abcam

Product datasheet

Anti-Hsc70 antibody [EP1531Y] ab51052





重组 RabMAb

★★★★★ 2 Abreviews 41 References 8 图像

概述

产品名称 Anti-Hsc70抗体[EP1531Y]

描述 兔单克隆抗体[EP1531Y] to Hsc70

宿主 Rabbit

经测试应用 适用于: IP, Flow Cyt (Intra), WB, IHC-P, ICC/IF

种属反应性 与反应: Mouse, Rat, Human

免疫原 Synthetic peptide within Human Hsc70 aa 600 to the C-terminus (C terminal). The exact

sequence is proprietary.

阳性对照 WB: A431, MCF7, and HeLa cell lysates. IHC-P: Human cerebral cortex tissue. ICC/IF: MCF7

cells. Flow Cyt (intra): MCF7 cells. IP: HeLa cell lysate.

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

Avoid freeze / thaw cycle.

存储溶液 pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 49% PBS, 50% Glycerol (glycerin, glycerine), 0.05% BSA

纯度 Protein A purified

克隆 单克隆 克隆编号 EP1531Y

同种型 IgG

应用

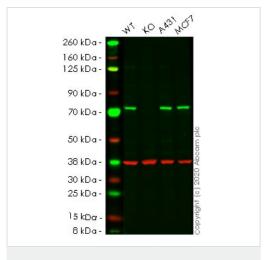
The Abpromise guarantee Abpromise™承诺保证使用ab51052于以下的经测试应用

"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
IP	★★★ ☆☆ <u>(1)</u>	1/10 - 1/20.
Flow Cyt (Intra)		1/30 - 1/70. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
WB		1/500 - 1/5000. Detects a band of approximately 71 kDa (predicted molecular weight: 71 kDa).
IHC-P		1/50. Perform heat mediated antigen retrieval before commencing with IHC staining protocol. See IHC antigen retrieval protocols.
ICC/IF	★★★★☆ (1)	1/100 - 1/250.

靶标	
功能	Acts as a repressor of transcriptional activation. Inhibits the transcriptional coactivator activity of CITED1 on Smad-mediated transcription. Chaperone. Isoform 2 may function as an endogenous inhibitory regulator of HSC70 by competing the co-chaperones.
组织 特异性	Ubiquitous.
序列相似性	Belongs to the heat shock protein 70 family.
结 构域	The N-terminal 1-386 residues constitute the ATPase domain, while residues 387-646 form the peptide-binding domain.
翻译后修饰	Phosphorylated upon DNA damage, probably by ATM or ATR. ISGylated.
细胞定位	Cytoplasm. Melanosome. Localized in cytoplasmic mRNP granules containing untranslated mRNAs. Translocates rapidly from the cytoplasm to the nuclei, and especially to the nucleoli, upon heat shock. Identified by mass spectrometry in melanosome fractions from stage I to stage IV.

图片



Western blot - Anti-Hsc70 antibody [EP1531Y] (ab51052)

All lanes : Anti-Hsc70 antibody [EP1531Y] (ab51052) at 1/500 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: HSPA8 knockout HeLa cell lysate

Lane 3 : A431 cell lysate

Lane 4 : MCF7 cell lysate

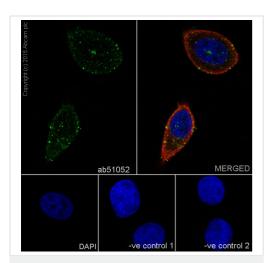
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 71 kDa **Observed band size:** 71 kDa

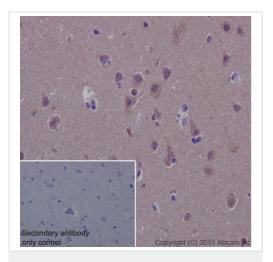
Lanes 1-4: Merged signal (red and green). Green - ab51052 observed at 71 kDa. Red - Anti-GAPDH antibody [6C5] - Loading Control (ab8245) observed at 37 kDa.

ab51052 was shown to react with Hsc70 in wild-type HeLa cells in western blot. Loss of signal was observed when knockout cell line ab265664 (knockout cell lysate ab256944) was used. Wild-type HeLa and HSPA8 knockout HeLa cell lysates were subjected to SDS-PAGE. Membrane was blocked for 1 hour at room temperature in 0.1% TBST with 3% non-fat dried milk. ab51052 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) overnight at 4°C at a 1 in 500 dilution and a 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.



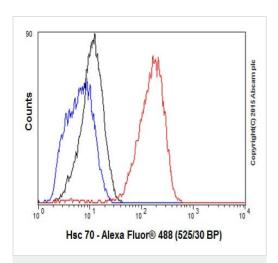
Immunocytochemistry/ Immunofluorescence - Anti-Hsc70 antibody [EP1531Y] (ab51052)

Immunofluorescence staining of MCF7 cells with purified ab51052 at a working dilution of 1/250, counter-stained with DAPI. The secondary antibody was Alexa Fluor[®] 488 goat anti-rabbit (ab150077), used at a dilution of 1/1000. ab7291, a mouse antitubulin antibody (1/1000), was used to stain tubulin along with ab150120 (Alexa Fluor[®] 594 goat anti-mouse, 1/1000), shown in the top right hand panel. The cells were fixed in 4% PFA and permeabilized using 0.1% Triton X 100. The negative controls are shown in bottom middle and right hand panels - for negative control 1, purified ab51052 was used at a dilution of 1/500 followed by an Alexa Fluor[®] 594 goat anti-mouse antibody (ab150120) at a dilution of 1/500. For negative control 2, ab7291 (mouse antitubulin) was used at a dilution of 1/500 followed by an Alexa Fluor[®] 488 goat anti-rabbit antibody (ab150077) at a dilution of 1/400.



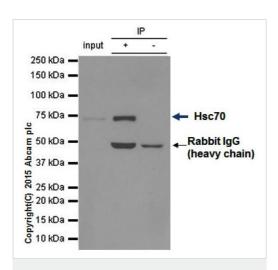
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Hsc70 antibody
[EP1531Y] (ab51052)

Immunohistochemical staining of paraffin embedded human cerebral cortex with purified ab51052 at a working dilution of 1/500. The secondary antibody used is HRP goat anti-rabbit IgG H&L (ab97051) at 1/500. The sample is counter-stained with hematoxylin. Antigen retrieval was perfomed using Tris-EDTA buffer, pH 9.0. PBS was used instead of the primary antibody as the negative control, and is shown in the inset.



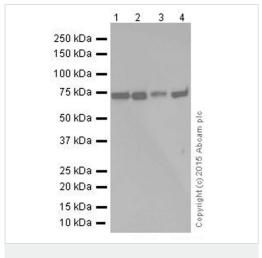
Flow Cytometry (Intracellular) - Anti-Hsc70 antibody [EP1531Y] (ab51052)

Overlay histogram showing MCF7 cells fixed in 4% PFA and stained with purified ab51052 at a dilution of 1 in 70 (red line). The secondary antibody used was FITC goat anti-rabbit at a dilution of 1 in 500. Rabbit monoclonal IgG was used as an isotype control (black line) and cells incubated in the absence of both primary and secondary antibody were used as a negative control (blue line).



Immunoprecipitation - Anti-Hsc70 antibody [EP1531Y] (ab51052)

ab51052 (purified) at 1/20 immunoprecipitating Hsc70 in 10 µg HeLa (Lanes 1 and 2, observed at 71 kDa). Lane 3 - PBS. For western blotting, HRP Veriblot for IP Detection Reagent (ab131366) was used for detection (1/10 000). Blocking buffer and concentration: 5% NFDM/TBST Dilution buffer and concentration: 5% NFDM/TBST



Western blot - Anti-Hsc70 antibody [EP1531Y] (ab51052)

All lanes : Anti-Hsc70 antibody [EP1531Y] (ab51052) at 1/5000 dilution (purified)

Lane 1: C6 cell lysate

Lane 2: PC-12 cell lysate

Lane 3 : Mouse heart tissue lysate

Lane 4 : Rat heart tissue lysate

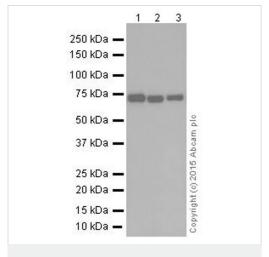
Lysates/proteins at 10 µg per lane.

Secondary

All lanes: HRP goat anti-rabbit lgG (H+L) at 1/20000 dilution

Predicted band size: 71 kDa **Observed band size:** 71 kDa

Blocking and dilution buffer: 5% NFDM/TBST.



Western blot - Anti-Hsc70 antibody [EP1531Y] (ab51052)

All lanes : Anti-Hsc70 antibody [EP1531Y] (ab51052) at 1/5000 dilution (purified)

Lane 1: human fetal brain lysate

Lane 2 : HeLa cell lysate
Lane 3 : HEK293 cell lysate

Lysates/proteins at 10 µg per lane.

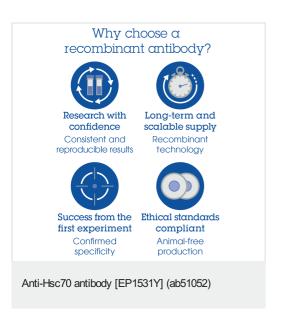
Secondary

All lanes: HRP goat anti-rabbit lgG (H+L) at 1/20000 dilution

Predicted band size: 71 kDa **Observed band size:** 71 kDa

Blocking buffer: 5% NFDM/TBST

Dilution buffer: 5% NFDM/TBST



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