tr>
<th>Phosphotungstic Acid Reagent (Times Present Strength)</th>
<th>Final Colour (Absorbance)</th>
</tr>
</thead>
<tbody>
<tr>
<td>10</td>
<td>0.166</td>
</tr>
<tr>
<td>8</td>
<td>0.133</td>
</tr>
<tr>
<td>6</td>
<td>0.175</td>
</tr>
<tr>
<td>5</td>
<td>0.199</td>
</tr>
<tr>
<td>4</td>
<td>0.201</td>
</tr>
<tr>
<td>3</td>
<td>0.209</td>
</tr>
<tr>
<td>2</td>
<td>0.222</td>
</tr>
<tr>
<td>3</td>
<td>0.222</td>
</tr>
<tr>
<td>0.9</td>
<td>0.222</td>
</tr>
</tbody>
</table>

EFFECT OF PIGMENTS AND OTHER CONSTITUENTS IN SERUM
Haemolysis must be avoided. Interfering materials are
found largely in the cells and the red cells contain much
less uric acid than serum. Bilirubin, up to 5.2 mg. %,
appears to exert no effect on the determination, nor does
lipaemia.

Sodium oxalate, ammonium heparin, sodium citrate,
and the disodium salt of ethylenediamine tetracetic acid
are all acceptable anticoagulants.

We found that salicylates spuriously raise serum values
as much as 60%. Many diuretic and antihypertensive
agents, such as chlorothiazide, hydrochlorothiazide,
bendroflumethiazide, and pyrazinamide, also have a
tendency to elevate serum levels to as high as 12 mg. %.
It is best to have the patient taken off all drug therapy for
at least one week before drawing blood for this determin-
ation.

RESULTS

RECOVERY OF ADDED URIC ACID. If aqueous standards
are admixed with serum before deproteinization,
apparently a metastable solution of uric acid forms,
with recoveries averaging only about 37 % at levels
of 2.5 to 5.0 mg. % added uric acid (calculated final
levels of 9.1 to 11.8 mg. %). However, when we admixed
diluted serum with a commercial control
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serum, the recoveries ranged from 97·5 to 102% (mean 99·0; s ± 1·1). Thus, uric acid appears to be adequately recovered from a physiological milieu. By a different approach, Caraway (1963) arrived at the same conclusion.

**COMPARISON OF METHODS** The present method was compared with that of Henry et al. (1957) and with the Caraway (1955) method. The results in Table III indicate excellent correlation between the three methods.

**TABLE III**

<table>
<thead>
<tr>
<th>Serum No.</th>
<th>Serum Uric Acid (mg./100 ml.)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Present</td>
</tr>
<tr>
<td>1</td>
<td>2·1</td>
</tr>
<tr>
<td>2</td>
<td>6·6</td>
</tr>
<tr>
<td>3</td>
<td>11·2</td>
</tr>
<tr>
<td>4</td>
<td>12·7</td>
</tr>
<tr>
<td>5</td>
<td>13·9</td>
</tr>
<tr>
<td>Mean</td>
<td>9·3</td>
</tr>
</tbody>
</table>

**REPRODUCIBILITY** Twenty-five sera were analysed in duplicate. The standard deviation, calculated by the formula of Henry and Dryer (1963), $s = \sqrt{\frac{\sum (d^2)}{N}}$ (d = difference between duplicates; N = total number of determinations), was ± 0·07 for the present method. A similar statistical treatment of 20 cerebrospinal fluid samples (with normal levels of uric acid) and 20 urine samples resulted in standard deviations of ± 0·10 and ± 0·08, respectively.

**STABILITY OF URIC ACID IN BODY FLUIDS** Uric acid is stable in serum and cerebrospinal fluid for at least 24 hours at room temperature (approx. 25°C), for at least one month in the refrigerator (approx. 5°C.), and at least four months in the freezer (approx. —15°C.).

The stability of uric acid in unpreserved urine is variable. By saturating with toluene, uric acid may be preserved in urine for several days in the refrigerator (approx. 5°C.).

**NORMAL VALUES** Ninety-five per cent limits were calculated using K factors as described by Henry and Dryer (1963).

The 95% limits for serum uric acid were 3·1 to 7·0 mg. % for males and 2·0 to 5·9 mg. % for females. In urine, the 95% limits were 0·1 to 2·0 g. per 24 hours but no sex difference was found.

The 95% ‘normal’ limits for cerebrospinal fluid uric acid were 0·2 to 0·3 mg. %. No sex difference was determined.

**DISCUSSION**

Lithium sulphate was included in the phosphotungstic acid reagent merely on the suggestion of Henry et al. (1957). We found its use unnecessary, in limited testing, since we experienced no turbidity in the final coloured mixture regardless of its use or deletion. We retained it, however, as prophylaxis against some yet unencountered serum interference which might produce problems with clarity in the final colour.

**REFERENCES**


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