Automated method for the estimation of serum protein-bound iodine following alkaline incineration

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SYNOPSIS. A method is described for the estimation of serum protein-bound iodine using alkaline incineration and an automated technique for the estimation of iodine in the ash. Pretreatment of the serum with an anion exchange resin avoids the need for precipitation and washing of the protein. The method is accurate, reproducible, and simple to perform.

A detailed investigation of the alkaline ashing method for the determination of serum protein-bound iodine was described by Foss, Hankes, and Van Slyke (1960). Their procedure included precipitation and washing of the serum protein, incineration of the protein with potassium hydroxide, and the colorimetric estimation of the inorganic iodine in the ash by its quantitative effect on the reduction of ceric sulphate with arsenious acid.

A method for the automation of the final colorimetric analysis was produced by the Technicon Instruments Company and papers have been published describing various modifications of this procedure by Widdowson and Northam (1963) and Benotti and Benotti (1963). However, these methods are only suitable for the analysis of samples after chloric acid digestion. Stevens and Levandoski (1963) described an AutoAnalyzer technique for the estimation of iodine in alkaline ash which involved the addition of a known excess of iodine to each sample. The method was considered to be too complicated, insensitive, and slow for the number of estimations normally carried out in a large laboratory.

This communication describes a simple method for the estimation of serum protein-bound iodine using the alkaline incineration technique of Foss et al. (1960) and an AutoAnalyzer procedure for the estimation of iodine in the ash which is rapid and reproducible. As an anion exchange resin is used to remove any inorganic iodine from the serum, precipitation and washing of the serum protein was found to be unnecessary.

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8 STOCK IODIDE SOLUTION A (20 mg. IODINE PER 100 ML.) Dissolve 262 mg. of potassium iodide in water and dilute to 1 litre.

9 STOCK IODIDE SOLUTION B (200 µG. IODINE PER 100 ML.) Dilute 10 ml. of iodide solution A to 1 litre with water.

10 STOCK STANDARD (EQUIVALENT TO 100 µG. IODINE PER 100 ML.) Dilute 10 ml. of iodide solution B with about 40 ml. of water, add 20 ml. of 2N potassium hydroxide solution, and 20 ml. of acid reagent. Make up to 100 ml. with water.

11 WORKING STANDARDS Dilute the stock standard (equivalent to 100 µg. iodine per 100 ml.) with blank reagent to give a series of standards equivalent to 5, 10, 15, and 20 µg. iodine per 100 ml. The standards are stable for at least four weeks if stored in amber bottles.

Use double-deionized water for the preparation of all solutions and standards.

PROCEDURE

Mix the serum thoroughly with a little of the anion exchange resin (approximately 0-2 g.) to remove any inorganic iodine. Centrifuge the serum and transfer 1 ml. to a Pyrex incineration tube, add 1 ml. of 2N potassium hydroxide, mix thoroughly and place the tubes in an oven at 115°C. overnight (at least 18 hours). When the tubes are completely dry transfer them to a cool muffle furnace and close the door. Allow the temperature to rise to 620°C. When the oven has reached the required temperature open the door for about 30 seconds; repeat this procedure at 20-minute intervals for one hour. Remove the tubes and allow to cool to room temperature. Gently add 5 ml. of acid diluent, mix thoroughly, and centrifuge at 2,000 r.p.m. for 10 minutes. Without delay decant a portion of each supernatant into an AutoAnalyzer sample cup and estimate the iodine in the solution using the AutoAnalyzer system shown in Figure 1. To obtain the correct base line equivalent to 0 µg. of iodine per 100 ml., pump the blank reagent through the sample line while the other reagents are passing through their respective flow tubes. The rate of determination is 60 samples per hour.

RESULTS AND DISCUSSION

Ten samples of a normal and abnormal Hyland control serum were analysed by the above procedure. The normal control serum (5-0 µg. iodine per 100 ml.) gave a mean result of 4-8 µg. per 100 ml. and the abnormal control serum (10-6 µg. iodine per 100 ml.) gave a mean result of 10-8 µg. per 100 ml. The standard deviations for the normal and abnormal controls were 0-12 and 0-24 respectively. The AutoAnalyzer record of this investigation is shown in Figure 2. Recovery experiments were carried out by adding a solution of sodium thyroxine to pooled serum corresponding to an increase of 10 µg. of iodine per 100 ml. The average recovery from 10 separate estimations was 95%. Resin treatment was found to be effective in removing any inorganic iodide likely to be present in sera. Using the recommended method the anion exchange resin absorbed 0-75 µg. of iodine per ml. of serum. A pooled serum from blood donors was analysed using both resin treatment and protein precipitation methods. The
mean result of 30 estimations using the resin technique was 4·4 µg. per 100 ml. (standard deviation 0·08) and 4·6 µg. per 100 ml. (standard deviation 0·15) when iodine is removed by precipitation and washing.

The possibility of iodine loss during the addition of the acid reagent to the alkaline ash was investigated. Contrary to the findings of Foss et al. (1960) the amount of iodine lost during the vigorous evolution of carbon dioxide was negligible.

After the acid reagent is added to the ash, the supernatant is immediately transferred to the sample cup and analysed. If the acid solution remains in the incineration tubes the iodide concentration is slowly reduced. The reason for this effect is unknown but is possibly due to adsorption of iodine on the etched surface of the tubes.

In the AutoAnalyzer technique the carry-over contamination from the 20 µg. per 100 ml. standard was less than 0·2 µg. per 100 ml. It was therefore unnecessary to alternate the test samples with blank solutions.

The semi-automated procedure allows at least 180 specimens, divided into five batches, to be estimated by one technician in five half-days. If a second rack for incineration tubes is used this volume of work can be doubled without any appreciable increase in total working time.

REFERENCES

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