abcam

Product datasheet

Anti-IFT88 antibody ab42497

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Overview

Product name Anti-IFT88 antibody

Description Goat polyclonal to IFT88

Host species Goat

Specificity This antibody is expected to recognise both reported isoforms (NP 783195.2; NP 006522.2).

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

Species reactivity Reacts with: Mouse, Human

Predicted to work with: Rat, Dog

Immunogen Synthetic peptide corresponding to Human IFT88 aa 804-816 (C terminal).

Sequence:

C-KKRIDEDDFADEE

Run BLAST with
Run BLAST with

Positive control WB: A549 cell lysate and Mouse testis tissue lysate; ICC/IF: HepG2 cells; Flow Cyt (intra): HepG2

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. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Storage buffer pH: 7.30

Preservative: 0.02% Sodium azide

Constituent: 0.5% BSA

Purity Immunogen affinity purified

Purification notes Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity

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chromatography using the immunizing peptide.

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab42497 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	★★★★ (1)	Use a concentration of 10 μg/ml.
Flow Cyt (Intra)		Use a concentration of 10 μg/ml.
WB		Use a concentration of 1 - 3 µg/ml. Detects a band of approximately 90 kDa (predicted molecular weight: 94 kDa). 1 hour primary incubation is recommended for this product.

Target

Function Involved in primary cilium biogenesis.

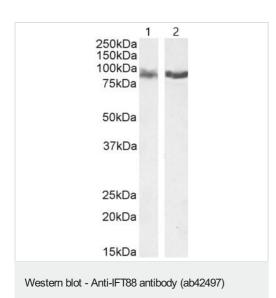
Tissue specificity Expressed in the heart, brain, liver, lung, kidney, skeletal muscle and pancreas.

Sequence similarities Contains 12 TPR repeats.

Cellular localization Cytoplasm > cytoskeleton > centrosome > centriole. Cytoplasm > cytoskeleton > cilium basal

body.

Images



All lanes: Anti-IFT88 antibody (ab42497) at 3 µg/ml

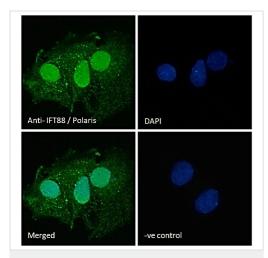
Lane 1: A549 (human lung carcinoma cell line) whole cell lysate

Lane 2: Mouse testis tissue lysate

Lysates/proteins at 35 µg per lane.

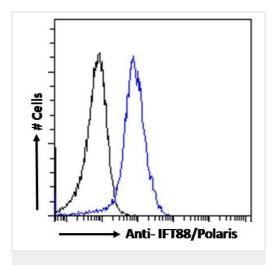
Predicted band size: 94 kDa

Primary incubation 1 hour at room temperature. Detected by chemiluminescence.



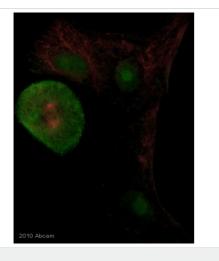
Immunocytochemistry/ Immunofluorescence - Anti-IFT88 antibody (ab42497)

Immunofluorescence analysis of paraformaldehyde fixed HepG2 cells labeling IFT88 with ab42497. Cells permeabilized with 0.15% Triton. Primary incubation 1hr (10 μ g/mL) followed by Alexa Fluor® 488 conjugated secondary antibody (2 μ g/mL), showing nuclear and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10 μ g/mL) followed by Alexa Fluor® 488 secondary antibody (2 μ g/mL).



Flow Cytometry (Intracellular) - Anti-IFT88 antibody (ab42497)

Flow cytometric analysis of paraformaldehyde fixed HepG2 cells labeling IFT88 with ab42497 (blue line). Cells permeabilized with 0.5% Triton. Primary incubation 1hr (10 μ g/mL) followed by Alexa Fluor 488® secondary antibody (1 μ g/mL). lgG control: Unimmunized goat lgG (black line) followed by Alexa Fluor® 488 secondary antibody.



Immunocytochemistry/ Immunofluorescence - Anti-IFT88 antibody (ab42497)

This image is a courtesy of Luke Hofkamp

ab42497 staining IFT88 in human prostate epithelial primary cells by Immunocytochemistry/ Immunofluorescence. Cells were fixed with paraformaldehyde and permeabilized with 0.5% Triton ×100. Samples were incubated with primary antibody (1/100:in DMEM w/ 10% FBS) for 1 hour at 200C. An Alexa Fluor ® 546-conjugated donkey polyclonal to goat IgG was used undiluted as secondary antibody.

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