abcam

Product datasheet

Anti-MMP3 antibody ab53015

★★★★★ 7 Abreviews 111 References 画像数 5

製品の概要

製品名 Anti-MMP3 antibody

製品の詳細 Rabbit polyclonal to MMP3

由来種 Rabbit

アプリケーション 適用あり: IHC-P, WB, ICC/IF

種交差性 交差種: Human

免疫原 Synthetic peptide derived from human MMP3 Peptide available as <u>ab134501</u>.

特記事項

The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

バッファー pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 50% Glycerol, 0.87% Sodium chloride, PBS

精製度 Immunogen affinity purified

ポリ/モノ ポリクローナル

アイソタイプ IgG

アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab53015の使用に適用されます

アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
IHC-P	★★★★ (3)	1/50 - 1/100.
WB	★★★★☆ (2)	1/500 - 1/1000. Detects a band of approximately 54 kDa (predicted molecular weight: 54 kDa).
ICC/IF	★★★★★ (1)	Use a concentration of 5 µg/ml.

ターゲット情報

機能 Can degrade fibronectin, laminin, gelatins of type I, III, IV, and V; collagens III, IV, X, and IX, and

cartilage proteoglycans. Activates procollagenase.

配列類似性 Belongs to the peptidase M10A family.

Contains 4 hemopexin-like domains.

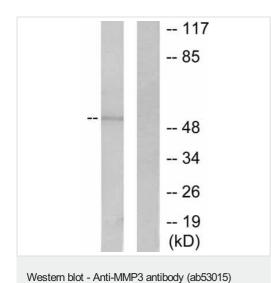
ドメイン The conserved cysteine present in the cysteine-switch motif binds the catalytic zinc ion, thus

inhibiting the enzyme. The dissociation of the cysteine from the zinc ion upon the activation-

peptide release activates the enzyme.

細胞内局在 Secreted > extracellular space > extracellular matrix.

画像



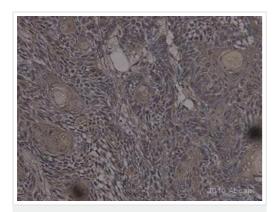
All lanes: Anti-MMP3 antibody (ab53015) at 1/500 dilution

Lane 1: Extracts from 293 cells, untreated.

Lane 2: Extracts from 293 cells, treated with the immunising

peptide.

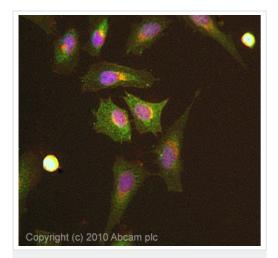
Predicted band size: 54 kDa **Observed band size:** 54 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-MMP3 antibody (ab53015)

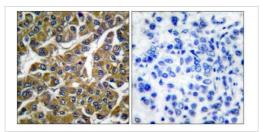
This image is courtesy of an anonymous abreview.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human lymphoma labeling MMP3 with ab53015 at 1/100 dilution. Tissue was fixed with formaldehyde; heat-mediated antigen retrieval was performed using citrate buffer (pH 6.0). The tissue was blocked with 1% serum for 1 hour at 25°C. A biotin conjugated polyclonal goat anti-rabbit secondary antibody was used at 1/200 dilution.



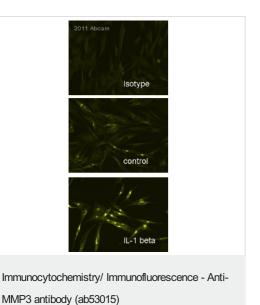
Immunocytochemistry/ Immunofluorescence - Anti-MMP3 antibody (ab53015)

ICC/IF image of ab53015 stained HeLa cells. The cells were 4% formaldehyde fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab53015, 1 μ g/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit lgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43 μ M.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-MMP3 antibody (ab53015)

Immunohistochemical analysis of paraffin-embedded human breast carcinoma tissue using MMP3 antibody at a dilution of 1/50. Left: without immunizing peptide; Right: with immunizing peptide.



Top image: Isotype control Middle image: Unstimulated cells Bottom image: IL-1 beta stimulated cells (24 hours, 10 ng/ml) ab53015 staining MMP3 in Human synovial fibroblasts by Immunocytochemistry/ Immunofluorescence. Cells were fixed in formaldehyde and permeabilized in methanol for 15 minutes at -20°C prior to blocking in 1% BSA for 1 hour at 25°C. The primary antibody was diluted 1/1000 and incubated with the sample for 2 hours at 25°C. The secondary antibody was Cy5®-conjugated Goat anti-Rabbit polyclonal, diluted 1/200.

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Torsten Lowin.

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