Determination of Iodoamino Acids and Inorganic Iodide by Thin-Layer Chromatography

CHUNG WU AND RONALD C. LING

Department of Internal Medicine, The University of Michigan Medical School, Ann Arbor, Michigan 48104

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Many methods for the determination of iodinated compounds in thyroid extracts by paper and thin-layer chromatography have been reported. Some of them do not separate inorganic iodide (1-3), diiodothyronine (4-8), or mono- and di-iodotyrosines (6, 9). Others use more than one solvent system (4, 7, 10-12), usually one system for iodothyrosines and another for iodothyronines. Still others show poor separation of the compounds (2, 7), or are not supported by adequate recovery data (1-3, 5-9, 11, 12). However, good separation (3, 5, 10, 11) and recovery (4, 10) have been achieved in several laboratories. In our study of the synthesis of iodinated compounds in thyroid tumors, we have developed a solvent system for the separation and quantitative determination of thyroidal iodine-containing compounds including inorganic iodide by thin-layer chromatography. This paper describes our method and its application to $^{131}$I experiments in vivo.

MATERIALS AND METHODS

Reagents

3-Monoiodo-L-tyrosine (MITyr) (Mann).
3,5-Diiodo-L-tyrosine (DITyr), 3,5-diiodo-L-thyronine (DIThy), 3,3',5-triiodo-L-thyronine (TIThy), and L-thyroxine (Sigma).

tert-Butyl alcohol (Eastman).
Methyl ethyl ketone, concentrated ammonium hydroxide, 28.9%, and potassium iodide (Baker).

tert-Amyl alcohol (Baker or Fisher).
Palladium chloride.

Ninhydrin (Nutritional Biochemicals).

Eastman chromatogram sheets #6060 (Silica gel with fluorescent indicator).
**Procedure**

_A. Thin-layer chromatography._ Eastman chromatogram sheets were used to make ascending runs. Iodoamino acids and KI were spotted at the origin, 2 cm from the lower edge of the sheet. The time of spotting was kept to a minimum to avoid degrading the amino acids. The standard iodoamino acid and KI solutions were prepared in 2N NH₄OH to contain 1–2 mg/ml. In spotting we used 0.5–1.5 μg MITyr, DITyr, and KI, and 1.5–2.5 μg DIThy, TIThy, and thyroxine.

The 20 × 20 cm chromatogram sheets were usually cut into 10 × 20 cm strips to get an even solvent front. They were held at the top by two stainless-steel clips which were in turn strung through two parallel stainless-steel wires 4 cm apart. Glass tubing 1 cm in length was inserted as a spacer between two successive clips. With stainless-steel supporting frame 20 × 16 × 22 cm high, a total of four parallel wires and 20 clips were mounted. A maximum of 10 sheets of 10 x 20 cm each can be hung on the frame.

A solvent system consisting of 295 ml tert-butyl alcohol, 160 ml tert-amyl alcohol, 75 ml 28.9% NH₄OH, 35 ml methyl ethyl ketone, and 85 ml distilled water was prepared and mixed thoroughly. It was allowed to stand for a few minutes, and the entire content was then poured into a Pyrex rectangular jar 24 × 18 × 32 cm high with a flat ground edge. The supporting frame with chromatogram sheets was lowered into the jar. The height of the sheets should be so adjusted beforehand that about 0.7 cm was submerged in the solvent. A piece of filter paper moistened with about 2 ml 28.9% NH₄OH was laid flat on the top of the rack. The jar was covered and sealed with a piece of ground-glass plate. It was kept at 26°.

The time it took for a complete run varied somewhat with the chromatogram sheets. It took less time and gave better separations with the sheets purchased in 1968 than those bought in 1969. With the “newer” sheets, however, it took 10 hr for the solvent front to travel 15 cm, at which point the run was discontinued. After drying the sheets in air, we sprayed them with 0.3% ninhydrin in n-butyl alcohol/glacial acetic acid (100:3) or viewed them under the ultraviolet light to mark the spots of iodotyrosines and iodothyronines. The iodide spot was sprayed with 0.1% PdCl₂ to give a brown color. The spots were cut and eluted in 4 ml water. The eluates were centrifuged, if necessary. The compounds were determined quantitatively in recovery experiments at their respective absorption maxima as follows: MITyr, 305 μm; DITyr, 223 and 310 (minor peak) μm; DIThy, 224 μm; TIThy, 224 and 320 (minor peak) μm; thyroxine, 226 and 325 (minor peak) μm; and KI, 226 μm.
B. Labeling experiments with $^{131}I$. About 300 $\mu$Ci $^{131}I$ was injected subcutaneously into a rat. At the end of 24 hr, the rat was killed by means of an overdose of pentobarbital. The thyroid glands and a piece of neck muscle serving as a control were dissected. A homogenate of thyroid tissue in ice-cold distilled water was prepared to contain 20 mg/ml. A sufficient quantity of 15% trichloroacetic acid was added so that the final concentration was 7%. After centrifugation at 27,000g for 15 min, the supernatant fluid was collected. The precipitate was washed with 0.5 ml cold distilled water and centrifuged again. This process of washing was repeated once. The two washings and the original fluid were combined and brought to dryness under reduced pressures. The residue was dissolved in sufficient 10% isopropyl alcohol in 2 N NII,OH to give a concentration equivalent to 30 mg thyroid tissue per milliliter.

Aliquots of the tissue extract, usually 1–3 $\mu$l, were applied on the chromatogram sheet. Unlabeled iodoamino acids and KI were added to the origin as carriers. The chromatograms were developed as described in the preceding section. The radioactive spots corresponding to the carriers and the area between two such spots were cut and placed in disposable test tubes (Packard Instruments Co.) for radioactivity counting. Direct gamma counting was done with a Packard autogamma counter.

RESULTS

Figure 1 shows the separation of MITyr, DITyr, DIThy, TIThy, thyroxine, and iodide on silica gel chromatogram. Their $R_f$ values with the “newer” sheets are as follows: 0.19 (DITyr), 0.23 (MITyr), 0.48 (thyroxine), 0.53 (iodide), 0.59 (TIThy), and 0.66 (DIThy). When tyrosine and thyronine are present, tyrosine will appear between MITyr and thyroxine, and thyronine, between iodide and TIThy on the chromatogram. The “older” chromatogram sheets yielded better separation of the compounds in one-half as much time, presumably because the particles used to prepare the sheets were not so fine.

To test the suitability of the method for quantitative determinations, we determined the recovery of each compound after thin-layer chromatography. The recovery values were similar whether the compounds were applied individually or in combination, and whether the compounds were added to a thyroid gland extract or not. Table 1 shows the results. For the two iodotyrosines, thyroxine, and inorganic iodide, the recoveries were excellent. Slightly lower recoveries, however, were obtained for the two iodothyronines.

We have used this method extensively to study incorporation of $^{131}I$
Fig. 1. Diagram of chromatograms of iodoamino acids and inorganic iodide on (A) "older" sheet, 5 hr, and (B) "newer" sheet, 10 hr. The numbers identify: DIThy (1), TTThy (2), iodide (3), thyroxine (4), MITyr (5), and DITyr (6).
TABLE 1

Recovery of Iodoamino Acids and Related Compounds after Chromatography

After chromatography, an amount of the compound equal to that previously applied at the origin was applied to an area adjacent to the spot of that compound on the chromatogram. Both spots were then cut and the compound extracted in 4 ml water. The amount in each extract from the chromatographed spot and from the newly applied spot was determined from a standard curve, which is linear at least through 30 μg/ml for each of these compounds. From these values percent recovery was calculated. Earlier, we had determined that the extraction was quantitative.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Wavelength used, μm</th>
<th>Amt. before chromatog., μg</th>
<th>Amt. after chromatog., μg</th>
<th>% recovery</th>
</tr>
</thead>
<tbody>
<tr>
<td>MITyr</td>
<td>305</td>
<td>60.0</td>
<td>58.8</td>
<td>98.0</td>
</tr>
<tr>
<td>DITyr</td>
<td>223</td>
<td>60.0</td>
<td>59.0</td>
<td>98.3</td>
</tr>
<tr>
<td>DITHy</td>
<td>224</td>
<td>10.0</td>
<td>9.4</td>
<td>94.0</td>
</tr>
<tr>
<td>TIThy</td>
<td>224</td>
<td>10.0</td>
<td>9.2</td>
<td>92.0</td>
</tr>
<tr>
<td>Thyroxine</td>
<td>226</td>
<td>10.0</td>
<td>10.0</td>
<td>100.0</td>
</tr>
<tr>
<td>KI</td>
<td>226</td>
<td>12.0</td>
<td>11.9</td>
<td>99.2</td>
</tr>
<tr>
<td>Tyrosine</td>
<td>281</td>
<td>20.0</td>
<td>18.8</td>
<td>94.0</td>
</tr>
<tr>
<td>Thyonine</td>
<td>230</td>
<td>20.0</td>
<td>18.6</td>
<td>93.0</td>
</tr>
</tbody>
</table>

into iodotyrosines and iodothyronines by thyroid gland and thyroid tumors of the rat under different conditions. Table 2 illustrates the information that can be obtained from this kind of experiment. The experiment was carried out as described under “Procedure.” After each compound has been counted, the data in counts per minute per milligram tissue were converted to percent of total counts in the thyroid extract, of which an aliquot was also used in counting. Table 2 lists the values obtained from 6 tumor-bearing rats together with the means and standard

TABLE 2

Incorporation of 131I into Iodoamino Acids by Thyroid Gland of Rat Bearing a Transplantable Thyroid Tumor

The values given are expressed as percent of total radioactivity present in the thyroid extract.

<table>
<thead>
<tr>
<th>Compounds</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>6</th>
<th>Mean</th>
<th>S.D.</th>
</tr>
</thead>
<tbody>
<tr>
<td>DIThy</td>
<td>1.0</td>
<td>1.2</td>
<td>1.4</td>
<td>1.4</td>
<td>1.2</td>
<td>1.1</td>
<td>1.2</td>
<td>0.2</td>
</tr>
<tr>
<td>TIThy</td>
<td>13.1</td>
<td>10.8</td>
<td>8.6</td>
<td>6.2</td>
<td>10.7</td>
<td>12.1</td>
<td>10.2</td>
<td>2.5</td>
</tr>
<tr>
<td>Iodide</td>
<td>30.5</td>
<td>26.7</td>
<td>31.4</td>
<td>33.0</td>
<td>32.5</td>
<td>27.4</td>
<td>30.2</td>
<td>2.6</td>
</tr>
<tr>
<td>Thyroxine</td>
<td>4.7</td>
<td>4.6</td>
<td>4.6</td>
<td>4.0</td>
<td>4.0</td>
<td>4.1</td>
<td>4.3</td>
<td>0.3</td>
</tr>
<tr>
<td>MITyr</td>
<td>5.9</td>
<td>4.7</td>
<td>4.8</td>
<td>5.5</td>
<td>5.8</td>
<td>5.4</td>
<td>5.4</td>
<td>0.5</td>
</tr>
<tr>
<td>DITyr</td>
<td>12.4</td>
<td>16.6</td>
<td>15.7</td>
<td>14.7</td>
<td>12.2</td>
<td>12.7</td>
<td>14.0</td>
<td>1.9</td>
</tr>
<tr>
<td>Total</td>
<td>67.6</td>
<td>64.6</td>
<td>66.5</td>
<td>64.8</td>
<td>66.4</td>
<td>62.8</td>
<td>65.4</td>
<td></td>
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</tbody>
</table>
deviations. As we would have expected, about one-third of the total radioactivity in the thyroid extract was recovered as inorganic iodide. Some 10 to 14% each of $^{131}$I in the extract appeared as TIThy and DITyr. Approximately 5% each was incorporated into MITyr and thyroxine. DIThy had the lowest incorporation of about 1%.

The sum of these percentages from the 6 compounds is about 65%. This leaves 35% of the total radioactivity in the extract not accounted for by the known compounds. Most of the radioactivity unaccounted for appeared at the origin and in the area between the origin and DITyr, though low levels of radioactivity were detected in areas in between two successive known spots. This portion of the total radioactivity could vary with the conditions of the experiment. Two factors may affect the outcome. First, the recovery of DIThy and TIThy was less than quantitative (Table 1). Second, the concentration of trichloroacetic acid used to prepare the extract may not precipitate certain $^{131}$I-labeled protein-like substances. This seems probable from the presence of radioactivity at the origin and in the area immediately above it.

**DISCUSSION**

The solvent system used in the present method contains several components. The proportions of these components have been so adjusted that the system gives a satisfactory separation of all the compounds. On the other hand, if only the iodotyrosines or the iodothyronines are to be studied, the system can be modified to give an even better separation of the group of compounds under consideration. Thus, increasing the proportion of tert-amyl alcohol increases the $R_f$ value for iodide, shortens the running time, and is good for separation of DIThy from TIThy. Ammonium hydroxide slows down the rate of movement of the iodide spot, thus improving the separation of TIThy from iodide. Increasing the proportion of methyl ethyl ketone also shortens the running time and makes good separation of MITyr from DITyr. The ketone, however, speeds the movement of iodide and interferes with separation of TIThy from DIThy and of TIThy from iodide. Increasing the proportion of tert-butyl alcohol or of water lengthens the running time but results in good separation of the iodotyrosines. In addition, water slows down the movement of iodide and facilitates TIThy separation. In this respect, its effect is like that of NH$_2$OH.

**SUMMARY**

A method has been presented for the separation of monoiodotyrosine diiodotyrosine, diiodothyronine, triiodothyronine, thyroxine, and inorganic iodide from one another by thin-layer chromatography on silica
iodo amino acids and iodide by TLC

The solvent system consisted of tert-butyl alcohol, tert-amyl alcohol, 28.9% NH₄OH, methyl ethyl ketone, and water (295:160:75:35:85). Recovery of the iodo compounds from chromatography gave satisfactory results. The method has been applied to study incorporation of ¹³¹I into these compounds by rat thyroid gland. Possible alterations of the five components in the solvent system to achieve certain special purposes are discussed.

ACKNOWLEDGMENTS

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REFERENCES