# abcam

## Product datasheet

# Anti-CD44 antibody [F10-44-2] ab6124

★★★★★ 5 Abreviews 65 References 画像数 4

#### 製品の概要

製品名 Anti-CD44 antibody [F10-44-2]

製品の詳細 Mouse monoclonal [F10-44-2] to CD44

由来種 Mouse

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

種交差性 交差種: Human

免疫原 Tissue, cells or virus. This information is proprietary to Abcam and/or its suppliers.

ポジティブ・コントロール In Flow Cytometry, this antibody gave a positive signal in peripheral blood lymphocytes. In IHC,

this antibody gave a positive signal in human kidney carcinoma sections. ICC/IF: A431 cell line.

特記事項

This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or

conjugation for your experiments, please contact orders@abcam.com.

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

80°C. Avoid freeze / thaw cycle.

**バッファー** pH: 7.40

Preservative: 0.02% Sodium azide Constituents: PBS, 6.97% L-Arginine

精製度 Protein G purified

**ポリ/モノ** モノクローナル

**クローン名** F10-44-2

アイソタイプ lgG2a

# **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ab6124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		Use $0.5$ -1 $\mu$ g for $10^6$ cells. <u>ab170191</u> - Mouse monoclonal lgG2a, is suitable for use as an isotype control with this antibody.
ICC/IF		Use a concentration of 5 µg/ml.
IHC-P	****(3)	Use a concentration of 5 $\mu$ g/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

### ターゲット情報

機能	Receptor for hyaluronic acid (HA). Mediates cell-cell and cell-matrix in	nteractions through its affinity
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for HA, and possibly also through its affinity for other ligands such as osteopontin, collagens, and matrix metalloproteinases (MMPs). Adhesion with HA plays an important role in cell migration, tumor growth and progression. Also involved in lymphocyte activation, recirculation and homing, and in hematopoiesis. Altered expression or dysfunction causes numerous pathogenic phenotypes. Great protein heterogeneity due to numerous alternative splicing and post-

translational modification events.

組織特異性 Isoform 10 (epithelial isoform) is expressed by cells of epithelium and highly expressed by

carcinomas. Expression is repressed in neuroblastoma cells.

配列類似性 Contains 1 Link domain.

ドメイン The lectin-like LINK domain is responsible for hyaluronan binding.

翻訳後修飾 Proteolytically cleaved in the extracellular matrix by specific proteinases (possibly MMPs) in

several cell lines and tumors.

N-glycosylated.

O-glycosylated; contains more-or-less-sulfated chondroitin sulfate glycans, whose number may

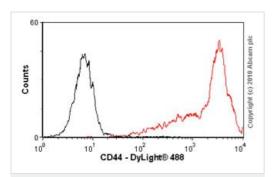
affect the accessibility of specific proteinases to their cleavage site(s).

Phosphorylated; activation of PKC results in the dephosphorylation of Ser-706 (constitutive

phosphorylation site), and the phosphorylation of Ser-672.

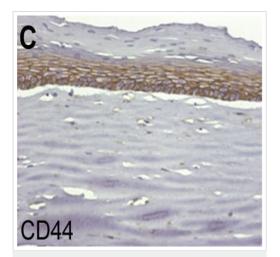
細胞内局在 Membrane.

#### 画像



Flow Cytometry - Anti-CD44 antibody [F10-44-2] (ab6124)

Overlay histogram showing peripheral blood lymphocytes stained with ab6124 (red line). The cells were incubated with the antibody (ab6124, 0.5µg/1x10<sup>6</sup> cells) for 30 min at 4°C. The secondary antibody used was DyLight® 488 goat anti-mouse lgG (H&L) (ab96879) at 1/200 dilution for 30 min at 4°C. Isotype control antibody (black line) was mouse lgG2a [ICIGG2A] (ab91361, 2µg/1x10<sup>6</sup> cells) used under the same conditions. Acquisition of >5,000 events was performed gating on peripheral blood lymphocytes.

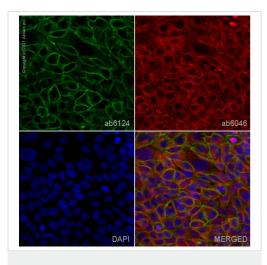


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD44 antibody [F10-44-2] (ab6124)

Image from Le Bras GF et al., PLoS One. 2011;6(11):e27063. Epub 2011 Nov 1. Fig 3.; doi:10.1371/journal.pone.0027063; November 1, 2011, PLoS ONE 6(11): e27063.

Immunohistochemical analysis of Human esophogeal keratinocytes, staining CD44 with ab6124.

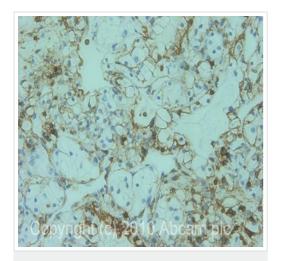
Keratinocytes were cultured *in vivo* to form an epithelium for 15 days before fixing in formaldehyde and embedding in paraffin. Primary antibody was incubated overnight at 4°C and secondary antibody for 30 minutes at 37°C. Staining was detected using DAB.



Immunocytochemistry/ Immunofluorescence - Anti-CD44 antibody [F10-44-2] (ab6124)

ab6124 staining CD44 in A431 cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% PBS-Tween for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at 4°C with ab6124 at 1µg/ml and ab6046, Rabbit polyclonal to beta Tubulin - Loading Control. Cells were then incubated with ab150117, Goat polyclonal Secondary Antibody to Mouse IgG H&L (Alexa Fluor<sup>®</sup> 488) preadsorbed at 1/1000 dilution (shown in green) and ab150080, Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor<sup>®</sup> 594) at 1/1000 dilution (shown in pseudocolour red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was acquired with a high-content analyser (Operetta CLS, Perkin Elmer) and a maximum intensity projection of confocal sections is shown.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD44 antibody [F10-44-2] (ab6124)

IHC image of CD44 staining in human kidney carcinoma formalin fixed paraffin embedded tissue section, performed on a Leica Bond TM system using the standard protocol F. The section was pretreated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab6124, 5µg/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.

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