abcam

Product datasheet

Anti-KIF5B antibody ab5629

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Overview

Product name Anti-KIF5B antibody

Description Rabbit polyclonal to KIF5B

Host species Rabbit

Specificity This antibody is specific for KIF5B and does not detect other kinesin isotypes.

Tested applications Suitable for: ICC/IF, WB

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide corresponding to Human KIF5B aa 376-396.

Sequence:

IDEQFDKEKANLEAFTVDKDI

Database link: P33176

(Peptide available as ab41782)

Run BLAST with
Run BLAST with

General notes

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

Properties

Form Liquid

Storage instructions 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.

Storage buffer Preservative: 0.05% Sodium azide

Constituents: 0.1% BSA, 99% PBS

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

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Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab5629 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	★★★★ <u>(1)</u>	1/250.
WB		1/2000 - 1/6000. Predicted molecular weight: 110 kDa.Can be blocked with Human KIF5B peptide (ab41782) .

Target

Function Microtubule-dependent motor required for normal distribution of mitochondria and lysosomes.

Sequence similarities Belongs to the TRAFAC class myosin-kinesin ATPase superfamily. Kinesin family. Kinesin

subfamily.

Contains 1 kinesin motor domain.

DomainComposed of three structural domains: a large globular N-terminal domain which is responsible

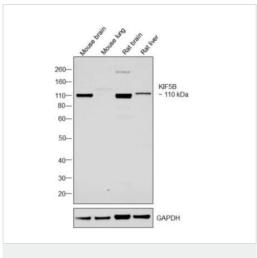
for the motor activity of kinesin (it hydrolyzes ATP and binds microtubule), a central alpha-helical coiled coil domain that mediates the heavy chain dimerization; and a small globular C-terminal domain which interacts with other proteins (such as the kinesin light chains), vesicles and

membranous organelles.

Cellular localization Cytoplasm, cytoskeleton. Uniformly distributed between soma and neurites in hippocampal

neurons.

Images



Western blot - Anti-KIF5B antibody (ab5629)

All lanes: Anti-KIF5B antibody (ab5629) at 1/2000 dilution

Lane 1: Mouse brain Whole cell extract

Lane 2: Mouse lung Whole cell extract

Lane 3: Rat brain Whole cell extract

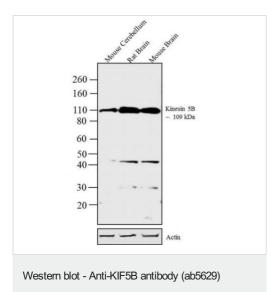
Lane 4: Rat liver Whole cell extract

Lysates/proteins at 40 µg per lane.

Secondary

All lanes: Goat anti-Rabbit lgG (H+L),HRP at 1/4000 dilution

Predicted band size: 110 kDa



All lanes: Anti-KIF5B antibody (ab5629) at 1/4000 dilution

Lane 1: Mouse Cerebellum whole cell extract

Lane 2: Rat Brain whole cell extract

Lane 3: Mouse Brain whole cell extract

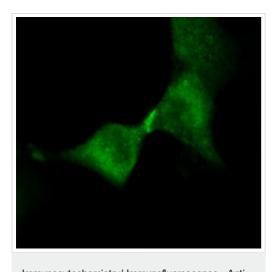
Lysates/proteins at 30 µg per lane.

Secondary

All lanes: Goat anti-Rabbit IgG (H+L) HRP conjugate at 1/2500

dilution

Predicted band size: 110 kDa



Immunocytochemistry/ Immunofluorescence - Anti-KIF5B antibody (ab5629) ab5629 labelling KIF5B (green) in HeLa cells. Cells fixed in 3.2% paraformaldehyde and permeabilized with 0.2% Triton X-100 for 30 minutes at 37°C were blocked with 3% BSA in PBS for 30 minutes at room temperature. Cells were incubated with the primary antibody (1:250) overnight at 4°C. A fluorescently-conjugated Goat anti-rabbit lgG was used as the secondary antibody (1:2000). Images were taken on wide-field fluorescent microscope at 63X magnification.

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

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