abcam

Product datasheet

Anti-Folate Binding Protein/FBP antibody ab67422

★★★★★ 2 Abreviews 8 References 4 图像

概述

产品名称 Anti-Folate Binding蛋白/FBP抗体

描述 兔多克隆抗体to Folate Binding蛋白/FBP

宿主 Rabbit

经测试应用 适用于: ICC/IF, WB, IHC-P

种属反应性 与反应: Mouse, Human

预测可用于: Rat, Chicken 📤

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

阳性对照 WB: Mouse cerebellum tissue lysate, JAR and HeLa cell lysates. ICC/IF: HepG2 cells.

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

80°C. Avoid freeze / thaw cycle.

存储溶液 pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising

scientific support team who will be happy to help.

纯**度** Immunogen affinity purified

克隆 多克隆

同种型 IgG

1

The Abpromise guarantee

Abpromise™承诺保证使用ab67422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		Use a concentration of 5 µg/ml.
WB		Use a concentration of 1 μ g/ml. Detects a band of approximately 28 kDa (predicted molecular weight: 30 kDa). Please note, we have been advised by some customers that ab67422 is unable to detect the human version of this protein in western blot. Please contact our scientific support services if you have any queries regarding this antibody.
IHC-P	*** <u>*</u>	Use a concentration of 10 µg/ml. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.

功能 Binds to folate and reduced folic acid derivatives and mediates delivery of 5-

methyltetrahydrofolate to the interior of cells.

组织**特异性** Exclusively expressed in tissues of epithelial origin. Expression is increased in malignant tissues.

Expressed in kidney, lung and cerebellum.

疾病相关 Defects in FOLR1 are the cause of neurodegeneration due to cerebral folate transport deficiency

(NCFTD) [MIM:613068]. NCFTD is an autosomal recessive disorder resulting from brain-specific

folate deficiency early in life. Onset is apparent in late infancy with severe developmental regression, movement disturbances, epilepsy, and leukodystrophy. Note=Recognition and diagnosis of this disorder is critical because folinic acid therapy can reverse the clinical

symptoms and improve brain abnormalities and function.

序列相似性 Belongs to the folate receptor family.

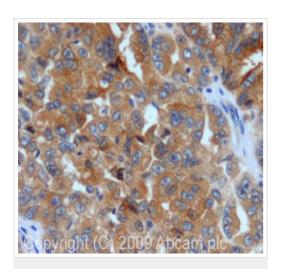
翻译**后修**饰 Eight disulfide bonds are present.

The secreted form is derived from the membrane-bound form either by cleavage of the GPI

anchor, or/and by proteolysis catalyzed by a metalloprotease.

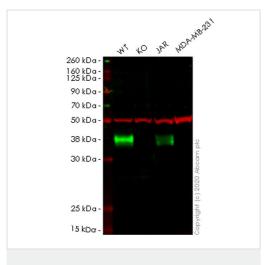
细**胞定位** Cell membrane. Secreted.

图片



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Folate Binding
Protein/FBP antibody (ab67422)

IHC image of Folate Binding Protein/FBP staining in human kidney carcinoma formalin fixed paraffin embedded tissue section, performed on a Leica BondTM system using the standard protocol F. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab67422, 10µg/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.



Western blot - Anti-Folate Binding Protein/FBP antibody (ab67422)

All lanes : Anti-Folate Binding Protein/FBP antibody (ab67422) at 1/500 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: FOLR1 knockout HeLa cell lysate

Lane 3: JAR cell lysate

Lane 4: MDA-MB-231 cell lysate

Lysates/proteins at 20 µg per lane.

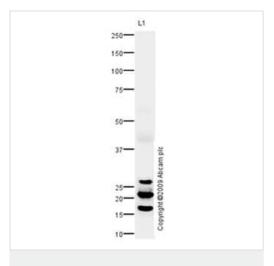
Performed under reducing conditions.

Predicted band size: 30 kDa **Observed band size:** 38 kDa

Lanes 1-4: Merged signal (red and green). Green - ab67422 observed at 38 kDa. Red - loading control, <u>ab7291</u> observed at 52 kDa.

ab67422 Anti-Folate Binding Protein/FBP antibody was shown to specifically react with Folate Binding Protein/FBP in wild-type HeLa

cells. Loss of signal was observed when knockout cell line <u>ab264921</u> (knockout cell lysate <u>ab257270</u>) was used. Wild-type and Folate Binding Protein/FBP knockout samples were subjected to SDS-PAGE. ab67422 and Anti-alpha Tubulin antibody [DM1A] - Loading Control (<u>ab7291</u>) were incubated overnight at 4°C at 1 in 500 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (<u>ab216773</u>) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (<u>ab216776</u>) secondary antibodies at 1 in 10000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-Folate Binding Protein/FBP antibody (ab67422)

Anti-Folate Binding Protein/FBP antibody (ab67422) at 1/1 dilution + Cerebellum Mouse Tissue Lysate at 10 µg

Secondary

Goat polyclonal to Rabbit lgG - H&L - Pre-Adsorbed (HRP) at 1/3000 dilution

Developed using the ECL technique.

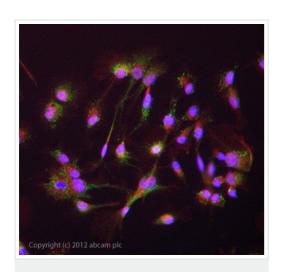
Performed under reducing conditions.

Predicted band size: 30 kDa Observed band size: 28 kDa

Additional bands at: 16 kDa (possible cleavage fragment), 22

kDa (possible cleavage fragment)

We hypothesize that the 28, 22 and 16 kDa bands represent the propertide with signal sequence, propertide without signal sequence and mature protein, respectively.



Immunocytochemistry/ Immunofluorescence - Anti-Folate Binding Protein/FBP antibody (ab67422)

ab67422 stained HepG2 cells. The cells were 4% formaldehyde fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody ab67422 at 5μ g/ml overnight at +4°C. The secondary antibody (green) was DyLight® 488 goat anti- rabbit (ab96899) lgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43μ M.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
- · Valid for 12 months from date of delivery
- · Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit https://www.abcam.cn/abpromise or contact our technical team.

Terms and conditions

· Guarantee only valid for products bought direct from Abcam or one of our authorized distributors