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Product datasheet

Anti-FHL2 antibody ab12328

1 References 画像数 5

製品の概要

製品名 Anti-FHL2 antibody

製品の詳細 Rabbit polyclonal to FHL2

由来種 Rabbit

アプリケーション 適用あり: IP, ICC/IF, IHC-P, WB

種交差性 交差種: Mouse, Human

交差が予測される動物種: Rat, Rabbit, Horse, Guinea pig, Cow, Dog, Chimpanzee, Rhesus

monkey, Gorilla, Chinese hamster, Orangutan, Elephant

免疫原 Synthetic peptide corresponding to Human FHL2.

Database link: Q14192

ポジティブ・コントロール WB: HeLa, TCMK-1 and NIH/3T3 whole cell lysates. IP: HeLa cells. IHC-P: Human breast and

ovarian carcinoma tissue.

特記事項

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

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

バッファー pH: 7

Preservative: 0.1% Sodium azide

Constituents: 0.021% PBS, 1.815% Tris, 1.764% Sodium citrate

精製度 Immunogen affinity purified

特記事項(精製) This antibody is affinity purified.

ポリ/モノ ポリクローナル

アイソタイプ IgG

1

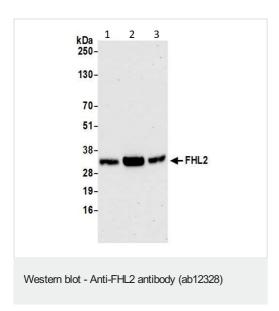
The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab12328の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
IP		Use a concentration of 1 µg/ml.
ICC/IF		Use a concentration of 1 µg/ml.
IHC-P		1/500 - 1/2000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		1/1000 - 1/10000. Detects a band of approximately 32 kDa (predicted molecular weight: 44.6 kDa).

ターゲット情報

機能	May function as a molecular transmitter linking various signaling pathways to transcriptional regulation. Negatively regulates the transcriptional repressor E4F1 and may function in cell growth.	
組織特異性	Expressed in skeletal muscle and heart.	
配列類似性	Contains 4 LIM zinc-binding domains.	
ドメイン	The third LIM zinc-binding mediates interaction with E4F1.	
細胞内局在	Cytoplasm. Nucleus.	

画像



All lanes: Anti-FHL2 antibody (ab12328) at 0.1 µg/ml

Lane 1 : HeLa (Human cervix adenocarcinoma epithelial cell)

whole cell lysate

Lane 2: TCMK-1 (Mouse kidney epithelial cell line) whole cell

lysate

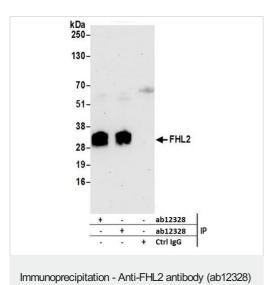
Lane 3: NIH/3T3 (Mouse embryonic fibroblast cell line) whole cell

lysate

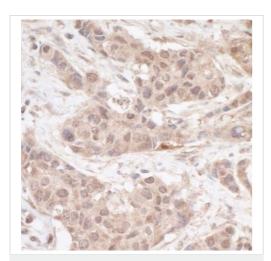
Lysates/proteins at 50 µg per lane.

Predicted band size: 44.6 kDa

Exposure time: 30 seconds



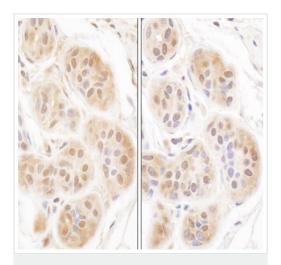
FHL2 was immunoprecipitated from HeLa (human epithelial cell line from cervix adenocarcinoma) whole cell lysate (1 mg for IP, 20% of IP loaded) with ab12328 at 3 μ g/mg lysate (different LOT numbers). Western blot was performed from the immunoprecipitate using ab12328 at 1 μ g/ml.



sections) analysis of human ovarian carcinoma tissue labelling FHL2 with ab12328 at 1/2000 (0.5 $\mu g/ml$). Detection: DAB.

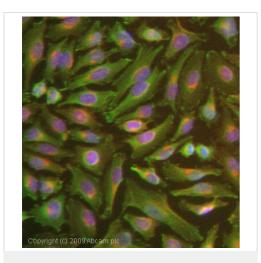
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded





Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-FHL2 antibody (ab12328)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human breast carcinoma tissue labelling FHL2 with ab12328 at 1/1000 (1 μ g/ml). Detection: DAB.



Immunocytochemistry/ Immunofluorescence - Anti-FHL2 antibody (ab12328)

ICC image of ab12328 stained HeLa cells. The cells were 4% PFA 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 (ab12328, 1 μ g/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit 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 1.43 μ M.

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