abcam

Product datasheet

Anti-TIMP1 antibody [EPR1550] ab109125



ייבע RabMAb

27 References 画像数9

製品の概要

製品名 Anti-TIMP1 antibody [EPR1550]

製品の詳細 Rabbit monoclonal [EPR1550] to TIMP1

由来種 Rabbit

アプリケーション 適用あり: WB, IHC-P

適用なし: Flow Cyt,ICC/IF or IP

種交差性 交差種: Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ポジティブ・コントロール HeLa cell lysate; HT1080 + TPA, HL60+TPA, PBMC lysates, human prostate lysate; Human liver

tissue; TIMP1 recombinant protein

特記事項 This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb® patents.

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with

these species. Please contact us for more information.

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.

Avoid freeze / thaw cycle. Stable for 12 months at -20°C.

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), PBS, 0.05% BSA

Protein A purified 精製度

ポリÆノ モノクローナル **ウローン名** EPR1550

アイソタイプ IgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab109125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		1/1000 - 1/10000. Predicted molecular weight: 23 kDa.
IHC-P		1/8000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. See IHC antigen retrieval protocols. For unpurified use at 1/250 - 1/500.

追加情報 Is unsuitable for Flow Cyt,ICC/IF or IP.

ターゲット情報

機能

Complexes with metalloproteinases (such as collagenases) and irreversibly inactivates them by binding to their catalytic zinc cofactor. Also mediates erythropoiesis in vitro; but, unlike IL-3, it is species-specific, stimulating the growth and differentiation of only human and murine erythroid progenitors. Known to act on MMP-1, MMP-2, MMP-3, MMP-7, MMP-8, MMP-9, MMP-10, MMP-

11, MMP-12, MMP-13 and MMP-16. Does not act on MMP-14.

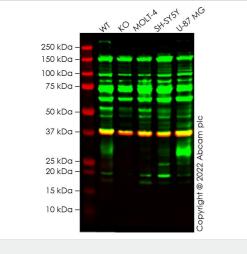
配列類似性 Belongs to the protease inhibitor I35 (TIMP) family.

Contains 1 NTR domain.

翻訳後修飾 The activity of TIMP1 is dependent on the presence of disulfide bonds.

細胞内局在 Secreted.

画像



Western blot - Anti-TIMP1 antibody [EPR1550] (ab109125)

All lanes: Anti-TIMP1 antibody [EPR1550] (ab109125) at 1/1000

dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: TIMP1 knockout HeLa cell lysate

Lane 3 : MOLT-4 cell lysate

Lane 4 : SH-SY5Y cell lysate

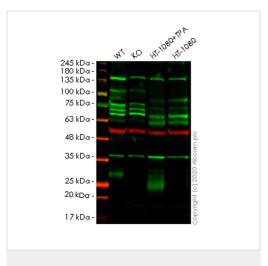
Lane 5 : U-87 MG cell lysate

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 23 kDa Observed band size: 30 kDa

False colour image of Western blot: Anti-TIMP1 antibody [EPR1550] staining at 1/1000 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] (ab8245) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab109125 was shown to bind specifically to TIMP1. A band was observed at 30 kDa in wild-type HeLa cell lysates with no signal observed at this size in TIMP1 knockout cell line ab264022 (knockout cell lysate ab260091). The identity of bands observed at higher molecular weights has not been determined. To generate this image, wild-type and TIMP1 knockout HeLa cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in fluorescent western blot (TBS-based) blocking solution before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L 800CW and Goat anti-Mouse IgG H&L 680RD at 1/20000 dilution.



Western blot - Anti-TIMP1 antibody [EPR1550] (ab109125)

All lanes : Anti-TIMP1 antibody [EPR1550] (ab109125) at 1/1000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: TIMP1 knockout HeLa cell lysate

Lane 3: HT-1080 treated with 200ng/ml 12-O-

Tetradecanoylphorbol-13-acetate (TPA) for 24 hours, cell lysate

Lane 4: Untreated HT-1080 cell lysate

Lysates/proteins at 20 µg per lane.

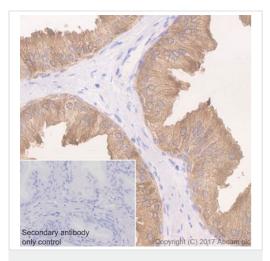
Secondary

All lanes : Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (**ab216773**) at 1/10000 dilution

Predicted band size: 23 kDa
Observed band size: 26 kDa

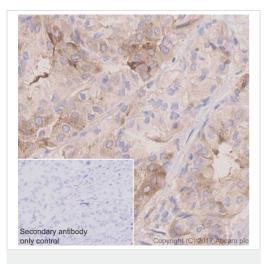
Lanes 1-4: Merged signal (red and green). Green - ab109125 observed at 26 kDa. Red - loading control <u>ab7291</u> observed at 50 kDa.

ab109125 Anti-TIMP1 antibody [EPR1550] was shown to specifically react with TIMP1 in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab261740 (knockout cell lysate ab257291) was used. Wild-type and TIMP1 knockout samples were subjected to SDS-PAGE. ab109125 and Anti-alpha Tubulin antibody [DM1A] - Loading Control (ab7291) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



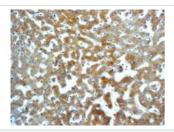
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-TIMP1 antibody
[EPR1550] (ab109125)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human prostatic hyperplasia tissue sections labeling TIMP1 with purified ab109125 at 1:8000 dilution (0.21 µg/ml). Heat mediated antigen retrieval was performed using EDTA buffer, pH 9.0. Tissue was counterstained with hematoxylin. ImmunoHistoProbe one step HRP Polymer (ready to use) secondary antibody was used at 1:0 dilution. PBS instead of the primary antibody was used as the negative control.



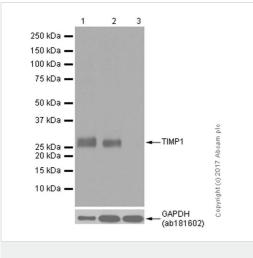
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-TIMP1 antibody
[EPR1550] (ab109125)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human thyroid carcinoma tissue sections labeling TIMP1 with purified ab109125 at 1:8000 dilution (0.21 µg/ml). Heat mediated antigen retrieval was performed using EDTA buffer, pH 9.0. Tissue was counterstained with hematoxylin. ImmunoHistoProbe one step HRP Polymer (ready to use) secondary antibody was used at 1:0 dilution. PBS instead of the primary antibody was used as the negative control.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-TIMP1 antibody
[EPR1550] (ab109125)

Unpurified ab109125, at 1/250 dilution staining TIMP1 in paraffinembedded human liver tissue, by Immunohistochemistry.



Western blot - Anti-TIMP1 antibody [EPR1550] (ab109125)



Lane 1: Human prostate lysates at 20 µg

Lane 2: HT-1080 (Human fibrosarcoma epithelial cell) treated with

200ng/ml TPA for 24 hours whole cell lysates at 15 µg

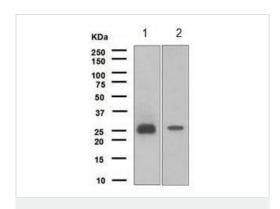
Lane 3 : Untreated with HT-1080 (Human fibrosarcoma epithelial cell) whole cell lysates at 15 μg

Secondary

All lanes : Anti-Rabbit $\lg G$ (HRP), specific to the non-reduced form of $\lg G$ at 1/2000 dilution

Predicted band size: 23 kDa

Blocking and diluting buffer: 5% NFDM/TBST



Western blot - Anti-TIMP1 antibody [EPR1550] (ab109125)

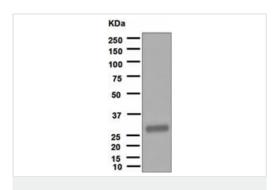
All lanes : Anti-TIMP1 antibody [EPR1550] (ab109125) at 1/1000 dilution (unpurified)

Lane 1: HL60+TPA cell lysates

Lane 2: PBMC cell lysates

Lysates/proteins at 10 µg per lane.

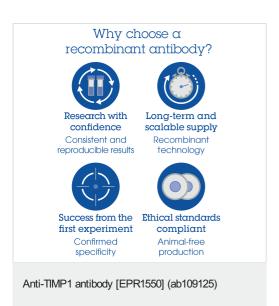
Predicted band size: 23 kDa



Western blot - Anti-TIMP1 antibody [EPR1550] (ab109125)

Anti-TIMP1 antibody [EPR1550] (ab109125) at 1/1000 dilution (unpurified) + TIMP1 recombinant protein at 10 µg

Predicted band size: 23 kDa



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