





MMP7 (Cleaved-Tyr95) mouse mAb

Catalog No	YP-mAb-02313
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	MMP7 MPSL1 PUMP1
Protein Name	MMP7 (Cleaved-Tyr95)
Immunogen	Synthesized peptide derived from human MMP7 (Cleaved-Tyr95)
Specificity	This antibody detects endogenous levels of Human,Mouse,Rat MMP7 (Cleaved-Tyr95, protein was cleaved amino acid sequence between 94-95)
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	Matrilysin (EC 3.4.24.23;Matrin;Matrix metalloproteinase-7;MMP-7;Pump-1 protease;Uterine metalloproteinase)
Observed Band	19 29kD
Cell Pathway	Secreted, extracellular space, extracellular matrix .
Tissue Specificity	
Function	proteolysis, collagen catabolic process, collagen metabolic process, regulation of cell proliferation, multicellular organismal metabolic process, multicellular organismal catabolic process, multicellular organismal macromolecule metabolic process,
Background	catalytic activity:Cleavage of 14-Ala- -Leu-15 and 16-Tyr- -Leu-17 in B chain of insulin. No action on collagen types I, II, IV, V. Cleaves gelatin chain alpha-2(I) > alpha-1(I).,cofactor:Binds 2 calcium ions per subunit.,cofactor:Binds 2 zinc ions per subunit.,domain:The conserved cysteine present in the cysteine-switch motif binds the catalytic zinc ion, thus inhibiting the enzyme. The dissociation of the cysteine from the zinc ion upon the activation-peptide release activates the enzyme.,function:Degrades casein, gelatins of types I, III, IV, and V, and fibronectin. Activates procollagenase.,similarity:Belongs to the peptidase M10A family.,



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matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

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