

#### FabGennix International, Inc.

9191 Kyser Way Bldg. 4 Suite 402 Frisco, TX 75033

Tel: (214)-387-8105, 1-800-786-1236 Fax: (214)-387-8105 Email: info@fabgennix.com Web: www.FabGennix.com

# Rabbit Polyclonal Anti-Myeloperoxidase antibody

Catalog Number: MPO-131AP

Lot Number:

#### **General Information**

Product	Myeloperoxidase Antibody C-terminus
Accession #	Uniprot: P05164
	GenBank: AAB25582.1
Verified Applications	CM, ELISA, ICC, IF, IHC, IP, WB
Species Cross Reactivity	Human, Mouse, Rat
Host	Rabbit
Immunogen	Synthetic peptide taken within amino acid region
	700-745 human Myeloperoxidase protein
Alternative Nomenclature	84 kDa myeloperoxidase antibody, 89 kDa
	myeloperoxidase antibody, EC 1.11.1.7 antibody,
	EC1.11.2.2 antibody, MPO antibody,
	Myeloperoxidase heavy chain antibody,
	Myeloperoxidase light chain antibody,
	PERM_HUMAN antibody

## **Physical Properties**

Quantity	100 μg
Volume	200 µl
Form	Affinity Purified Immunoglobulins
Determinant	Extreme C-epitope
Immunoglobulin & Concentration	0.75-1.2 mg/ml IgG in antibody stabilization buffer
Storage	Store at -20°C for long term storage.

## **Recommended Dilutions**

DOT Blot	1:10,000
ELISA	1:10,000
Immunocytochemistry	1:200
Immunofluorescence	1:200
Immunohistochemistry	1:200
Immunoprecipitation	1:200
Western Blot	1:500

Tel: (214)-387-8105, 1-800-786-1236 Fax: (214)-387-8105 Email: info@fabgennix.com Web: www.FabGennix.com

### Related Products

FITC-Conjugated	MPO.131-FITC
BIOTIN-Conjugated	MPO.131-BIOTIN
Antigenic Blocking Peptide	P-MPO.131
Western Blot Positive Control	PC-MPO
N-epitope	MPO-101AP
Mid-Region	MPO-112AP
C-epitope	MPO-121AP
Mixed-epitope	MPO-141AP

Catalog #

#### Overview:

Myeloperoxidase (MPO) is a functionally important component of the normal human neutrophil azurophil granule of polymorphonuclear leukocyte's host defense system. MPO is a hemoprotein that is abundantly expressed in neutrophils and secreted during their activation. The inflammatory component of allergic reactions can be measured by eosinophil cationic protein (ECP, specific for eosinophils) and myeloperoxidase (MPO, specific for neutrophils) measurement (1). Neutrophils, especially in acute infection or the myeloid leukemias, may shed platelet-sized particles that can readily be distinguished from true platelets because they contain neutrophil myeloperoxidase, an enzyme that is not inhibited by glutaraldehyde. Traditionally Myeloperoxidase was considered as a serological marker for granulomatosis (Churg-Strauss syndrome), the main target of anti-neutrophil cytoplasm antibodies (ANCA), Low to moderate anti-Myeloperoxidase autoantibody levels are also reported in rheumatoid arthritis. Recently it was shown that MPO participates in the initiation and progression of cardiovascular disease. It possesses potent pro-inflammatory properties and may contribute directly to tissue injury. Now Myeloperoxidase is uncertain systemic vasculitis e.g. periarteritis nodosa, microscopic polyarteritis and pulmonary eosinophilic grander consideration as one of the most promising cardiac markers. Human myeloperoxidase, a 745 amino acid protein, is produced by a gene composed of 12 exons and 11 introns. The sequence was found to contain an open reading frame, 2,235 nucleotides coding for a protein of 745 amino acids with a calculated Mr of 83,868. The heavy chain of myeloperoxidase, consisting of 467 amino acids, was located on the COOH terminus half of the protein. Native Myeloperoxidase is represented as a covalently bound tetrameric complex of two glycosylated alpha chains (MW 59 - 64 kDa) and two unglycosylated beta chains (MW 14 kDa) with total MW 150 kDa and theoretical pl 9.2, each of which is associated with a heme-like prosthetic group. The heme environment, defined by X-ray crystallographic analyses of MPO appears to be highly conserved in four mammalian peroxidases (lactoperoxidase, thyroid, eosinophil and Myeloperoxidase). MPO when heated under non-reducing, denaturing conditions, MPO is cleaved to produced 22 and a 38kDa proteins, these two species are produced via a novel autolytic cleavage at Meth409 and Pro410 (2).

#### References:

- 1. Linder A, Venge P, Deuschl H.Allergy. Eosinophil cationic protein and myeloperoxidase in nasal secretion as markers of inflammation in allergic rhinitis. 1987 Nov;42(8):583-90.
- 2. Taylor KL, Pohl J, Kinkade JM Jr. Unique autolytic cleavage of human myeloperoxidase. Implications for the involvement of active site MET409. J Biol Chem. 1992 Dec 15;267(35):25282-8.

Tel: (214)-387-8105, 1-800-786-1236 Fax: (214)-387-8105 Email: info@fabgennix.com Web: www.FabGennix.com

<sup>\*</sup> For users who may require large amounts of the products listed above, please inquire about bulk material discounts. This Product is for Research Use Only and is NOT intended for use in humans or clinical diagnosis