Contact points:

Pervaz Mohammed BCPS Specialist services clinical lead Tel: 0121 507 5353 Email: Pervaz.Mohammed@nhs.net

Dudley Laboratory

Tel: 01384 244082 Opening times: week days: 9.00am to 17:30pm Weekends & Bank Holidays: closed

Mailing Address

Department of Clinical Biochemistry Russell's Hall Hospital Pensnett Road Dudley, West Midlands DY1 2HQ





TRAB Test

Trace Elements Laboratory Clinical Biochemistry





Provided by Sandwell and West Birmingham NHS Trust, The Dudley Group NHS Foundation Trust, The Royal Wolverhampton NHS Trust and Walsall Healthcare NHS Trust.



A Teaching Trust of The University of Birmingham

Incorporating City, Sandwell and Rowley Regis Hospitals

© Sandwell and West Birmingham Hospitals NHS Trust ML6209 Issue Date: March 2021 Review Date: March 2023

Sample requirements

- Minimum of 250µl of serum.
- Allow sample to clot adequately before centrifugation.
- Keep tubes stoppered at all times.
- Refrigerate sample at 2-8°C until transport.

Clinical Use

Autoantibodies to the thyroid stimulation hormone receptor (TRAb) are beneficial in diagnosis and management of hyperthyroidism in Graves' disease (autoimmune hyperthyroidism).

The thyroid stimulating hormone (TSH) receptor (serves as an antigen) is a twosubunit glycoprotein; the extracellular A subunit is recognised by thyroid stimulating antibodies, while those antibodies recognising the B subunit, located much nearer the cell surface, appear to function as blocking antibodies. Based on the mechanism of action TSH autoantibodes can be classified as stimulating blocking or neutral.

Although TSHR stimulating antibodies have a similar action to TSH they are not subjective to the negative feedback mechanism correlated with TSH, leading to continual prolonged activation of the TSHR. Resulting in elevated thyroid hormone level and thyrotoxicosis associated with Graves' disease. Indications for TRAb determination include:

- By way of exclusion, the presence of TSH receptor antibodies (TRAbs) in the serum sample differentiate between thyrotoxicosis with hyperthyroidism (for example, Graves' disease rather than toxic nodular goitre) and thyrotoxicosis without hyperthyroidism (for example, transient thyroiditis).
- 2. Absence of TRAbs.
- 3. Can be used as an important decision making for the monitoring, treatment and prediction of relapse of Graves' disease. During treatment with antithyroid drugs the level of TRAbs tends to fall may indicate remission thereby allowing the withdrawal of therapy.
- 4. TRAb are IgG-class antibodies that cross the placenta and can cause neonatal thyroid disease. The importance of measuring TRAbs in the last trimester of pregnancy aids in assessing the risk of thyroid disease in the neonate.

Reference ranges:

Interpretation of results	
<1.1 IU/L	Normal
1.11-1.75 IU/L	Intermediate
>1.75 IU/L	Raised

Method:

TRAb is measured in serum using Cobas e 801 employing a (competitive principle) immunoassay system. We offer electronic reporting of results by PDF and NPEX.

Turn round

We aim to analyse and report the results within 5 working days from receipt of sample.

References

- 1. Sinclair D. Analytical aspects of thyroid antibodies estimation. Autoimmunity. 2008;41(1):46-54. doi:10.1080/08916930701619466.
- Kamijo K. TSH-receptor antibodies determined by the first, second and third generation assays and thyroidstimulating antibody in pregnant patients with Graves' disease. Endocr J 2007;54(4):619-624.
- Schott M, Seißler J, Scherbaum WA. Diagnostic testing for autoimmune thyroid diseases. J Lab Med 2006;34(4):254-257.
- 4. NICE guideline [NG145] Published date: 20 November 2019 NICE clinical guidelines CG122.

UKAS Accredited

