Elabscience®

MMP-1 Polyclonal Antibody

catalog number: D-AB-10358L

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description			
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Reactivity	Human Becombinent Human MMB 1 protein supressed by Easli		
Immunogen	Recombinant Human MMP-1 protein expressed by E.coli		
Host	Rabbit		
Isotype	IgG		
Purification	Antigen Affinity Purification		
Conjugation	Unconjugated		
Buffer	PBS with 0.05% Proclin300	PBS with 0.05% Proclin300, 1% protective protein and 50% glycerol, pH7.4	
Applications	Recommended Dilution		
IHC	1:300-1:600		
Data			
•	of paraffin-embedded Human lung Polyclonal Antibody at dilution of	Immunohistochemistry of paraffin-embedded Human cervical cancer using MMP-1 Polyclonal Antibody at	
cancer using MMP-1 P	-		
cancer using MMP-1 P Preparation & Storage	Polyclonal Antibody at dilution of 1:600	cervical cancer using MMP-1 Polyclonal Antibody at dilution of 1:600	
cancer using MMP-1 P Preparation & Storage Storage	Polyclonal Antibody at dilution of 1:600 Store at -20°C Valid for 12 i	cervical cancer using MMP-1 Polyclonal Antibody at dilution of 1:600 months. Avoid freeze / thaw cycles.	
cancer using MMP-1 P Preparation & Storage	Polyclonal Antibody at dilution of 1:600 Store at -20°C Valid for 12 i	cervical cancer using MMP-1 Polyclonal Antibody at dilution of 1:600	

Background

Matrix metalloproteinases are a family of zinc and calcium dependent endopeptidases with the combined ability to degrade all the components of the extracellular matrix MMP-1 (interstitial collagenase), can degrade a broad range of substrates including types I, II, III, VII, VIII, and X collagens as well as casein, gelatin, alpha -1 antitrypsin, myelin basic protein, L-Selectin, pro-TNF, IL-1 beta, IGFBP-3, IGFBP-5, pro-MMP-2, and pro-MMP-9. A significant role of MMP-1 is the degradation of fibrillar collagens in extracellular matrix remodeling, characterized by the cleavage of the interstitial collagen triple helix into ³/₄, ¹/₄ fragments. However, as the list of substrates above illustrates, the role of MMP-1 is more diverse than originally envisaged, and may involve enzyme cascades, cytokine regulation, and cell surface molecule modulation. MMP-1 is expressed by fibroblasts, keratinocytes, endothelial cells, monocytes, and macrophages. Structurally, MMP-1 may be divided into several distinct domains; a pro-domain which is cleaved upon activation; a catalytic domain containing the zinc binding site; a short hinge region and a carboxyl terminal (hemopexin-like) domain.