

Product datasheet

Anti-Thyroglobulin antibody [EPR3614(2)] - BSA and Azide free ab248992

Recombinant RabMAb

7 Images

Overview		
Product name	Anti-Thyroglobulin antibody [EPR3614(2)] - BSA and Azide free	
Description	Rabbit monoclonal [EPR3614(2)] to Thyroglobulin - BSA and Azide free	
Host species	Rabbit	
Tested applications	Suitable for: WB, IHC-P, mIHC	
Species reactivity	Reacts with: Human	
Immunogen	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.	
General notes	ab248992 is the carrier-free version of <u>ab151539</u> .	
	Our <u>carrier-free</u> antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.	
	This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.	
	Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.	
	This product is compatible with the Maxpar [®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar [®] is a trademark of Fluidigm Canada Inc.	
	 This product is a recombinant monoclonal antibody, which offers several advantages including: High batch-to-batch consistency and reproducibility Improved sensitivity and specificity Long-term security of supply Animal-free production For more information <u>see here</u>. Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <u>RabMAb[®] patents</u>. 	

Form	Liquid
Storage instructions	Shipped at 4°C. Store at +4°C. Do Not Freeze.
Storage buffer	pH: 7.2 Constituent: PBS
Carrier free	Yes
Purity	Protein A purified
Clonality	Monoclonal
Clone number	EPR3614(2)
lsotype	lgG

Applications

The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab248992 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

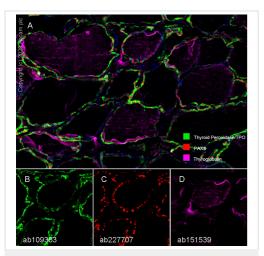
Application	Abreviews	Notes
WB		Use at an assay dependent concentration. Predicted molecular weight: 305 kDa.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
mIHC		Use at an assay dependent concentration.

Target		
Function	Precursor of the iodinated thyroid hormones thyroxine (T4) and triiodothyronine (T3).	
Tissue specificity	Thyroid gland specific.	
Involvement in disease	 Defects in TG are the cause of congenital hypothyroidism due to dyshormonogenesis type 3 (CHDH3) [MIM:274700]. A disorder due to thyroid dyshormonogenesis, causing large goiters of elastic and soft consistency in the majority of patients. Although the degree of thyroid dysfunction varies considerably among patients with defective thyroglobulin synthesis, patients usually have a relatively high serum free triiodothyronine (T3) concentration with disproportionately low free tetraiodothyronine (T4) level. The maintenance of relatively high free T3 levels prevents profound tissue hypothyroidism except in brain and pituitary, which are dependent on T4 supply, resulting in neurologic and intellectual defects in some cases. Variations in TG are associated with susceptibility to autoimmune thyroid disease type 3 (AITD3) [MIM:608175]. AITDs including Graves disease (GD) and Hashimoto thyroiditis (HT), are among the most common human autoimmune diseases. They are complex diseases, which are caused by an interaction between susceptibility genes and nongenetic factors, such as infection. 	
Sequence similarities	Belongs to the type-B carboxylesterase/lipase family. Contains 11 thyroglobulin type-1 domains.	
Post-translational	Sulfated tyrosines are desulfated during iodination.	

Cellular localization

Secreted.

Images



Multiplex immunohistochemistry - Anti-Thyroglobulin antibody [EPR3614(2)] - BSA and Azide free (ab248992) Fluorescence multiplex immunohistochemical analysis of Human thyroid gland tissue (Formalin/PFA-fixed paraffin-embedded sections).

Panel A: merged staining of anti-Thyroglobulin (<u>ab151539</u>, magenta; Opal[™]690), anti-Thyroid Peroxidase/TPO (<u>ab109383</u>, green; Opal[™]520) and anti-PAX8 (<u>ab227707</u>, red; Opal[™]570) on human thyroid gland. Panel B: anti-Thyroid Peroxidase/TPO stained on cytoplasm of follicular epithelial cells. Panel C: anti-PAX8 stained on nucleus of follicular epithelial cells. Panel D: anti-Thyroglobulin stained on colloid. Opal Polymer HRP Ms + Rb was used as a secondary antibody.

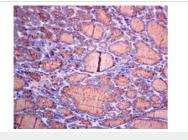
The section was incubated in three rounds of staining: in the order of **<u>ab151539</u>** at 1/5000 dilution (0.167 µg/ml), **<u>ab109383</u>** at 1/1500 dilution (0.072 µg/ml), and **<u>ab227707</u>** at 1/200 dilution (5.22 µg/ml) for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal[™] 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope. DAPI (blue) was used as a nuclear counter stain.

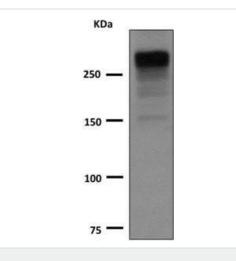
Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

This data was developed using <u>ab151539</u>, the same antibody clone in a different buffer formulation

This data was developed using **ab151539**, the same antibody clone in a different buffer formulation.Immunohistochemical analysis of paraffin-embedded Human thyroid gland follicular carcinoma tissue labeling Thyroglobulin with **ab151539** at 1/250 dilution. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Thyroglobulin antibody [EPR3614(2)] - BSA and Azide free (ab248992)



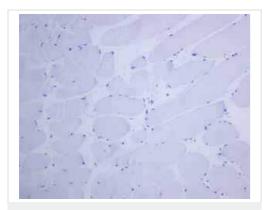
Western blot - Anti-Thyroglobulin antibody [EPR3614(2)] - BSA and Azide free (ab248992) Anti-Thyroglobulin antibody [EPR3614(2)] (ab151539) at 1/100000 dilution + Human thyroid tissue lysate at 10 µg

Secondary

Goat anti-rabbit HRP conjugated at 1/2000 dilution

Predicted band size: 305 kDa

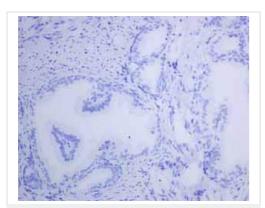
This data was developed using <u>ab151539</u>, the same antibody clone in a different buffer formulation.



This data was developed using <u>ab151539</u>, the same antibody clone in a different buffer formulation.Immunohistochemical analysis of paraffin embedded Human Skeletal muscle tissue using <u>ab151539</u> showing -ve staining.

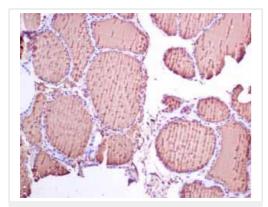
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Thyroglobulin antibody [EPR3614(2)] - BSA and Azide free (ab248992)

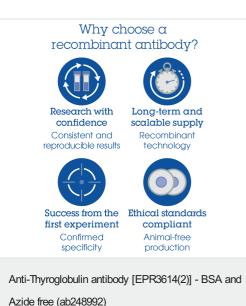


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Thyroglobulin antibody [EPR3614(2)] - BSA and Azide free (ab248992) This data was developed using <u>ab151539</u>, the same antibody clone in a different buffer formulation.Immunohistochemical analysis of paraffin embedded normal Human prostate tissue using <u>ab151539</u> showing -ve staining.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Thyroglobulin antibody [EPR3614(2)] - BSA and Azide free (ab248992)



This data was developed using <u>ab151539</u>, the same antibody clone in a different buffer formulation.Immunohistochemical analysis of paraffin embedded Human Thyroid gland tissue using <u>ab151539</u> showing +ve staining.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

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