abcam

Product datasheet

HRP Anti-alpha Tubulin antibody [EPR13478(B)] - Loading Control ab185067

יעלאעבע RabMAb

5 References 画像数6

製品の概要

製品名 HRP Anti-alpha Tubulin antibody [EPR13478(B)] - Loading Control

製品の詳細 HRP Rabbit monoclonal [EPR13478(B)] to alpha Tubulin - Loading Control

由来種 Rabbit 標識 HRP

アプリケーション 適用あり: IHC-P, WB

種交差性 交差種: Mouse, Rat, Human

交差が予測される動物種: African green monkey 🔷

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ポジティブ・コントロール HeLa, Jurkat, A431 and K562 cell lysates; Human kidney and uterus tissues; A431 and Jurkat

cells. IHC: normal human colon tissue, normal human spleen tissue, rat and mouse spleen tissue.

特記事項 This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle. Store In the Dark.

バッファー pH: 7.40

Preservative: 0.1% 10% Proclin 300 Solution

Constituents: PBS, 1% BSA, 30% Glycerol (glycerin, glycerine)

精製度 Protein A purified

ポリ/モノ モノクローナル **ウローン名** EPR13478(B)

アイソタイプ IgG

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ab185067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
IHC-P		Use a concentration of 1 μ g/ml. Perform heat mediated antigen retrieval before commencing with IHC staining protocol. <u>ab199507</u> - Rabbit monoclonal lgG (HRP), is suitable for use an as isotype control with this antibody.
WB		1/5000. Predicted molecular weight: 50 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.

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

exclusively on glutamate residues and results in polyglutamate chains on the gamma-carboxyl 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 modifications is still unclear but they regulate the assembly and dynamics of axonemal

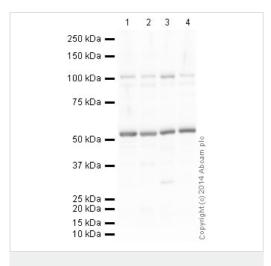
modifications is still unclear but they regulate the assembly and dynamics of a microtubules.

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.

細胞内局在 Cytoplasm > cytoskeleton.

画像



Western blot - HRP Anti-alpha Tubulin antibody [EPR13478(B)] - Loading Control (ab185067)

All lanes : HRP Anti-alpha Tubulin antibody [EPR13478(B)] - Loading Control (ab185067) at 1/5000 dilution

Lane 1: HeLa whole cell lysate (ab150035)

Lane 2 : Jurkat (Human T cell lymphoblast-like cell line) Whole Cell

Lane 3: A431 (Human epithelial carcinoma cell line) Whole Cell Lysate

Lane 4 : K562 (Human erythromyeloblastoid leukemia cell line) Whole Cell Lysate

Lysates/proteins at 10 µg per lane.

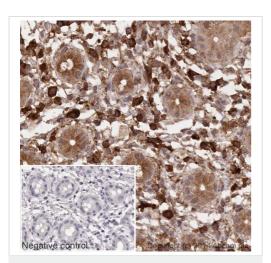
Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 50 kDa **Observed band size:** 52 kDa

Exposure time: 10 seconds

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Bovine Serum Albumin before being incubated with ab185067 overnight at 4°C. Antibody binding was visualised using ECL development solution ab133406



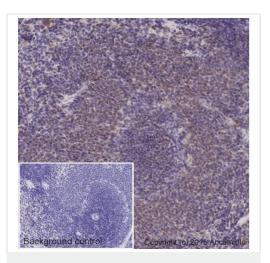
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - HRP Anti-alpha Tubulin antibody [EPR13478(B)] - Loading Control (ab185067)

IHC image of alpha Tubulin staining in a section of formalin-fixed paraffin-embedded normal human colon*. The section was pretreated using pressure cooker heat mediated antigen retrieval with sodium citrate buffer (pH6) for 30mins. The section was incubated with ab185067, 1 μ g/ml overnight at +4°C. The section was counterstained with haematoxylin and mounted with DPX.

The inset negative control image is taken from an identical assay without primary antibody.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

*Tissue obtained from the Human Research Tissue Bank, supported by the NIHR Cambridge Biomedical Research Centre

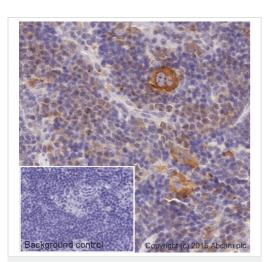


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - HRP Anti-alpha Tubulin antibody [EPR13478(B)] - Loading Control (ab185067)

IHC image of alpha tubulin staining in a section of formalin-fixed paraffin-embedded normal rat spleen, performed on a Leica BONDTM. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20mins. The section was then incubated with ab185067, 1/200 dilution, for 15 mins at room temperature. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

The inset background control image is taken from an identical assay without primary antibody.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

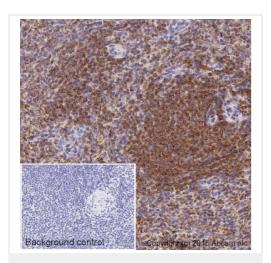


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - HRP Anti-alpha Tubulin antibody [EPR13478(B)] - Loading Control (ab185067)

IHC image of alpha tubulin staining in a section of formalin-fixed paraffin-embedded normal mouse spleen, performed on a Leica BONDTM. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20mins. The section was then incubated with ab185067, 1/200 dilution, for 15 mins at room temperature. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

The inset background control image is taken from an identical assay without primary antibody.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - HRP Anti-alpha Tubulin antibody [EPR13478(B)] - Loading Control (ab185067)

IHC image of alpha tubulin staining in a section of formalin-fixed paraffin-embedded normal human spleen*, performed on a Leica BONDTM. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20mins. The section was then incubated with ab185067, 1/200 dilution, for 15 mins at room temperature. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

The inset background control image is taken from an identical assay without primary antibody.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

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