bs-4943R

[Primary Antibody]

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MPO Rabbit pAb

DATASHEET -

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

GeneID: 4353 **SWISS:** P05164

Target: MPO

Immunogen: KLH conjugated synthetic peptide derived from human

Myeloperoxidase heavy chain: 678-745/745.

Purification: affinity purified by Protein A

Concentration: 1mg/ml

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50%

Glycerol.

Shipped at 4°C. Store at -20°C for one year. Avoid repeated

freeze/thaw cycles.

Background: Myeloperoxidase (MPO) is a heme protein synthesized during myeloid differentiation that constitutes the major component of neutrophil azurophilic granules. Produced as a single chain precursor, myeloperoxidase is subsequently cleaved into a light and heavy chain. The mature myeloperoxidase is a tetramer composed of 2 light chains and 2 heavy chains. This enzyme produces hypohalous acids central to the microbicidal activity of

neutrophils. [provided by RefSeq, Nov 2014]

Applications: IHC-P (1:100-500)

IHC-F (1:100-500) **IF** (1:100-500) ICC/IF (1:100-500)

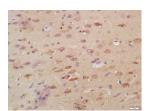
Reactivity: Human, Mouse

(predicted: Rat, Rabbit, Dog, GuineaPig, Horse)

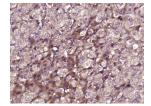
Predicted 84 kDa

Subcellular Cytoplasm Location:

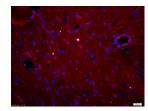
VALIDATION IMAGES



Tissue/cell: mouse brain tissue: 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Incubation: Anti-MPO Polyclonal Antibody, Unconjugated(bs-4943R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Paraformaldehyde-fixed, paraffin embedded (Human liver carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min: Antibody incubation with (MPO) Polyclonal Antibody, Unconjugated (bs-4943R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Tissue/cell: mouse brain tissue:4% Paraformaldehyde-fixed and paraffinembedded: Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min: Incubation: Anti-MPO Polyclonal Antibody, Unconjugated(bs-4943R) 1:200, overnight at 4°C: The secondary antibody was Goat Anti-Rabbit IgG, Cy3 conjugated(bs-0295G-Cy3)used at 1:200 dilution for 40 minutes at 37°C. DAPI(5ug/ml,blue,C-0033) was used to stain the cell nuclei

— SELECTED CITATIONS ——

- [IF=11.53] Ye, Buging. et al. Induction of functional neutrophils from mouse fibroblasts by thymidine through enhancement of Tet3 activity. Cell Mol Immunol. 2022 Mar;:1-15 IF; E.Coli. 35301470
- [IF=5.6] Yaxi Zhou. et al. Silkworm pupa protein peptide improved DSS-induced colitis in C57BL/6 mice through the МАРК/NF-кВ signaling pathway. J FUNCT FOODS. 2023 Nov;110:105852 IHC; Mouse. 10.1016/j.jff.2023.105852
- [IF=5.7] Yang Li. et al. Deciphering hub genes and immune landscapes related to neutrophil extracellular traps in rheumatoid arthritis: insights from integrated bioinformatics analyses and experiments.FRONTIERS IN IMMUNOLOGY.2025

Jan 8:15:1521634. IF ; Human. 39845946

- [IF=5.595] Chen B et al. A self-organized actomyosin drives multiple intercellular junction disruption and directly promotes neutrophil recruitment in lipopolysaccharide-induced acute lung injury.FASEB J. 2018 Jun 7:fj201701506RR. ICC ;Human. 29879372
- [IF=4.755] Bae Ju-Eun. et al. Effects of erythropoietin on osteoblast in the tooth extraction socket in mice periodontitis model. FRONT PHYSIOL. 2022 Oct;0:2117 IHC; Mouse. 36277197