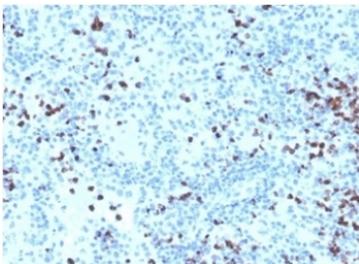


## MPO Antibody / Myeloperoxidase (V9351)

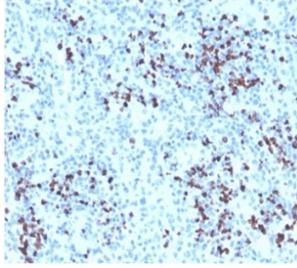
Catalog No.	Formulation	Size
V9351-100UG	0.2 mg/ml in 1X PBS with 0.1 mg/ml BSA (US sourced), 0.05% sodium azide	100 ug
V9351-20UG	0.2 mg/ml in 1X PBS with 0.1 mg/ml BSA (US sourced), 0.05% sodium azide	20 ug
V9351SAF-100UG	1 mg/ml in 1X PBS; BSA free, sodium azide free	100 ug

[Bulk quote request](#)

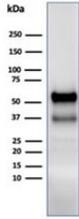
<b>Availability</b>	1-3 business days
<b>Species Reactivity</b>	Human
<b>Format</b>	Purified
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal (rabbit origin)
<b>Isotype</b>	Rabbit IgG, kappa
<b>Purity</b>	Protein A/G affinity
<b>UniProt</b>	P05164
<b>Localization</b>	Cytoplasmic
<b>Applications</b>	Western Blot : 2-4ug/ml Immunohistochemistry (FFPE) : 1-2ug/ml
<b>Limitations</b>	This Myeloperoxidase antibody is available for research use only.



IHC staining of FFPE human spleen tissue with Myeloperoxidase antibody. HIER: boil tissue sections in pH 9 10mM Tris with 1mM EDTA for 20 min and allow to cool before testing.



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Western blot testing of human spleen lysate using Myeloperoxidase antibody. Expected molecular weight: 59-64 kDa (alpha chain, may be observed at higher molecular weights due to glycosylation), 150+ kDa (glycosylated mature form).

## Description

The heme protein myeloperoxidase (MPO) is a major component of azurophilic granules of neutrophils and polymorphonuclear leukocytes. Optimal oxygen-dependent microbiocidal activity depends on MPO as the critical enzyme for the generation of hypochlorous acid and other toxic oxygen products. The MPO precursor is synthesized during the promyelocytic stage of myeloid differentiation and is subsequently processed and transported intracellularly to the lysosomes. The precursor undergoes cotranslational N-linked glycosylation to produce a glycoprotein. Glucosidases in the endoplasmic reticulum (ER) or early cisGolgi convert the pro-MPO to a form which is sorted into a prelysosomal compartment, which undergoes final proteolytic maturation to native MPO, a pair of heavy-light protomers. In normal neutrophils, MPO is expressed as a dimer. Calreticulin, a calcium-binding protein residing in the ER, interacts specifically with fully glycosylated apopro-MPO. MPO mRNA is abundant in human promyelocytic HL-60 and mouse myeloid leukemia NFS-60 cells. MPO is expressed at high levels in circulating neutrophils and monocytes but is not detectable in microglia, brain-specific macrophages or normal brain tissue. MPO, which has a molecular weight of approximately 140 kD, is a homodimer that can be split into two halves that still have enzymatic activity. These hemi-MPO monomers consist of a 59-kD alpha chain and a 13.5-kD beta chain.

## Application Notes

Optimal dilution of the Myeloperoxidase antibody should be determined by the researcher.

## Immunogen

A portion of amino acids 150-250 of human Myeloperoxidase protein was used as the immunogen for the Myeloperoxidase antibody.

## Storage

Aliquot the Myeloperoxidase antibody and store frozen at -20°C or colder. Avoid repeated freeze-thaw cycles.

