abcam

Product datasheet

Anti-CRABP2 antibody [EPR17376] ab211927



重组 RabMAb

★★★★★ 1 Abreviews 4 References 11 图像

概述

产品名称 Anti-CRABP2抗体[EPR17376]

描述 兔单克隆抗体[EPR17376] to CRABP2

宿主 Rabbit

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

种属反应性 与反应: Mouse, Rat, Human

免疫原 Recombinant full length protein. This information is proprietary to Abcam and/or its suppliers.

阳性对照 WB: Human, mouse and rat skin lysates; HT-29 and MCF7 whole cell lysates. IHC-P: Human

oesophagus, skin and pancreatic ductal adenocarcinoma tissues; mouse and rat skin tissues.

ICC/IF: MCF7 and HT-29 cells. Flow Cyt (intra): MCF7 cells.

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 long

term. Avoid freeze / thaw cycle.

存储溶液 pH: 7.2

Preservative: 0.01% Sodium azide

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

纯度 Protein A purified

克隆 单克隆 克隆编号 EPR17376

同种型 lgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab211927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。

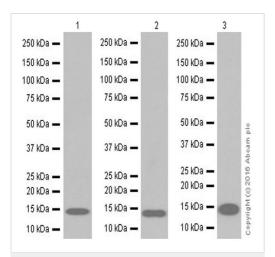
应用	Ab评论	说明
ICC/IF		1/250.
IHC-P	★★★★☆ (1)	1/1000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		1/1000. Detects a band of approximately 14 kDa (predicted molecular weight: 16 kDa).
Flow Cyt (Intra)		1/600. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.

靶标		
功能	Transports retinoic acid to the nucleus. Regulates the access of retinoic acid to the nuclear retinoic acid receptors.	
序列相似性	Belongs to the calycin superfamily. Fatty-acid binding protein (FABP) family.	
结 构域	Forms a beta-barrel structure that accommodates hydrophobic ligands in its interior.	

细胞定位 Cytoplasm. Nucleus. Upon ligand binding, a conformation change exposes a nuclear localization

motif and the protein is transported into the nucleus.

图片



Western blot - Anti-CRABP2 antibody [EPR17376] (ab211927)

Lanes 1 & 3: Anti-CRABP2 antibody [EPR17376] (ab211927) at 1/5000 dilution

Lane 2: Anti-CRABP2 antibody [EPR17376] (ab211927) at 1/1000 dilution

Lane 1: Human skin lysate

Lane 2 : MCF7 (Human breast adenocarcinoma cell line) whole cell lysate

Lane 3: HT-29 (Human colorectal adenocarcinoma cell line) whole cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

Lane 1 : Goat Anti-Rabbit IgG Peroxidase Conjugate, specific to the non-reduced form of IgG at 1/10000 dilution

Lane 2 : Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Lane 3 : Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/10000 dilution

Predicted band size: 16 kDa
Observed band size: 14 kDa

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

ab211927 MERGED

Immunocytochemistry/ Immunofluorescence - Anti-CRABP2 antibody [EPR17376] (ab211927)

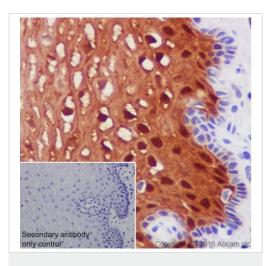
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized MCF7 (Human breast adenocarcinoma cell line) cells labeling CRABP2 with ab211927 at 1/250 dilution, followed by Goat Anti-Rabbit lgG (Alexa Fluor[®] 488) (ab150077) secondary at 1/1000 dilution (green). Confocal image showing nuclear and cytoplasmic staining on MCF7 cell line. The nuclear counter stain is DAPI (blue).

Tubulin is detected with Anti-alpha Tubulin mouse MAb (<u>ab7291</u>) at 1/1000 dilution followed by Goat Anti-Mouse IgG H&L (Alexa Fluor[®] 594) (<u>ab150120</u>) secondary antibody at 1/1000 dilution (red).

The negative controls are as follows:

-ve control 1: ab211927 at 1/250 dilution, followed by Goat Anti-Mouse lgG H&L (Alexa Fluor $^{\otimes}$ 594) (ab150120) secondary antibody at 1/1000 dilution.

-ve control 2: Anti-alpha Tubulin mouse MAb (<u>ab7291</u>) at 1/1000 dilution followed, by followed by Goat Anti-Rabbit lgG (Alexa Fluor[®] 488) (<u>ab150077</u>) secondary at 1/1000 dilution.

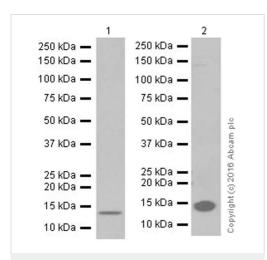


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CRABP2 antibody
[EPR17376] (ab211927)

Immunohistochemical analysis of paraffin-embedded human oesophagus tissue labeling CRABP2 with ab211927 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Cytoplasmic and nuclear staining on the stratified squamous epithelium of the human oesophagus is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Western blot - Anti-CRABP2 antibody [EPR17376] (ab211927)

All lanes : Anti-CRABP2 antibody [EPR17376] (ab211927) at 1/1000 dilution

Lane 1 : Mouse skin lysate

Lane 2 : Rat skin lysate

Lysates/proteins at 10 µg per lane.

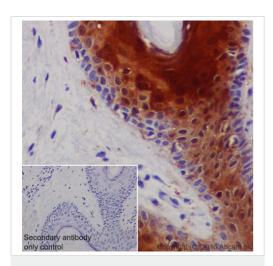
Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/100000 dilution

Predicted band size: 16 kDa **Observed band size:** 14 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

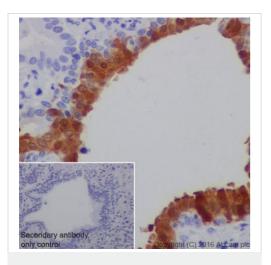


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CRABP2 antibody
[EPR17376] (ab211927)

Immunohistochemical analysis of paraffin-embedded human skin tissue labeling CRABP2 with ab211927 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Cytoplasmic and nuclear staining on the stratified squamous epithelium and hair follicle cells of the human skin is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

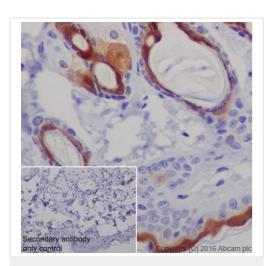


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CRABP2 antibody
[EPR17376] (ab211927)

Immunohistochemical analysis of paraffin-embedded human pancreatic ductal adenocarcinoma tissue labeling CRABP2 with ab211927 at 1/1000 dilution, followed by Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution. Cytoplasmic and nuclear staining on the human pancreatic ductal adenocarcinoma is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

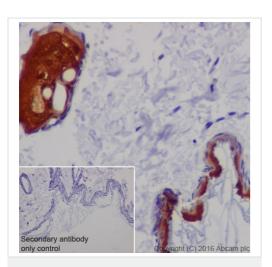


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CRABP2 antibody
[EPR17376] (ab211927)

Immunohistochemical analysis of paraffin-embedded mouse skin tissue labeling CRABP2 with ab211927 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Cytoplasmic and nuclear staining on the stratified squamous epithelium, hair follicle cells and sweat gland cells of the mouse skin is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

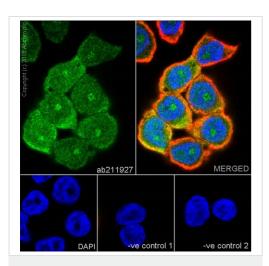


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CRABP2 antibody
[EPR17376] (ab211927)

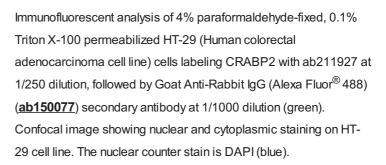
Immunohistochemical analysis of paraffin-embedded rat skin tissue labeling CRABP2 with ab211927 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Cytoplasmic and nuclear staining on the stratified squamous epithelium and sweat gland cells of the rat skin is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



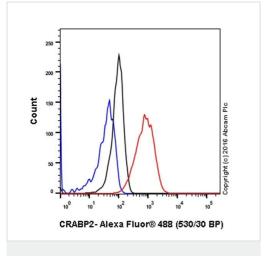
Immunocytochemistry/ Immunofluorescence - Anti-CRABP2 antibody [EPR17376] (ab211927)



Tubulin is detected with Anti-alpha Tubulin mouse MAb (ab7291) at 1/1000 dilution, followed by Goat Anti-Mouse IgG H&L (Alexa Fluor[®] 594) (ab150120) secondary antibody at 1/1000 dilution (red).

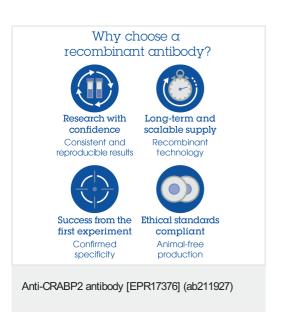
The negative controls are as follows:

- -ve control 1: ab211927 at 1/250 dilution, followed by Goat Anti-Mouse IgG H&L (Alexa Fluor[®] 594) (ab150120) secondary antibody at 1/1000 dilution.
- -ve control 2: Anti-alpha Tubulin mouse MAb (<u>ab7291</u>) at 1/1000 dilution, followed by Goat Anti-Rabbit lgG (Alexa Fluor[®] 488) (<u>ab150077</u>) secondary antibody at 1/1000 dilution.



Flow Cytometry (Intracellular) - Anti-CRABP2 antibody [EPR17376] (ab211927)

Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed MCF7 (Human breast adenocarcinoma cell line) cells labeling CRABP2 with ab211927 at 1/600 dilution (red) compared with a rabbit monoclonal IgG isotype control (ab172730; black) and an unlabeled control (cells without incubation with primary antibody and secondary antibody; blue). Goat Anti-Rabbit IgG (Alexa Fluor® 488) at 1/2000 dilution was used as the secondary antibody.



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