abcam

Product datasheet

Anti-CaMKII antibody [EP1829Y] - BSA and Azide free ab271841

יעלאעבע RabMAb

画像数 4

製品の概要

ポジティブ・コントロール

特記事項

製品名 Anti-CaMKII antibody [EP1829Y] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EP1829Y] to CaMKII - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: ICC/IF, IHC-P, WB

適用なし: Flow Cyt or IP

種交差性 交差種: Rat, Human

交差が予測される動物種: Mouse

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

IHC-P: Human hepatocellulas carcinoma tissue. ICC/IF: U87-MG and PC12 cells.

ab271841 is the carrier-free version of ab52476.

Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.

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. Do Not Freeze.

バッファー pH: 7.20

Constituent: PBS

キャリア・フリー はい

精製度 Protein A purified

ポリモノクローナル **ウローン名** EP1829Y

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ab271841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。

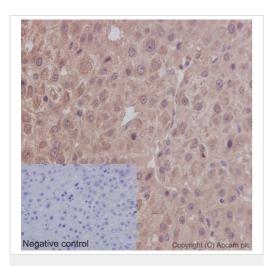
アプリケーション	Abreviews	特記事項
ICC/IF		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 54 kDa. For unpurified use at 1/20000.

追加情報 Is unsuitable for Flow Cyt or IP.

ターゲット情報

機能	CaM-kinase II (CAMK2) is a prominent kinase in the central nervous system that may function in long-term potentiation and neurotransmitter release. Member of the NMDAR signaling complex in excitatory synapses it may regulate NMDAR-dependent potentiation of the AMPAR and synaptic plasticity.
配列類似性	Belongs to the protein kinase superfamily. CAMK Ser/Thr protein kinase family. CaMK subfamily. Contains 1 protein kinase domain.
細胞内局在	Cell junction > synapse > presynaptic cell membrane. Cell junction > synapse. Postsynaptic lipid rafts.

画像

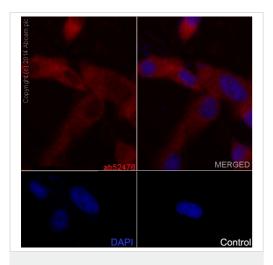


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CaMKII antibody

[EP1829Y] - BSA and Azide free (ab271841)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human hepatocellular carcinoma tissue labelling CaMKII with purified <u>ab52476</u> at 1/500. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit IgG was used as the secondary antibody. Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab52476).

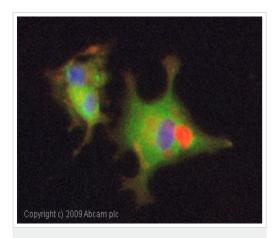


Immunocytochemistry/ Immunofluorescence - Anti-CaMKII antibody [EP1829Y] - BSA and Azide free (ab271841)

Immunocytochemistry/Immunofluorescence analysis of U87-MG cells labelling CaMKII with purified <u>ab52476</u> at 1/250. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% aTriton X-100. An Alexa Fluor[®] 555-conjugated goat anti-rabbit IgG (1/500) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain.

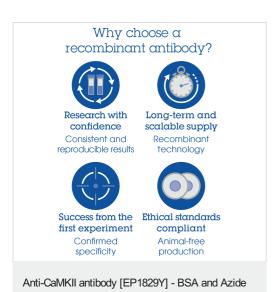
Control: primary antibody (1/250) and secondary antibody, **ab150113**, an Alexa Fluor[®] 488-conjugated goat anti-mouse IgG (1/500).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab52476</u>).



Immunocytochemistry/ Immunofluorescence - Anti-CaMKII antibody [EP1829Y] - BSA and Azide free (ab271841) ICC/IF image of unpurified <u>ab52476</u> stained PC12 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 (<u>ab52476</u>, 1μg/ml) overnight at +4°C. The secondary antibody (green)ÿwas Alexa Fluor[®] 488 goat anti-rabbit lgG (H+L) used at a 1/1000 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 1.43μM.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab52476).



free (ab271841)

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