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Product datasheet

Anti-alpha Tubulin (acetyl K40) antibody [6-11B-1] ab24610

★★★★★ 33 Abreviews 193 References 画像数 5

製品の概要

製品名 Anti-alpha Tubulin (acetyl K40) antibody [6-11B-1]

製品の詳細 Mouse monoclonal [6-11B-1] to alpha Tubulin (acetyl K40)

由来種 Mouse

特異性 ab24610 detects acetylated alpha tubulin.

アプリケーション 適用あり: Flow Cyt, ICC, WB 種交差性 交差種: Mouse, Rat, Human

交差が予測される動物種: Cow 🔷

免疫原 Tissue, cells or virus corresponding to alpha Tubulin (acetyl K40).

エピトープ The antibody recognizes an epitope located on the a3 isoform of Chlamydomonas axonemal a-

tubulin, within four residues of Lys40 when this amino acid is acetylated.

ポジティブ・コントロール In Western Blot, this antibody gave a positive signal in mouse brain tissue lysate and in the

following whole cell lysates: HeLa; NIH3T3; PC12.

特記事項 Production of this antibody has been changed on 8th April 2016. This antibody is now purified

from tissue culture supernatant. This shouldn't affect the use of this antibody but if you have any

issues, please contact our Scientific Support team.

This antibody binds to primary cilia, centrioles, mitotic spindles, midbodies and to subsets of

cytoplasmic microtubules in 3T3 and HeLa cells.

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

バッファー pH: 7.40

1

Preservative: 0.097% Sodium azide

Constituent: PBS

精製度 Proprietary Purification

特記事項(精製) Purified from Tissue culture supernatant.

ポリ/モノ モノクローナル

クローン名6-11B-1アイソタイプIgG2b軽鎖の種類kappa

アプリケーション

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ab24610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 $1\mu I$ for 10^6 cells. <u>ab170192</u> - Mouse monoclonal lgG2b, is suitable for use as an isotype control with this antibody.
ICC		Use at an assay dependent concentration.
WB	★★★★ ★ <u>(12)</u>	Use a concentration of 0.03 - 0.06 µg/ml. Detects a band of approximately 55 kDa (predicted molecular weight: 55 kDa).

ターゲット情報

機能 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.

exclusively on glutamate residues and results in polyglutamate chains on the gamma-carboxyl

配列類似性 Belongs to the tubulin family.

翻訳後修飾 Some glutamate residues at the C-terminus are polyglutamylated. This modification occurs

group. Also 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) whereas glutamylation is prevalent in neuronal cells, centrioles, axonemes, and the mitotic spindle. Both modifications can coexist on the same protein on adjacent residues, and lowering glycylation levels increases polyglutamylation, and reciprocally. The precise function of such

microtubules.

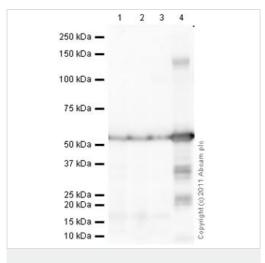
Acetylation of alpha chains at Lys-40 stabilizes microtubules and affects affinity and processivity of microtubule motors. This modification has a role in multiple cellular functions, ranging from cell

motility, cell cycle progression or cell differentiation to intracellular trafficking and signaling.

modifications is still unclear but they regulate the assembly and dynamics of axonemal

細胞内局在 Cytoplasm > cytoskeleton.

画像



Western blot - Anti-alpha Tubulin (acetyl K40) antibody [6-11B-1] (ab24610)

All lanes : Anti-alpha Tubulin (acetyl K40) antibody [6-11B-1] (ab24610) at 5 µg/ml

Lane 1 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

Lane 2: NIH 3T3 (Mouse embryonic fibroblast cell line) Whole Cell Lysate

Lane 3: PC12 (Rat adrenal pheochromocytoma cell line) Whole Cell Lysate

Lane 4: Brain (Mouse) Tissue Lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat Anti-Mouse lgG H&L (HRP) preadsorbed (ab97040) at 1/5000 dilution

Performed under reducing conditions.

Predicted band size: 55 kDa **Observed band size:** 55 kDa

Additional bands at: 140 kDa, 25 kDa, 35 kDa. We are unsure as

to the identity of these extra bands.

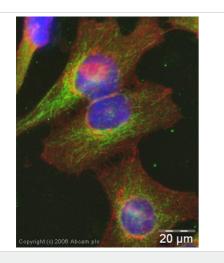
Exposure time: 150 seconds

10/µm 2014 Abcam

Immunocytochemistry - Anti-alpha Tubulin (acetyl K40) antibody [6-11B-1] (ab24610)

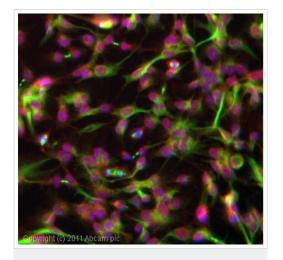
This image is courtesy of an Abreview submitted by Aaron Halpen

ab24610 staining Acetylated alpha Tubulin in monkey kidney cells by ICC/IF (Immunocytochemistry/immunofluorescence). Cells were fixed with 3% PFA + 0.1% GA and blocked with 3% BSA + 0.5% Triton X-100 for 45 minutes at 25°C. Samples were incubated with primary antibody (1/100 in 3% BSA + 0.5% Triton X-100) for 1 hour at 21°C. An Alexa Fluor $^{\circledR}$ 647-conjugated donkey anti-rabbit IgG polyclonal (2 µg/ml) was used as the secondary antibody.



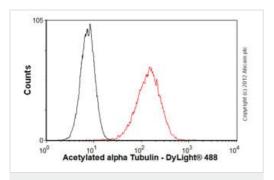
Immunocytochemistry - Anti-alpha Tubulin (acetyl K40) antibody [6-11B-1] (ab24610)

ICC/IF image of ab24610 stained human HeLa cells. The cells were methanol fixed (5 min) and incubated with the antibody (ab24610, 1µg/ml) for 1h at room temperature. The secondary antibody (green) was Alexa Fluor[®] 488 goat anti-mouse IgG (H+L) used at a 1/1000 dilution for 1h. Image-iT[™] FX Signal Enhancer was used as the primary blocking agent, 5% BSA (in TBS-T) was used for all other blocking steps. DAPI was used to stain the cell nuclei (blue). Alexa Fluor[®] 594 WGA was used to label plasma membranes (red).



Immunocytochemistry - Anti-alpha Tubulin (acetyl K40) antibody [6-11B-1] (ab24610)

ICC/IF image of ab24610 stained HepG2 cells. The cells were 4% PFA 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 (ab24610, 5 μ g/ml) overnight at +4°C. The secondary antibody (green) was DyLight® 488 goat anti-mouse lgG - H&L, preadsorbed (ab96879) used at a 1/250 dilution for 1h. 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.



Flow Cytometry - Anti-alpha Tubulin (acetyl K40) antibody [6-11B-1] (ab24610)

Overlay histogram showing HeLa cells stained with ab24610 (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 (ab24610, 1µg/1x10⁶ cells) for 30 min at 22°C. (This data was generated from a purified version of the antibody. Some lots are produced as ascites fluid. We suggest 1µl/1x10⁶ cells for ascites preparations). The secondary antibody used was DyLight[®] 488 goat anti-mouse IgG (H+L) (ab96879) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse IgG2b [PLPV219] (ab91366, 2µg/1x10⁶ cells) used under the same conditions. Acquisition of >5,000 events was performed. This

antibody gave a positive signal in HeLa cells fixed with 80% methanol (5 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.

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