

Our Experience Using T₃ RIA KIT (PEG Method)

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Measuring T₃ (triiodothyronine) in the blood is rapidly becoming a common and indispensable clinical procedure. We had the opportunity to use the T₃ RIA KIT developed by Dinabott RI laboratory. This kit uses polyethyren-glycol (PEG) to separate antibody combined with hormone labeled RI from isolated hormone labeled RI. We intend to report the foundermental and clinical findings that we obtained with this kit.

(Object and Methods)

Foundermental Examination—In accordance with standard measuring method, we examined the time and temperature of incubation, and the influence of light. We also tested diluted T₃ high concentration serum, its recovery and its ability to reappear.

Clinical Examination—We measured T₃ in serum taken from 58 patients who have various thyroid diseases, 9 pregnancies, 3 nephrosis, and 33 healthy persons by means of this method. We then also measured T₃ in all these cases by means of the usual T₃ RIA KIT (dextran-charcoal

method) and compared the results obtained from both.

(Results)

We obtained a good standard curve which was taken after 2 hours of incubation at 25°C, and made these our assay condition. The dilution curve of T₃ high concentration serum run parallel to the standard curve. When we added standard 0.25–4.00 ng/ml of T₃ to the pooled serum, the rate of recovery was 87–120%, 98.6% on the average. In healthy persons, the amount of T₃ in serum was 1.20–1.73 ng/ml, 1.44 ± 0.15 ng/ml on the average; in hyperthyroidism patients, 2.65–8.00 ng/ml; in hypothyroidism patients, 0.09–1.14 ng/ml; in pregnancies, 1.05–1.98 ng/ml; in patients with nephrosis, 1.30–1.85 ng/ml. The correlation coefficient of the T₃ of the dextran-charcoal method and T₃ of the PEG method was $\gamma = 0.927$. This means that these correlate very well.

This new method is simple and can be performed quickly. For these reasons, we think this method is very useful to measure T₃ in serum.

Radioimmunoassay for Measurement of Triiodothyronine and Thyroxine in Human Serum

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Our experience with radioimmunoassay (RIA) used for measuring serum triiodothyronine (T₃) and thyroxine (T₄) is reported. The method has good specificity for T₃, T₄ and its reproducibility is satisfactory. RIA T₃ values ranged from 0.80 to 2.14 ng/ml (1.30 ± 0.31) in normal adults. RIA T₄ values ranged from 4.7 to 14.7 $\mu\text{g}/100 \text{ ml}$ (9.4 ± 2.7) in 50 normals cases.

In all clinical states, serum T₃ values obtained by RIA afforded excellent agreement (correlation coefficient -0.91) with those obtained by triiodo-

thyronine binding globulin index-thyopac 3. Serum T₄ values obtained by RIA afforded excellent agreement (correlation coefficient 0.91) with those obtained by competitive protein binding assay-thyopac 4.

Hemolytic serum are light higher than normal serum between the two techniques.

Advantages of the method over commercially available RIA kits or CPBA and TBG-index are outlined.