

# Novocastra™ Liquid Mouse Monoclonal Antibody Thyroid Peroxidase

**Product Code: NCL-L-TPO**

<b>Intended Use</b>	FOR RESEARCH USE ONLY.
<b>Specificity</b>	Human thyroid peroxidase.
<b>Clone</b>	AC25
<b>Ig Class</b>	IgG1
<b>Antigen Used for Immunizations</b>	Prokaryotic recombinant protein containing 105 amino acids corresponding to a portion of the N-terminus of the human thyroid peroxidase molecule.
<b>Hybridoma Partner</b>	Mouse myeloma (p3-NS1-Ag4-1).
<b>Preparation</b>	Liquid tissue culture supernatant containing 15 mM sodium azide. Volume as indicated on vial label.
<b>Effective on Frozen Tissue</b>	Not evaluated.
<b>Effective on Paraffin Wax Embedded Tissue</b>	Yes (using heat induced epitope retrieval with Tris/EDTA based buffer, pH 9.0: see overleaf).
<b>Recommendations on Use</b>	Immunohistochemistry: Typical working dilution 1:80. Heat induced epitope retrieval with Tris/EDTA-based buffer, pH 9.0. 30 minutes primary antibody incubation at 25 °C. Polymer detection recommended. Western Blotting: Not recommended.
<b>Positive Controls</b>	Immunohistochemistry: Thyroid.
<b>Staining Pattern</b>	Cytoplasmic.
<b>Storage and Stability</b>	Store liquid antibody at 4 °C. Under these conditions, there is no significant loss in product performance up to the expiry date indicated on the vial label. Prepare working dilutions on the day of use.
<b>General Overview</b>	Thyroid peroxidase (TPO) is a membrane-bound protein that is involved in two important reactions in the biosynthesis of thyroid hormones; the iodination of tyrosine residues on thyroglobulin and the intramolecular coupling reaction of iodinated tyrosines, leading to the formation of 3-3'-5-triiodothyronine (T3) and thyroxine (T4). The thyroid peroxidase gene expression is under the regulation of thyroid stimulating hormone (TSH). In normal thyroid, expression of TPO described immunohistochemically is reported to produce a diffuse, fine granular cytoplasmic stain in all follicular cells.
<b>General References</b>	Czarnocka B, Pastuszko D, Janota-Bzowski M et al. Breast Journal Cancer 85(6): 875-880 (2001) Christensen L, Blichert-Toft M, Brandt M et al. Clinical Endocrinology 53: 161-169 (2000) Tanaka T, Umeki K, Yamamoto I et al. Journal of Pathology 179: 89-94 (1996) De Micco C, Ruf J, Chrestian MA et al. Cancer 67(12): 3036-3041 (1991) Kimura S, Kotani T, McBride O et al. Proceedings of the National Academy of Science USA 84, 5555-5559, (1987).



# Instructions for Use

## Heat Induced Epitope Retrieval Combined With Polymer Detection For Immunohistochemical Demonstration On Paraffin Sections

1. Cut and mount sections on slides coated with a suitable tissue adhesive.
2. Deparaffinize sections and rehydrate to distilled water.
3. Place sections in 0.5% hydrogen peroxide/methanol for 10 minutes (or use other appropriate endogenous peroxidase blocking procedure). Wash sections in tap water.
4. Heat 1500 mL of the recommended epitope retrieval solution (Citrate based pH 6.0 - Epitope Retrieval Solution unless otherwise indicated overleaf) in a stainless steel pressure cooker until boiling. Cover but do not lock lid.
5. Position slides into metal staining racks (do not place slides close together as uneven staining may occur) and lower into pressure cooker ensuring slides are completely immersed in epitope retrieval solution. Lock lid.
6. When the pressure cooker reaches operating temperature and pressure (after about 5 minutes) start a timer for 1 minute (unless otherwise indicated on the data sheet).
7. When the timer rings, remove pressure cooker from heat source and run under cold water with lid on. DO NOT OPEN LID UNTIL THE INDICATORS SHOW THAT PRESSURE HAS BEEN RELEASED. Open lid, remove slides and place immediately into a bath of tap water.
8. Wash sections once using fresh Tris-Buffered Saline (TBS, pH 7.6) buffer for 5 minutes.
9. Place sections in diluted normal serum (eg NCL-G-SERUM) for 10 minutes.
10. Incubate sections with primary antibody.
11. Wash twice, each time using fresh TBS buffer for 5 minutes.
12. For visualization of the bound primary antibody, follow instructions supplied with the Polymer Detection System.
13. Counterstain with hematoxylin (if required), dehydrate and mount.

*\* (In most applications, Phosphate Buffered Saline, pH 7.6, can be used instead of TBS, pH 7.6).*

### Safety Note

To ensure the correct and safe use of your pressure cooker, PLEASE READ THE MANUFACTURER'S INSTRUCTIONS.