

## Product datasheet

# Anti-Thyroid Peroxidase antibody [6H7] ab10246

### 1 References

#### Overview

<b>Product name</b>	Anti-Thyroid Peroxidase antibody [6H7]
<b>Description</b>	Mouse monoclonal [6H7] to Thyroid Peroxidase
<b>Specificity</b>	This antibody recognizes specifically human thyroid peroxidase.
<b>Tested applications</b>	<b>Suitable for:</b> WB
<b>Species reactivity</b>	<b>Reacts with:</b> Human
<b>Immunogen</b>	Recombinant full length protein (Human).
<b>General notes</b>	Concentration varies from lot to lot and can be provided on request.

#### Properties

<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
<b>Storage buffer</b>	Preservative: 0.1% Sodium Azide Constituents: PBS, pH 7.4
<b>Purity</b>	Protein A purified
<b>Purification notes</b>	Purity tested by electrophoresis.
<b>Clonality</b>	Monoclonal
<b>Clone number</b>	6H7
<b>Myeloma</b>	Sp2/0
<b>Isotype</b>	IgG1

#### Applications

Our [Abpromise guarantee](#) covers the use of **ab10246** 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
AP		Use at an assay dependent dilution.

Application	Abreviews	Notes
WB		Use at an assay dependent dilution. Predicted molecular weight: 103 kDa.

## Target

<b>Function</b>	Iodination and coupling of the hormonogenic tyrosines in thyroglobulin to yield the thyroid hormones T(3) and T(4).
<b>Pathway</b>	Hormone biosynthesis; thyroid hormone biosynthesis.
<b>Involvement in disease</b>	Note=An alternative splicing in the thyroperoxidase mRNA can cause Graves' disease. Defects in TPO are the cause of congenital hypothyroidism due to dyshormonogenesis type 2A (CHDH2A) [MIM:274500]; also called genetic defect in thyroid hormonogenesis 2A or thyroid hormone organification defect II. CHDH2A is due to defective conversion of accumulated iodide to organically bound iodine. The iodide organification defect can be partial or complete.
<b>Sequence similarities</b>	Belongs to the peroxidase family. XPO subfamily. Contains 1 EGF-like domain. Contains 1 Sushi (CCP/SCR) domain.
<b>Post-translational modifications</b>	Glycosylated. Heme is covalently bound through a H(2)O(2)-dependent autocatalytic process. Heme insertion is important for the delivery of protein at the cell surface. Cleaved in its N-terminal part.
<b>Cellular localization</b>	Membrane and Cell surface.

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