

Anti-Met (c-Met) antibody [4AT44] ab59884

★★★★★ [1 Abreviews](#) [5 References](#) [画像数 6](#)

製品の概要

製品名	Anti-Met (c-Met) antibody [4AT44]
製品の詳細	Mouse monoclonal [4AT44] to Met (c-Met)
由来種	Mouse
アプリケーション	適用あり: WB, ELISA, IHC-P, ICC/IF
種交差性	交差種: Mouse, Human
免疫原	Recombinant full length protein corresponding to Human Met (c-Met). This monoclonal antibody is generated from mice immunized with purified recombinant protein encoding the catalytic domain of human Met Database link: 4233
ポジティブ・コントロール	HepG2 cell lysate.

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.
バッファー	Preservative: 0.09% Sodium azide Constituent: 99% PBS
精製度	Protein G purified
特記事項 (精製)	ab59884 is purified through a protein G column and eluted out with both high and low pH buffers and neutralized immediately after elution then followed by dialysis against PBS.
ポリ/モノ	モノクローナル
クローン名	4AT44
アイソタイプ	IgG1

アプリケーション

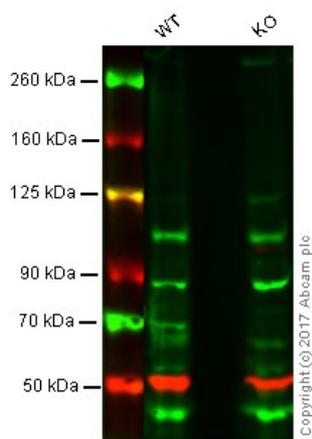
The Abpromise guarantee **Abpromise保証は、** 次のテスト済みアプリケーションにおけるab59884の使用に適用されます
アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB		1/100 - 1/500. Detects a band of approximately 135 kDa (predicted molecular weight: 156 kDa).
ELISA		1/1000.
IHC-P	★★★★★ (1)	1/50 - 1/100.
ICC/IF		1/100.

ターゲット情報

機能	Receptor for hepatocyte growth factor and scatter factor. Has a tyrosine-protein kinase activity. Functions in cell proliferation, scattering, morphogenesis and survival.
関連疾患	<p>Note=Activation of MET after rearrangement with the TPR gene produces an oncogenic protein.</p> <p>Note=Defects in MET may be associated with gastric cancer.</p> <p>Defects in MET are a cause of hepatocellular carcinoma (HCC) [MIM:114550].</p> <p>Defects in MET are a cause of renal cell carcinoma papillary (RCCP) [MIM:605074]. It is a subtype of renal cell carcinoma tending to show a tubulo-papillary architecture formed by numerous, irregular, finger-like projections of connective tissue. Renal cell carcinoma is a heterogeneous group of sporadic or hereditary carcinoma derived from cells of the proximal renal tubular epithelium. It is subclassified into common renal cell carcinoma (clear cell, non-papillary carcinoma), papillary renal cell carcinoma, chromophobe renal cell carcinoma, collecting duct carcinoma with medullary carcinoma of the kidney, and unclassified renal cell carcinoma.</p> <p>Note=A common allele in the promoter region of the MET shows genetic association with susceptibility to autism in some families. Functional assays indicate a decrease in MET promoter activity and altered binding of specific transcription factor complexes.</p> <p>Note=MET activating mutations may be involved in the development of a highly malignant, metastatic syndrome known as cancer of unknown primary origin (CUP) or primary occult malignancy. Systemic neoplastic spread is generally a late event in cancer progression. However, in some instances, distant dissemination arises at a very early stage, so that metastases reach clinical relevance before primary lesions. Sometimes, the primary lesions cannot be identified in spite of the progresses in the diagnosis of malignancies.</p>
配列類似性	<p>Belongs to the protein kinase superfamily. Tyr protein kinase family.</p> <p>Contains 3 IPT/TIG domains.</p> <p>Contains 1 protein kinase domain.</p> <p>Contains 1 Sema domain.</p>
ドメイン	The kinase domain is involved in SPSB1 binding.
翻訳後修飾	Dephosphorylated by PTPRJ at Tyr-1349 and Tyr-1365.
細胞内局在	Membrane.

画像



Western blot - Anti-Met (c-Met) antibody [4AT44] (ab59884)

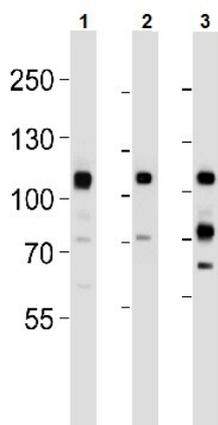
Lane 1: Wild type HAP1 whole cell lysate (20 µg)

Lane 2: Empty knockout HAP1 whole cell lysate (0 µg)

Lane 3: MET whole cell lysate (20 µg)

Lanes 1 - 3: Merged signal (red and green). Green - ab59884 observed at 155 kDa. Red - loading control, **ab176560**, observed at 50 kDa.

ab59884 was shown not to specifically react with Met (c-Met) when Met (c-Met) knockout samples were used. Wild-type and Met (c-Met) knockout samples were subjected to SDS-PAGE. Ab59884 and **ab176560** (Rabbit anti alpha Tubulin loading control) were incubated overnight at 4°C at 100 dilution and 1/10000 dilution respectively. Blots were developed with Goat anti-Mouse IgG H&L (IRDye® 800CW) preabsorbed **ab216772** and Goat anti-Rabbit IgG H&L (IRDye® 680RD) preabsorbed **ab216777** secondary antibodies at 1/10000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-Met (c-Met) antibody [4AT44] (ab59884)

All lanes : Anti-Met (c-Met) antibody [4AT44] (ab59884) at 1/1000 dilution

Lane 1 : HeLa Cell Lysate

Lane 2 : HepG2 Cell Lysate

Lane 3 : Cos-7 Cell Lysate

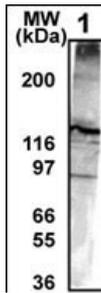
Lysates/proteins at 35 µg per lane.

Secondary

All lanes : Goat anti-mouse IgG H+L HRP conjugated at 1/3000 dilution

Predicted band size: 156 kDa

Incubation time was overnight at 4°C. Blocking/Dilution buffer: 5% NFDM/TBST.

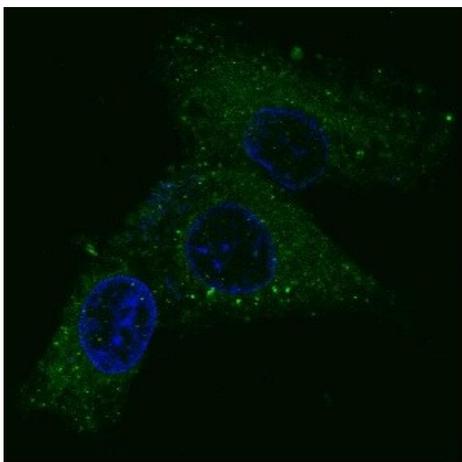


Western blot - Anti-Met (c-Met) antibody [4AT44] (ab59884)

Anti-Met (c-Met) antibody [4AT44] (ab59884) at 1/100 dilution + HepG2 cell lysate at 10 µg

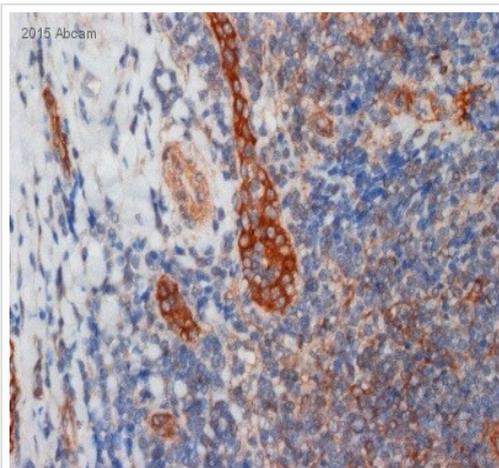
Predicted band size: 156 kDa

Observed band size: 135 kDa



Immunocytochemistry/ Immunofluorescence - Anti-Met (c-Met) antibody [4AT44] (ab59884)

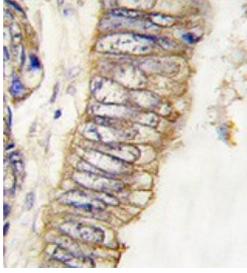
Fluorescent confocal image of HepG2 cells stained with ab59884 antibody. HepG2 cells were fixed with 4% PFA (20 min), permeabilized with Triton X-100 (0.2%, 30 min). Cells were then incubated with ab59884 primary antibody (1:100, 2 h at room temperature). For secondary antibody, Alexa Fluor® 488 conjugated donkey anti-mouse antibody (**ab150105**) (green) was used (1:1000, 1h). Nuclei were counterstained with Hoechst 33342 (blue) (10 µg/ml, 5 min).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Met (c-Met) antibody [4AT44] (ab59884)

This image is courtesy of an anonymous Abreview

ab59884 staining Met (c-Met) in human tonsil tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections). Tissue was fixed with formaldehyde; antigen retrieval was by heat mediation. Samples were incubated with the undiluted primary antibody. An undiluted biotin-conjugated goat anti-mouse IgG polyclonal was used as the secondary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Met (c-Met) antibody [4AT44] (ab59884)

Immunohistochemical staining of Met (c-Met) in Human colon carcinoma tissue sections (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections) with ab59884 at a dilution of 1/25. Tissue was fixed with formaldehyde and blocked with 3% BSA for 0.5 hours at 38°C. Antigen retrieval was heat mediation with a citrate buffer (pH6). Samples were incubated with primary antibody (1/25) for 1 hours at 37°C. A HRP-conjugated goat anti-mouse polyclonal (ready to use) was used as the secondary antibody.

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