abcam

Product datasheet

FITC Anti-Cytokeratin 18 antibody [DC-10] ab72813

3 References 3 Images

Overview

Product name FITC Anti-Cytokeratin 18 antibody [DC-10]

Description FITC Mouse monoclonal [DC-10] to Cytokeratin 18

Host species Mouse

Conjugation FITC. Ex: 493nm, Em: 528nm

Tested applications Suitable for: ICC/IF, Flow Cyt (Intra)

Species reactivity Reacts with: Human

Does not react with: Mouse, Rat, Sheep, Hamster, Cow, Dog, Pig

Immunogen Tissue, cells or virus corresponding to Human Cytokeratin 18. Human breast carcinoma cell line

PMC-42

Positive control ICC/IF: HCT116 cells. Flow Cyt (Intra): HeLa cells.

General notesThe 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

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C.

Storage buffer pH: 7.4

Preservative: 0.097% Sodium azide

Constituent: PBS

Purity Size exclusion

Purification notes The purified antibody is conjugated with Fluorescein isothiocyanate (FITC) under optimum

conditions. The reagent is free of unconjugated FITC.

Clonality Monoclonal

Clone number DC-10

1

Isotype lgG1

Applications

The Abpromise guarantee

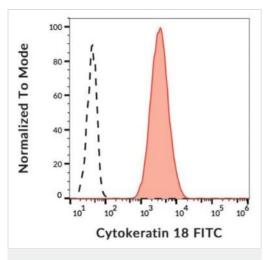
Our <u>Abpromise guarantee</u> covers the use of ab72813 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
ICC/IF		Use a concentration of 1 µg/ml.
Flow Cyt (Intra)		Use a concentration of 1 - 5 μg/ml.

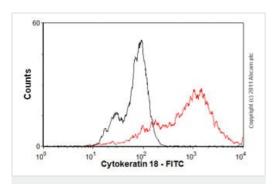
Target		
Function	Involved in the uptake of thrombin-antithrombin complexes by hepatic cells (By similarity). When phosphorylated, plays a role in filament reorganization. Involved in the delivery of mutated CFTR to the plasma membrane. Together with KRT8, is involved in interleukin-6 (IL-6)-mediated barrier protection.	
Tissue specificity	Expressed in colon, placenta, liver and very weakly in exocervix. Increased expression observed in lymph nodes of breast carcinoma.	
Involvement in disease	Defects in KRT18 are a cause of cirrhosis (CIRRH) [MIM:215600].	
Sequence similarities	Belongs to the intermediate filament family.	
Post-translational modifications	Phosphorylation at Ser-34 increases during mitosis. Hyperphosphorylated at Ser-53 in diseased cirrhosis liver. Phosphorylation increases by IL-6. Proteolytically cleaved by caspases during epithelial cell apoptosis. Cleavage occurs at Asp-238 by either caspase-3, caspase-6 or caspase-7. O-glycosylated at multiple sites; glycans consist of single N-acetylglucosamine residues.	
Cellular localization	Cytoplasm > perinuclear region.	

Images



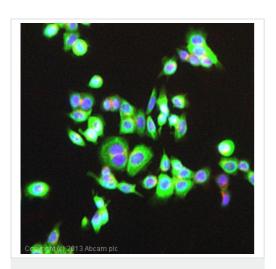
Flow Cytometry (Intracellular) - FITC Anti-Cytokeratin 18 antibody [DC-10] (ab72813)

Flow cytometry analysis (intracellular staining) of HeLa cells with ab72813 at 5 $\mu g/ml$.



Flow Cytometry (Intracellular) - FITC Anti-Cytokeratin 18 antibody [DC-10] (ab72813)

Overlay histogram showing MCF-7 cells stained with ab72813 (red line). The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab72813, 0.5 μ g/1x10⁶ cells) for 30 min at 22°C. Isotype control antibody (black line) was mouse IgG1 (FITC) (2 μ g/1x10⁶ cells) for 30 min at 22°C. Acquisition of >5,000 events was performed.



Immunocytochemistry/ Immunofluorescence - FITC
Anti-Cytokeratin 18 antibody [DC-10] (ab72813)

ICC/IF image of ab72813 stained HCT116 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 (ab72813, 1 μ g/ml) overnight at +4°C. 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.

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

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