abcam

Product datasheet

Biotin Anti-beta Tubulin antibody [BT7R] ab173839

5 Images

Overview

Product name Biotin Anti-beta Tubulin antibody [BT7R]

DescriptionBiotin Mouse monoclonal [BT7R] to beta Tubulin

Host species Mouse

Conjugation Biotin

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

Species reactivity Reacts with: Mouse, Rat, Human, Non human primates, African green monkey

Immunogen Synthetic peptide corresponding to Human beta Tubulin (N terminal) conjugated to Keyhole

Limpet Haemocyanin (KLH).

Database link: P07437

Positive control HeLa, 293T, COS7, NRK and C2C12 cell line lysates.

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.

Storage buffer Preservative: 0.02% Sodium azide

Constituent: 99% PBS

Clonality Monoclonal

Clone number BT7R

Isotype IgG2a

Applications

1

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab173839 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		1/500 - 1/2000.
Flow Cyt		Use at an assay dependent concentration. 1 µg/test. <u>ab97679</u> - Mouse monoclonal lgG2a, is suitable for use as an isotype control with this antibody.
WB		1/500 - 1/1000. Predicted molecular weight: 50 kDa.

Target

Function Tubulin is the major constituent of microtubules. It binds two moles of GTP, one at an

exchangeable site on the beta chain and one at a non-exchangeable site on the alpha chain.

Tissue specificityUbiquitously expressed with highest levels in spleen, thymus and immature brain.

Involvement in disease Cortical dysplasia, complex, with other brain malformations 6

Skin creases, congenital symmetric circumferential, 1

Sequence similarities Belongs to the tubulin family.

Domain The highly acidic C-terminal region may bind cations such as calcium.

Post-translational modifications

Some glutamate residues at the C-terminus are polyglutamylated, resulting in polyglutamate chains on the gamma-carboxyl group (PubMed:26875866). Polyglutamylation plays a key role in microtubule severing by spastin (SPAST). SPAST preferentially recognizes and acts on microtubules decorated with short polyglutamate tails: severing activity by SPAST increases as the number of glutamates per tubulin rises from one to eight, but decreases beyond this

glutamylation threshold (PubMed:26875866).

Some glutamate residues at the C-terminus are monoglycylated but not polyglycylated due to the absence of functional TTLL10 in human. Monoglycylation is mainly limited to tubulin incorporated into axonemes (cilia and flagella). Both polyglutamylation and monoglycylation can coexist on the same protein on adjacent residues, and lowering glycylation levels increases polyglutamylation,

and reciprocally. The precise function of monoglycylation is still unclear.

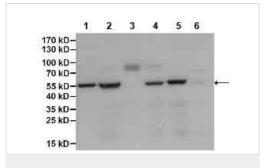
Phosphorylated on Ser-172 by CDK1 during the cell cycle, from metaphase to telophase, but not

in interphase. This phosphorylation inhibits tubulin incorporation into microtubules.

Cellular localization

Cytoplasm, cytoskeleton.

Images



Western blot - Biotin Anti-beta Tubulin antibody
[BT7R] (ab173839)

All lanes: Biotin Anti-beta Tubulin antibody [BT7R] (ab173839) at

1/1000 dilution

Lane 1 : HeLa cell lysate
Lane 2 : 293T cell lysate

Lane 3: A431 cell lysate

Lane 4 : COS7 cell lysate

Lane 5 : C2C12 cell lysate

Lane 6: NRK cell lysate

Lysates/proteins at 50 µg per lane.

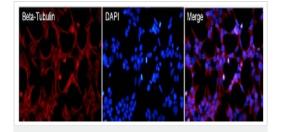
Secondary

All lanes: Streptavidin-HRP at 1/20000 dilution

Developed using the ECL technique.

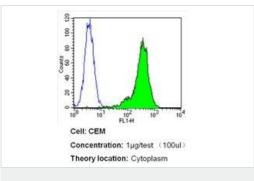
Predicted band size: 50 kDa

4-20% Tris-HCI polyacrylamide gel.

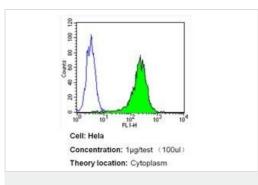


Immunocytochemistry/ Immunofluorescence - Biotin
Anti-beta Tubulin antibody [BT7R] (ab173839)

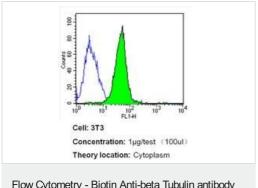
Immunofluorescent analysis of Beta-Tubulin (red) in HEK293T cells. Cells fixed in 4% formaldehyde were permeabilized and blocked with 1X PBS containing 5% BSA and 0.3% Triton X-100 for 1 hour at room temperature. Cells were probed with a Beta-Tubulin monoclonal antibody (ab173839) at a dilution of 1:100 overnight at 4°C in 1X PBS containing 1% BSA and 0.3% Triton X-100, washed with 1X PBS, and incubated with a fluorophore-conjugated goat anti-mouse IgG secondary antibody at a dilution of 1:200 for 1 hour at room temperature. Nuclei (blue) were stained with DAPI. Images were taken at 40X magnification.



Flow Cytometry - Biotin Anti-beta Tubulin antibody [BT7R] (ab173839)



Flow Cytometry - Biotin Anti-beta Tubulin antibody [BT7R] (ab173839)



Flow Cytometry - Biotin Anti-beta Tubulin antibody [BT7R] (ab173839)

Flow cytometry analysis of Beta Tubulin in CEM cells (green) compared to an isotype control (blue). Cells were harvested, adjusted to a concentration of 1-5x10^6 cells/ml, fixed with 2% paraformaldehyde and washed with PBS. Cells were blocked with a 2% solution of BSA-PBS for 30 min at room temperature and incubated with a Beta Tubulin loading control antibody (ab173839) at a dilution of 1 ug/test for 40 min at room temperature. Cells were then incubated for 40 min at room temperature in the dark using a Dylight 488-conjugated secondary antibody and re-suspended in PBS for FACS analysis.

Flow cytometry analysis of Beta Tubulin in Hela cells (green) compared to an isotype control (blue). Cells were harvested, adjusted to a concentration of 1-5x10^6 cells/ml, fixed with 2% paraformaldehyde and washed with PBS. Cells were blocked with a 2% solution of BSA-PBS for 30 min at room temperature and incubated with a Beta Tubulin loading control antibody (ab173839) at a dilution of 1 ug/test for 40 min at room temperature. Cells were then incubated for 40 min at room temperature in the dark using a Dylight 488-conjugated secondary antibody and re-suspended in PBS for FACS analysis.

Flow cytometry analysis of Beta Tubulin in NIH-3T3 cells (green) compared to an isotype control (blue). Cells were harvested, adjusted to a concentration of 1-5x10^6 cells/ml, fixed with 2% paraformaldehyde and washed with PBS. Cells were blocked with a 2% solution of BSA-PBS for 30 min at room temperature and incubated with a Beta Tubulin loading control antibody (ab173839) at a dilution of 1 ug/test for 40 min at room temperature. Cells were then incubated for 40 min at room temperature in the dark using a Dylight 488-conjugated secondary antibody and re-suspended in PBS for FACS analysis.

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

Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you

• We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit https://www.abcam.com/abpromise or contact our technical team.

Terms and conditions

• Guarantee only valid for products bought direct from Abcam or one of our authorized distributors