abcam

Product datasheet

Anti-UFM1 antibody [EPR4264(2)] ab109305



重组 RabMAb

★★★★ 4 Abreviews 29 References 9图像

概述

产品名称 Anti-UFM1抗体[EPR4264(2)]

描述 兔单克隆抗体[EPR4264(2)] to UFM1

宿主 Rabbit

经测试应用 适用于: WB, IP, IHC-P

不适用于: Flow Cyt or ICC/IF

种属反应性 与反应: Mouse, Rat, Human

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

阳性对照 Fetal kidney, 293T, SH-SY5Y, and HepG2 whole cell lysate (ab7900); Human brain tissue

常规说明 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**.

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

存储溶液 pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), 59% PBS, 0.05% BSA

纯度 Protein A purified

克隆 单克隆

克隆编号 EPR4264(2)

同种型 ΙgG

The Abpromise guarantee

Abpromise™承诺保证使用ab109305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。

应用	Ab评论	说明
WB	★★★★★ (3)	1/1000 - 1/2000. Predicted molecular weight: 9 kDa.
IP		1/10 - 1/40.
IHC-P		1/250 - 1/1000. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

应用说明

Is unsuitable for Flow Cyt or ICC/IF.

靶标

功能

Ubiquitin-like modifier protein which binds to a number of target proteins, such as DDRGK1.

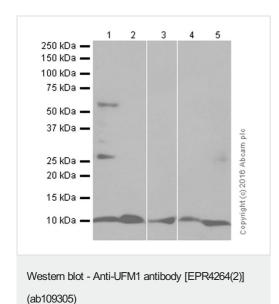
序列相似性

Belongs to the UFM1 family.

细胞定位

Nucleus. Cytoplasm. Predominantly nuclear. Also expressed diffusely in the cytoplasm.

图片



All lanes : Anti-UFM1 antibody [EPR4264(2)] (ab109305) at 1/2000 dilution

Lane 1: SH-SY5Y (Human neuroblastoma epithelial cell) whole cell

lysate

Lane 2: HepG2 (Human hepatocellular carcinoma epithelial cell)

whole cell lysate

Lane 3: Mouse spleen cell lysate

Lane 4: Mouse heart cell lysate

Lane 5: Rat spleen cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000

dilution

Performed under reducing conditions.

Predicted band size: 9 kDa **Observed band size:** 28,9 kDa Mouse heart cell lysate Blocking and diluting buffer: 5% NFDM/TBST

The band at around 28kDa might be UFM1 conjugates [PMID: 23152784].

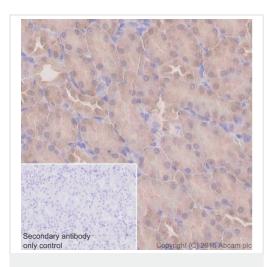
Immunoprecipitation - Anti-UFM1 antibody [EPR4264(2)] (ab109305)

ab109305 immunoprecipitating UFM1. 10µg of cell lysate was incubated with primary antibody at a dilution of 1/20 and VeriBlot for IP Detection Reagent (HRP) (ab131366) at a dilution of 1/1000.

Lane 1: 293 (Human embryonic kidney epithelial cell) whole cell lysate 10ug

Lane 2: 293 (Human embryonic kidney epithelial cell) whole cell lysate

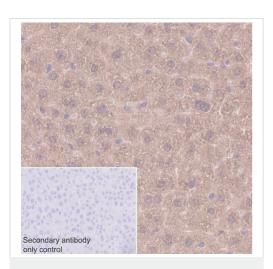
Lane 3: Rabbit monoclonal IgG (<u>ab172730</u>) instead of ab109305 in 293 (Human embryonic kidney epithelial cell) whole cell lysate



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-UFM1 antibody
[EPR4264(2)] (ab109305)

ab109305 staining UFM1 in rat kidney tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffinembedded sections). Tissue was fixed with paraformaldehyde and antigen retrieval was by heat mediation in a EDTA buffer. Samples were incubated with primary antibody at a dilution of 1/1000. A goat anti-rabbit IgG H&L (HRP) ab97051 was used as the secondary antibody at a dilution of 1/500.

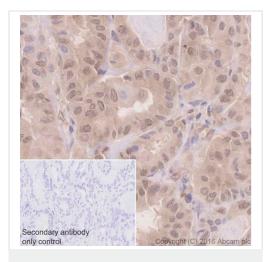
Negative control 1: PBS in place of primary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-UFM1 antibody
[EPR4264(2)] (ab109305)

ab109305 staining UFM1 in mouse mouse liver tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffinembedded sections). Tissue was fixed with paraformaldehyde and antigen retrieval was by heat mediation in a EDTA buffer. Samples were incubated with primary antibody at a dilution of 1/100. A goat anti-rabbit IgG H&L (HRP) <u>ab97051</u> was used as the secondary antibody at a dilution of 1/500.

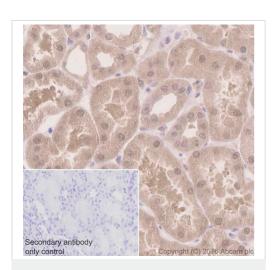
Negative control 1: PBS in place of primary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-UFM1 antibody
[EPR4264(2)] (ab109305)

ab109305 staining UFM1 in human thyroid carcinoma tissue sections by Immunohistochemistry (IHC-P - paraformaldehydefixed, paraffin-embedded sections). Tissue was fixed with paraformaldehyde and antigen retrieval was by heat mediation in a EDTA buffer. Samples were incubated with primary antibody at a dilution of 1/100. A goat anti-rabbit IgG H&L (HRP) ab97051 was used as the secondary antibody at a dilution of 1/500.

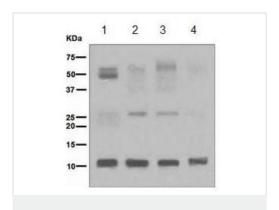
Negative control 1: PBS in place of primary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-UFM1 antibody
[EPR4264(2)] (ab109305)

ab109305 staining UFM1 in human kidney tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffinembedded sections). Tissue was fixed with paraformaldehyde and antigen retrieval was by heat mediation in a EDTA buffer. Samples were incubated with primary antibody at a dilution of 1/1000. A goat anti-rabbit IgG H&L (HRP) <u>ab97051</u> was used as the secondary antibody at a dilution of 1/500.

Negative control 1: PBS in place of primary antibody.



Western blot - Anti-UFM1 antibody [EPR4264(2)] (ab109305)

All lanes : Anti-UFM1 antibody [EPR4264(2)] (ab109305) at 1/1000 dilution

Lane 1: Fetal kidney lysate

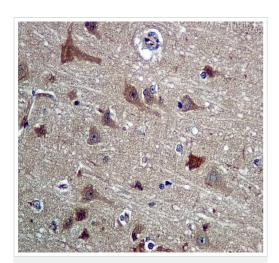
Lane 2: 293T cell lysate

Lane 3: SH-SY5Y cell lysate

Lane 4: HepG2 cell lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 9 kDa **Observed band size:** 9 kDa

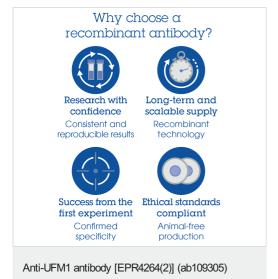


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-UFM1 antibody

[EPR4264(2)] (ab109305)

Immunohistochemical analysis of paraffin-embedded Human brain tissue using ab109305 at 1/250 dilution.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



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