# abcam

# Product datasheet

# Anti-MMP14 antibody - Hinge region ab38971

★★★★★ 5 Abreviews 39 References 画像数 3

#### 製品の概要

製品名 Anti-MMP14 antibody - Hinge region

製品の詳細 Rabbit polyclonal to MMP14 - Hinge region

由来種 Rabbit

特異性 This antibody binds to MMP14, but does not cross react with the other MMP family members

(MMP1, MMP2, MMP3, MMP9).

 アプリケーション
 適用あり: WB

 種交差性
 交差種: Human

免疫原 Synthetic peptide corresponding to Human MMP14.

(Peptide available as ab41091)

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

term. Avoid freeze / thaw cycle.

パッファー Preservative: 0.05% Sodium azide

Constituent: 50% Glycerol

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ab38971の使用に適用されます

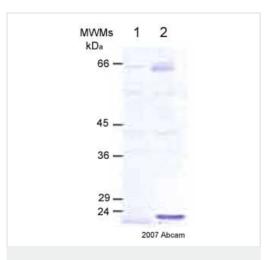
アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB	★★★★☆ (3)	1/2000 - 1/5000. Predicted molecular weight: 66 kDa. When used against the reduced protein this antibody identifies bands at 65 Kd and 63 Kd (the pro-form and active form) as well as activation/breakdown products.

## ターゲット情報

機能	Seems to specifically activate progelatinase A. May thus trigger invasion by tumor cells by activating progelatinase A on the tumor cell surface. May be involved in actin cytoskeleton reorganization by cleaving PTK7.
組織特異性	Expressed in stromal cells of colon, breast, and head and neck. Expressed in lung tumors.
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.
翻訳後修飾	The precursor is cleaved by a furin endopeptidase.
細胞内局在	Membrane. Melanosome. Identified by mass spectrometry in melanosome fractions from stage I to stage IV.

### 画像



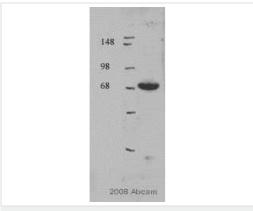
Western blot - Anti-MMP14 antibody - Hinge region (ab38971)

**All lanes :** Anti-MMP14 antibody - Hinge region (ab38971)

Lane 1 : Cell lysates from human dermal fibroblasts neonatal (no

**Lane 2**: Cell lysates from human dermal fibroblasts *neonatal* (treated with II1beta)

Predicted band size: 66 kDa Observed band size: 66 kDa



(ab38971)

This image is courtesy of an anonymous Abreview

Western blot - Anti-MMP14 antibody - Hinge region

1 2

Western blot - Anti-MMP14 antibody - Hinge region (ab38971)

This image is a courtesy of an anonymous Abreview

Anti-MMP14 antibody - Hinge region (ab38971) at 1/1000 dilution + HT1080 whole cell lysate at 30 µg

#### Secondary

HRP-conjugated donkey anti-rabbit

Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 66 kDa Observed band size: 64 kDa

Exposure time: 1 minute

All lanes: Anti-MMP14 antibody - Hinge region (ab38971) at 1/5000 dilution

Lane 1: Whole cell lysate prepared from HT1080 cells

Lane 2: Whole cell lysate prepared from human vascular smooth muscle cells

Lysates/proteins at 5 µg per lane.

#### **Secondary**

All lanes: HRP conjugated donkey anti-rabbit lgG at 1/2000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 66 kDa Observed band size: 63,65 kDa

Additional bands at: 20 kDa (possible non-specific binding)

Exposure time: 30 seconds

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

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