abcam

Product datasheet

Anti-MTR4 antibody ab70551

★★★★★ 3 Abreviews 11 References 4 图像

概述

产品名称 Anti-MTR4抗体

描述 兔多克隆抗体to MTR4

宿主 Rabbit

经测试应用 适用于: IP, IHC-P, ICC, WB

种属反应性 与反应: Mouse, Human

免疫原 Synthetic peptide corresponding to Human MTR4.

Database link: NP 056175.2

常规说明

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

性能

形式 Liquid

存放说明 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

存储溶液 pH: 6.8

Preservative: 0.09% Sodium azide

Constituents: 0.1% BSA, Tris buffered saline

纯**度** Immunogen affinity purified

应用

The Abpromise guarantee Abpromise™承诺保证使用ab70551于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

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应用	Ab评论	说明
IP		Use at an assay dependent concentration.
IHC-P		Use a concentration of 5 µg/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
ICC		Use at an assay dependent concentration. Epitope retrieval with Tris-EDTA pH9.0 is recommended for formalin-fixed paraffin-embedded cells.
WB	★★★★☆ (3)	1/2000 - 1/10000. Detects a band of approximately 118 kDa (predicted molecular weight: 118 kDa).

靶标

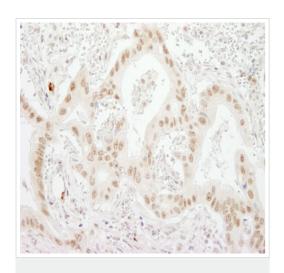
相关性 MTR4, or superkiller viralicidic activity 2 like 2, may be involved in pre-mRNA splicing. MTR4 is an

exosome-associated protein and may be involved in the recruitment of the exosome to pre-rRNA,

to mediate the 3' end processing of 5.8S rRNA.

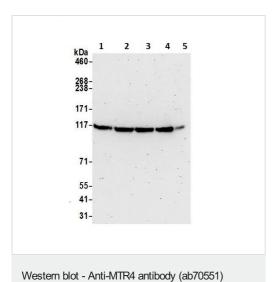
细胞定位 Nucleus, nucleolus.

图片



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-MTR4 antibody (ab70551)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human lung carcinoma tissue labelling MTR4 with ab70551 at 1/200 (1 μ g/ml). Detection: DAB.



All lanes: Anti-MTR4 antibody (ab70551) at 0.1 µg/ml

Lane 1 : HeLa (human epithelial cell line from cervix adenocarcinoma) whole cell lysate

Lane 2: HEK-293T (human epithelial cell line from embryonic kidney transformed with large T antigen) whole cell lysate

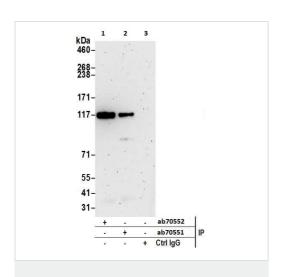
Lane 3: Jurkat (human T cell leukemia cell line from peripheral blood) whole cell lysate

Lane 4 : TCMK-1 (mouse kidney epithelial cell line) whole cell lysate

Lane 5 : NIH/3T3 (mouse embryo fibroblast cell line) whole cell lysate

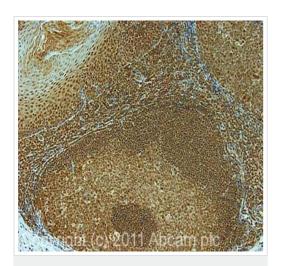
Predicted band size: 118 kDa

Detection: Chemiluminescence with an exposure time of 3 minutes.



Immunoprecipitation - Anti-MTR4 antibody (ab70551)

Detection of human MTR4 by western blot of immunoprecipitates. Samples: Whole cell lysate (1.0 mg per IP reaction; 20% of IP loaded) from HeLa cells prepared using NETN lysis buffer. Antibodies: Affinity purified rabbit anti-MTR4 antibody ab70551 used for IP at 3 μ g per reaction. MTR4 was also immunoprecipitated by rabbit anti-MTR4 antibody ab70552. For blotting immunoprecipitated MTR4, ab70551 was used at 1 μ g/ml. Detection: Chemiluminescence with an exposure time of 3 minutes.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-MTR4 antibody (ab70551)

IHC image of ab70551 staining in human normal tonsil formalin fixed paraffin embedded tissue section, performed on a Leica BondTM system using the standard protocol F. The section was pretreated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab70551, 5µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

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