Iodine contamination of the serum protein-bound iodine: Incidence and clinical significance

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SYNOPSIS Iodine contamination as defined by the combination of a raised (PBI-T₄I) difference and low ¹³¹I neck uptake was found in 38 (17·5%) of 217 euthyroid patients. Of these, 17 had PBI levels of greater than 20 µg/dl but in the remainder levels were clinically feasible. In only 21 was there a history of exposure to iodine. Two of 12 hypothyroid patients had PBI levels well within normal limits. False elevation of the PBI is thus shown to be common. It is neither always obvious nor can it be easily avoided. The PBI is not an acceptable alternative to T₄ estimation by other methods.

Thyroid disease is probably the commonest endocrine abnormality encountered in clinical practice and enters into the differential diagnosis of a number of conditions, particularly obesity and anxiety states. When considered, it must be confirmed or excluded by appropriate laboratory investigations (Bayliss and Hall, 1973). The simplest method is the estimation of serum thyroxine (T₄), preferably with an assessment of thyroxine-binding proteins by a resin-uptake test. The product of serum thyroxine and resin uptake, the calculated free thyroxine index (FTI), is accepted as the most reliable single indicator of thyroid secretory function (Bayliss and Hall, 1973).

For the calculation of the FTI either the protein-bound iodine (PBI) or the serum T₄, usually determined by a method not measuring iodine, may be used. Recent reviews on the subject (Flynn and Hobbs, 1971; Bayliss and Hall, 1973), while warning against falsely elevated PBI values, nevertheless list this test as an alternative to serum T₄ measurement. The PBI, which is cheaper and more easily automated than the T₄ estimation, remains a popular test of thyroid function despite the well known risk of spuriously high results due to iodine contamination (Acland, 1971). This may be due to the belief that contamination is easily recognized by the unrealistically high values obtained, or that it can be avoided by excluding patients who have recently ingested iodine in any form or who have undergone diagnostic procedures involving iodine-containing radioopaque media.

The present study was undertaken to test the acceptability of the PBI as an alternative to T₄ estimation. There were three main aims: (1) to assess the incidence of iodine contamination in patients referred for thyroid function studies; (2) to see whether such interference is easily recognized or could be prevented; (3) to assess, within the limits of the study, the diagnostic implications of such interference with special reference to the detection of hypothyroidism.

Patients and Methods

Most methods of PBI estimation employ steps to remove or reduce exogenous iodine contamination, with varying success. It was felt therefore that the best index of a falsely high PBI result would be the difference between the thyroxine iodine measured in the PBI and that calculated from thyroxine measured by a competitive protein-binding technique. This difference, representing non-thyroxine iodine, will be referred to as dl. Exogenous iodine may also cause a falsely low ¹³¹I neck uptake by enlarging the body iodide pool. It seems therefore reasonable to consider that in euthyroid patients the combination of a low neck uptake and a high dl is evidence of iodine contamination. Such a group was sought among patients referred to us for thyroid function studies.

Patients
All patients referred were carefully questioned about possible iodine intake during the preceding three months, in the form of cough mixtures, throat...
lozenges, 'traveller's diarrhoea' preparations, eg, Enterovioform (Ciba), and iodine itself. Information on all previous radiological examinations and where possible a detailed history of all drugs was obtained. Patients were also asked whether they habitually used municipal water or well water.

Venous blood was drawn with precautions to avoid exogenous iodine contamination. Disposable syringes and needles were used and blood for PBI estimation was put into clean disposable containers.

**METHODS**

Serum T₄ was measured by a competitive protein-binding technique (Tetrasorb-125, Abbott Laboratories) and the T₃-resin uptake with the Triosorb-125 kit (Abbott Laboratories). The PBI was estimated by a standard AutoAnalyzer (Technicon) method after preliminary removal of inorganic iodide with IRA-400 Amberlite; a rapid sampling screening run detected all grossly elevated values.

The ¹³¹I neck uptake was measured 24 hours after an oral dose of ¹³¹I obtained from the Radiochemical Centre, Amersham, using a Nuclear Chicago scintillation counter with collimator according to the specifications of the International Atomic Energy Commission.

**CALCULATIONS**

Thyroxine iodine (T₄I) was calculated as T₄ (µg/dl) × 0.635. The FTI was calculated as the product of the serum T₄ (or PBI) and T₃-resin uptake and is referred to as FTI(T₄) or FTI(PBI) respectively; as one calculation is based on iodine and the other on thyroxine different numerical values are obtained. The difference (PBI-T₄I) was calculated for each sample and is referred to as dI.

**Results**

Two hundred and eighty-seven patients were seen. Of these 14 were judged hyperthyroid and 12 hypothyroid on the basis of the test results including, where indicated, stimulation or suppression tests. Twenty-four patients were being treated for thyroid disease and in a further 20 patients the final data were incomplete.

Two hundred and seventeen patients with normal FTI(T₄) values remained for evaluation and, unless otherwise indicated, all results refer to this group.

**THE INCIDENCE OF IODINE CONTAMINATION**

The distribution of the dI is shown in figure 1. The scatter and mean of the neck uptakes of different groups of dI are shown in fig 2 and the results summarized in table I. The group of two patients with dI of less than −2.0 is too small for analysis.

The mean neck uptakes of the subgroups of group A (dI -1.9 to +2.0 µg/dl) did not differ significantly. The mean neck uptakes of the two subgroups of group B, both separately and together, were significantly lower than that of group A. The mean FTI(T₄) did not differ significantly between groups A and B whereas the mean FTI(PBI) was significantly higher in group B. The T₄ was within the normal range in all the patients of group B, except two who had mildly elevated values of 13.3 and 13.4 µg/dl (normal 5.6–12.6).

It was considered therefore that the patients of
**Iodine contamination of the serum protein-bound iodine: incidence and clinical significance**

<table>
<thead>
<tr>
<th>dl (µg/dl)</th>
<th>Group A (dl &lt; 2.0)</th>
<th>Group B (dl &gt; 2.0)</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>-1.9 to 0.0</td>
<td>0.1 to 1.1</td>
<td>1.1</td>
</tr>
<tr>
<td></td>
<td>-1.0 to 0.0</td>
<td>1.0 to 2.0</td>
<td>2.1</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>24-hr neck uptake (%)</th>
<th>Mean and (SD)</th>
<th>FTI(T₄) mean ± SD</th>
<th>FTI(PBI) mean ± SD</th>
<th>Number of patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>29.8 to 30.9</td>
<td>(11.0) to (10.5)</td>
<td>2.53 (±0.54)</td>
<td>1.74 (±0.22)</td>
<td>179</td>
</tr>
<tr>
<td>30.9 to 30.2</td>
<td>(10.5) to (8.6)</td>
<td>2.46 (±0.52)</td>
<td>2.69 (±0.74)</td>
<td>38</td>
</tr>
</tbody>
</table>

Table I Differences in indices of thyroid function in euthyroid patients with dl (PBI-T₄I) less and greater than 2.0 µg/dl

1The neck uptakes of both subgroups of B differ significantly from the mean neck uptake of group A. The mean values of FTI(T₄) and FTI(PBI) differ numerically as the former is calculated on thyroxine and the latter on iodine.

**group B**, i.e., those with a dl of greater than 2.0 µg/dl showed evidence of iodine contamination. This group contained 38 patients (17.5%). In 16 the PBI was greater than 20 µg/dl while in the remaining 22 it was less than 16 µg/dl. Of these, 13 patients had values of less than 10 µg/dl.

**SOURCE OF CONTAMINATION**

The history of iodine exposure is shown in table II. The incidence of exposure to radioopaque media and iodine-containing drugs was significantly greater in group B than group A, whereas that of exposure to other drugs or well water did not differ significantly. The time elapsed since exposure to radioopaque media and the type of investigation are shown in table III. All four patients of group B who had had intravenous pyelograms had also had a cholecystogram performed (one in the previous six months and three more than five years earlier). The evidence of iodine contamination in patients who had had cholecystograms more than five years before, while those in the intervening period appear to be unaffected, is interesting but no explanation could be found. In this small number of cases it may be fortuitous. Three had also used iodine-containing drugs.

**Table II Exposure of patients to possible sources of iodine contamination**

<table>
<thead>
<tr>
<th>I-containing Drugs</th>
<th>Radioopaque Media</th>
<th>Other Drugs</th>
<th>Well Water</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group A</td>
<td>12.3</td>
<td>16.2</td>
<td>46.8</td>
</tr>
<tr>
<td>Group B</td>
<td>31.6</td>
<td>39.5</td>
<td>44.7</td>
</tr>
<tr>
<td>Significance</td>
<td>p &lt; 0.05</td>
<td>p &lt; 0.05</td>
<td>NS</td>
</tr>
</tbody>
</table>

**Discussion**

To establish whether our group A patients are similar to those found elsewhere the findings may be compared with those of other authors. The mean dl of this group was 0.29 (±0.9) µg/dl. Farran,
Haiste, and Hoffenberg (1971) found a mean dl of 0.35 μg/dl in 217 euthyroid patients and their diagram indicates a scatter of results similar to ours. Keeling and Williams (1972) found a mean 24-hour 131I neck uptake of 31.9% in 67 euthyroid patients (the corresponding figure in our group A is 30.2%). These findings indicate that our group of euthyroid patients is comparable regardless of iodine contamination. Farran et al drew attention to an increasing dl when levels of both T4 and PBI are elevated, regardless of cause, which they ascribed to a circulating non-thyroxine iodinated compound. The elevated dl values in the present series cannot be accounted for on the same basis as only two patients had T4 levels slightly above the normal range. The combination of elevated dl and low 131I neck uptake in patients who are not taking thyroid supplements can only be satisfactorily explained on the basis of iodine excess. This being so, the results indicate that almost one in five PBI estimations is falsely elevated.

The significance of this finding is emphasized by the small series of hypothyroid patients, two of whom (16.7%) had PBI and FTI(PBI) values well within the normal range. It is realized that without serum TSH estimation or TSH stimulation tests in all clinically suspected patients, early hypothyroidism could be missed. This reservation, however, applies to both T4 and PBI estimations.

Similarly, there were 16 euthyroid patients with PBI values ranging from 8.4 to 15.6 μg/dl, levels which could cause diagnostic confusion or misdiagnosis. The 16 patients with PBI values greater than 20 μg/dl would almost certainly be recognized as contaminated.

There is no sure way of predicting which patients will have falsely raised PBI values. Certainly all who have had cholecystograms or angiograms in the previous six months should be suspected whereas, in this small series, intravenous pyelography seems less troublesome. Other procedures using iodine-containing radioopaque media such as myelography are known to affect thyroid function tests for many years (Acland, 1971). Excluding all patients with any history of such procedures or of iodine intake, would limit the problem but in 45% of those affected in the present series no such history was obtained on direct questioning. In many contamination was massive.

As serum thyroxine estimation by methods not involving measurement of iodine are widely available there seems no good reason to continue using a test with such a high incidence of false results. Protein-bound iodine estimation should not be considered as an acceptable alternative to thyroxine estimation either directly or for use in calculating the FTI.

References

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