abcam

Product datasheet

Anti-Fibronectin antibody [F1] - BSA and Azide free ab271831

יעלאעבע RabMAb

★★★★★ 1 Abreviews 画像数 4

製品の概要

特記事項

製品名 Anti-Fibronectin antibody [F1] - BSA and Azide free

製品の詳細 Rabbit monoclonal [F1] to Fibronectin - BSA and Azide free

由来種 Rahhit

アプリケーション 適用あり: Flow Cyt (Intra), WB, IHC-P, IHC-Fr, ICC/IF

種交差性 交差種: Human

免疫原 Recombinant full length protein corresponding to Fibronectin aa 1-2400.

Database link: P02751

ポジティブ・コントロール IHC-P: Human breast carcinoma tissue. ICC/IF: HepG2. Flow Cyt (intra): HepG2 cells.

ab271831 is the carrier-free version of ab32419.

Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.

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**.

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with these species. Please contact us for more information.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C. Do Not Freeze.

バッファー pH: 7.20

Constituent: PBS

キャリア・フリー はい

精製度 Protein A purified

ポリ/モノ モノクローナル

クローン名 F1 **Pイソタイプ** IgG

アプリケーション

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ab271831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 |
|------------------|-----------------|---|
| Flow Cyt (Intra) | | Use at an assay dependent concentration. |
| WB | | Use at an assay dependent concentration. Predicted molecular weight: 262 kDa. |
| IHC-P | **** <u>(1)</u> | Use at an assay dependent concentration. |
| IHC-Fr | | Use at an assay dependent concentration. |
| ICC/IF | | Use at an assay dependent concentration. |

ターゲット情報

機能 Fibronectins bind cell surfaces and various compounds including collagen, fibrin, heparin, DNA,

and actin. Fibronectins are involved in cell adhesion, cell motility, opsonization, wound healing, and maintenance of cell shape. Involved in osteoblast compaction through the fibronectin fibrillogenesis cell-mediated matrix assembly process, essential for osteoblast mineralization.

Participates in the regulation of type I collagen deposition by osteoblasts.

Anastellin binds fibronectin and induces fibril formation. This fibronectin polymer, named superfibronectin, exhibits enhanced adhesive properties. Both anastellin and superfibronectin inhibit tumor growth, angiogenesis and metastasis. Anastellin activates p38 MAPK and inhibits

lysophospholipid signaling.

組織特異性 Plasma FN (soluble dimeric form) is secreted by hepatocytes. Cellular FN (dimeric or cross-

linked multimeric forms), made by fibroblasts, epithelial and other cell types, is deposited as fibrils

in the extracellular matrix. Ugl-Y1, Ugl-Y2 and Ugl-Y3 are found in urine.

関連疾患 Glomerulopathy with fibronectin deposits 2

配列類似性 Contains 12 fibronectin type-I domains.

Contains 2 fibronectin type-II domains.

Contains 16 fibronectin type-III domains.

発生段階 Ugl-Y1, Ugl-Y2 and Ugl-Y3 are present in the urine from 0 to 17 years of age.

翻訳後修飾 Sulfated.

It is not known whether both or only one of Thr-2064 and Thr-2065 are/is glycosylated.

Forms covalent cross-links mediated by a transglutaminase, such as F13A or TGM2, between a

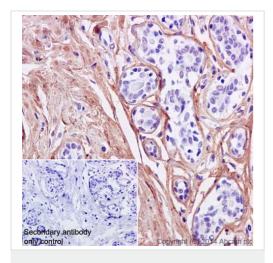
glutamine and the epsilon-amino group of a lysine residue, forming homopolymers and heteropolymers (e.g. fibrinogen-fibronectin, collagen-fibronectin heteropolymers).

Phosphorylated by FAM20C in the extracellular medium.

Proteolytic processing produces the C-terminal NC1 peptide, anastellin.

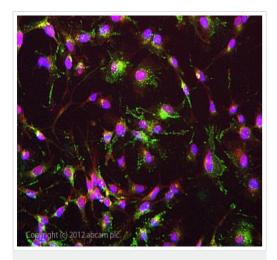
細胞内局在 Secreted, extracellular space, extracellular matrix.

画像

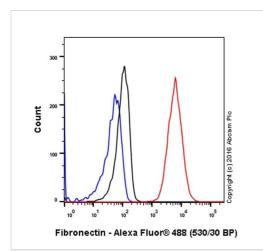


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Fibronectin antibody [F1] - BSA and Azide free (ab271831)

Immunohistochemical staining of paraffin embedded human breast carcinoma with purified <u>ab32419</u> at a dilution of 1/250. The secondary antibody used is <u>ab97051</u>, a HRP-conjugated goat antirabbit lgG (H+L), at a dilution of 1/500. The sample is counterstained with hematoxylin. Antigen retrieval was performed 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. This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab32419</u>).



Immunocytochemistry/ Immunofluorescence - Anti-Fibronectin antibody [F1] - BSA and Azide free (ab271831)



Flow Cytometry (Intracellular) - Anti-Fibronectin antibody [F1] - BSA and Azide free (ab271831)

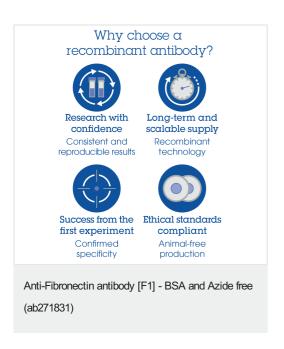
ICC/IF image of unpurified <u>ab32419</u> stained HepG2 (Human liver hepatocellular carcinoma cell line) cells. The cells were fixed in 4% formaldehyde (10 minutes) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1 hour to permeabilize the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody <u>ab32419</u> at 1/100 dilution overnight at +4°C. The secondary antibody (green) was DyLight[®] 488 goat anti- rabbit (<u>ab96899</u>) lgG (H+L) used at a 1/1000 dilution for 1 hour. Alexa Fluor[®] 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1 hour. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43 μ M.

Alexa Fluor[®] 488 (<u>ab198933</u>) and Alexa Fluor[®] 647 (<u>ab198934</u>) conjugated versions are available for this clone.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab32419).

Intracellular Flow Cytometry analysis of HepG2 (Human liver hepatocellular carcinoma cell line) cells labeling Fibronectin with purified ab32419 at 1/20 dilution (10 µg/mL) (red). Cells were fixed with 4% paraformaldehyde and permeabilized with 90% methanol. A Goat anti rabbit lgG (Alexa Fluor[®]488) at 1/2000 dilution was used as the secondary antibody. Rabbit monoclonal lgG (Black) was used as the isotype control, cells without incubation with primary antibody and secondary antibody (Blue) was used as the unlabeled control.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab32419).



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