abcam

Product datasheet

Anti-MMP9 antibody [56-2A4] ab58803

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Overview

Product name Anti-MMP9 antibody [56-2A4]

Description Mouse monoclonal [56-2A4] to MMP9

Host species Mouse

Specificity Recognizes the ~92 kDa latent and the ~83 kDa active forms of MMP9. Does not react with

MMP1, MMP2, MMP3 or MMP13. The antibody recognizes precursor (92 kDa) and active (83

kDa) forms of human and guinea pig MMP-9.

Tested applications Suitable for: IHC-P, IHC-Fr, ICC/IF, WB

Species reactivity Reacts with: Mouse, Rat, Rabbit, Guinea pig, Human

Immunogen Synthetic peptide corresponding to Human MMP9 aa 626-644.

Sequence:

LFSGRRLWRFDVKAQMVDP

Database link: P14780

Run BLAST with
Run BLAST with

Positive control Breast carcinoma tissue. Conditioned medium of human fibrosarcoma HT1080 cells The protein

is secreted so you may need to concentrate the protein from cell culture media. Concentrate protein by TCA, acetone or ammonium sulphate precipitation. HL60 cell lysate, U937 cell lysate,

HT1080 cell lysate, human spleen lysate

General notesThe 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

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

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Storage buffer pH: 7.00

Preservative: 0.1% Sodium azide

Constituents: 1.64% Sodium phosphate, 0.1% BSA

Purity Immunogen affinity purified

Purification notesIs purified.ClonalityMonoclonalClone number56-2A4IsotypeIgG1Light chain typekappa

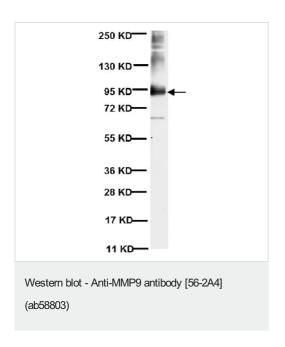
Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab58803 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

| Application | Abreviews | Notes |
|-------------|------------------------|--|
| IHC-P | ★★★★ (3) | Use at an assay dependent concentration. |
| IHC-Fr | ★★★★ <u>(3)</u> | Use at an assay dependent concentration. |
| ICC/IF | *** <u>*</u> | Use at an assay dependent concentration. PubMed: 22496779 |
| WB | ★★★★☆ (2) | Use a concentration of 2 µg/ml. Detects a band of approximately 92 kDa (predicted molecular weight: 78 kDa). |

| May play an essential role in local proteolysis of the extracellular matrix and in leukocyte migration. Could play a role in bone osteoclastic resorption. Cleaves KiSS1 at a Gly- -Leu bond. Cleaves type IV and type V collagen into large C-terminal three quarter fragments and |
|---|
| shorter N-terminal one quarter fragments. Degrades fibronectin but not laminin or Pz-peptide. |
| Produced by normal alveolar macrophages and granulocytes. |
| Intervertebral disc disease Metaphyseal anadysplasia 2 |
| Belongs to the peptidase M10A family. Contains 3 fibronectin type-II domains. Contains 4 hemopexin repeats. |
| 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. |
| Processing of the precursor yields different active forms of 64, 67 and 82 kDa. Sequentially processing by MMP3 yields the 82 kDa matrix metalloproteinase-9. N- and O-glycosylated. |
| Secreted, extracellular space, extracellular matrix. |
| |



1:500 dilution of ab58803 detected MMP-9 on 10 μ g of Human spleen lysates. Arrow indicates protein MMP-9 (~92 kDa). Lower bands indicate active forms of MMP-9

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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