



TIMP-3 Monoclonal Antibody

Catalog No	YP-mAb-02798
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	TIMP3
Immunogen	The antiserum was produced against synthesized peptide derived from human TIMP3. AA range:91-140
Specificity	TIMP-3 Monoclonal Antibody detects endogenous levels of TIMP-3 protein.
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	TIMP3;Protein MIG-5;Protein MIG 5;Metalloproteinase inhibitor 3;K222TA2;TIMP-3;Tissue inhibitor of metalloproteinases 3
Observed Band	25kD
Calculated Molecular Weight	25kD
Cell Pathway	Secreted, extracellular space, extracellular matrix.
Tissue Specificity	Kidney,Mammary carcinoma,Pancreas,Placenta,Retina,Uterus,
Function	disease:Defects in TIMP3 are the cause of Sorsby fundus dystrophy (SFD) [MIM:136900]. SFD is a rare autosomal dominant macular disorder with an age of onset in the fourth decade. It is characterized by loss of central vision from subretinal neovascularization and atrophy of the ocular tissues. Generally, macular disciform degeneration develops in the patients eye within 6 months to 6 years.,function:Complexes with metalloproteinases (such as collagenases) and irreversibly inactivates them. May form part of a tissue-specific acute response to remodeling stimuli. Known to act on MMP-1, MMP-2, MMP-3, MMP-7, MMP-9, MMP-13, MMP-14 and MMP-15.,online information:Retina International's Scientific Newsletter,similarity:Belongs to the protease inhibitor I35 (TIMP) family.,similarity:Contains 1 NTR domain.,



Background

This gene belongs to the TIMP gene family. The proteins encoded by this gene family are inhibitors of the matrix metalloproteinases, a group of peptidases involved in degradation of the extracellular matrix (ECM). Expression of this gene is induced in response to mitogenic stimulation and this netrin domain-containing protein is localized to the ECM. Mutations in this gene have been associated with the autosomal dominant disorder Sorsby's fundus dystrophy. [provided by RefSeq, Jul 2008],

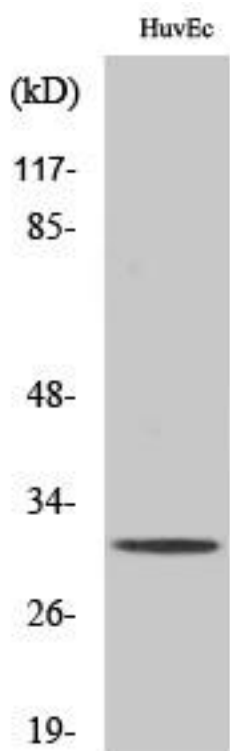
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

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

Products Images



Western Blot analysis of various cells using TIMP-3 Monoclonal Antibody