

Product Data Sheet

Anti-Thyroid Peroxidase Antibody

Catalog #	Source	Reactivity	Applications		
CPA9637	Mouse	Н	IH		
Description	N	Mouse monoclonal antibody to Thyroid Peroxidase			
Immunogen	K	LH-conjugated synthetic p	eptide encompassing a sequence within human Thyroid		
	Pe	eroxidase. The exact sequ	ence is proprietary.		
Purification	TI	he antibody was purified	by immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous lev	els of Thyroid Peroxidase protein.		
Clonality	Ν	Ionoclonal			
Conjugation					
Form	N	1ouse IgG2b. Liquid in PB	containing 50% glycerol, 0.2% BSA and 0.01% sodium		
	az	zide.			
Dilution	IF	H (1/100 - 1/300)			
Gene Symbol	TI	РО			
Alternative Na	ames Tl	hyroid peroxidase; TPO			
Entrez Gene	7	173 (Human)			
SwissProt	P	07202 (Human)			
Storage/Stabi	lity Sł	hipped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid		
	fr	eeze/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

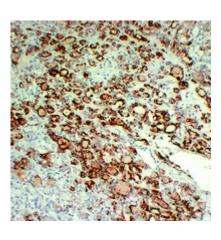
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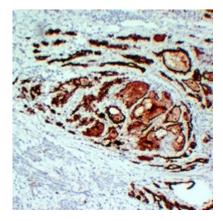


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Immunohistochemical analysis of Thyroid Peroxidase staining in human follicular thyroid carcinoma formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The s



Immunohistochemical analysis of Thyroid Peroxidase staining in human papillary thyroid carcinoma formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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