abcam

Product datasheet

Anti-DARPP32 antibody [EP720Y] - BSA and Azide free ab220808

重组 RabMAb

<u>18 References</u> 7 图像

概述		
产品名称	Anti-DARPP32 抗体 [EP720Y] - BSA and Azide free	
描述	兔单 克隆抗体 [EP720Y] to DARPP32 - BSA and Azide free	
宿主	Rabbit	
经测试应 用	适用于: WB, IHC-P, ICC/IF, IP	
种属反应性	与反应: Mouse, Rat, Human	
	预测可用于: Pig 🔺	
免疫原	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.	
阳性 对照	Rat brain, cerebral cortex and hippocampus tissue; Mouse brain and cerebral cortex tissue; Human breast adenocarcinoma; Human fetal brain tissue lysate; Human colon tissue; pig brain tissue.	
常 规说 明	ab220808 is the carrier-free version of <u>ab40801</u> .	
	Our <u>carrier-free</u> 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 cell-based 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.	
	Our RabMAb [®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <u>RabMAb</u>® patents.	

形式	Liquid	
存放说明	Shipped at 4°C. Store at +4°C. Do Not Freeze.	
存储溶液	pH: 7.20 Constituent: PBS	
无载体	是	
纯 度	Protein A purified	
克隆	单 克隆	
克 隆 编号	EP720Y	
同种型	lgG	

应用

The Abpromise guarantee Abpromise™承诺保证使用ab220808于以下的经测试应用

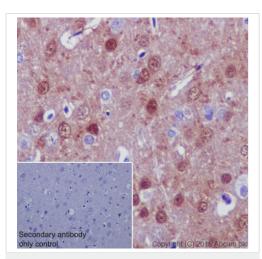
"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		Use at an assay dependent concentration. Detects a band of approximately 32 kDa (predicted molecular weight: 32 kDa).
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. See IHC antigen retrieval protocols .
ICC/IF		Use at an assay dependent concentration.
IP		Use at an assay dependent concentration.

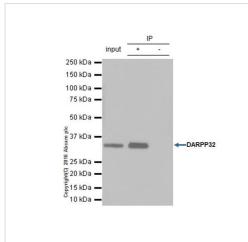
靶标

Inhibitor of protein-phosphatase 1.
Belongs to the protein phosphatase inhibitor 1 family.
Dopamine- and cyclic AMP-regulated neuronal phosphoprotein. Phosphorylation of Thr-34 is required for activity.
Cytoplasm.

图片



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DARPP32 antibody [EP720Y] - BSA and Azide free (ab220808)



Immunoprecipitation - Anti-DARPP32 antibody [EP720Y] - BSA and Azide free (ab220808) Immunohistochemical analysis of paraffin-embedded Rat cerebral cortex sections labelling DARPP32 with purified <u>ab40801</u> at dilution of 1/50. The secondary antibody used was <u>ab97051</u>; a goat anti-rabbit IgG H&L (HRP) at dilution of 1/500. The sample was counterstained with hematoxylin. Antigen retrieval was performed using EDTA Buffer; pH 9.0. PBS was used instead of the primary antibody as the negative control and is shown in the inset.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab40801**).

<u>ab40801</u> at 1/20 immunoprecipitating DARPP32 in rat brain whole cell lysate observed at 32 KDa (lanes 1 and 2).

Lane 1 (input): Rat brain whole cell lysate 10µg

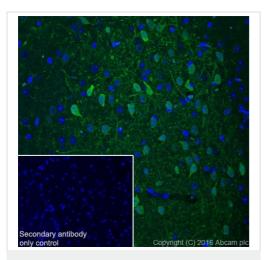
Lane 2 (+): <u>ab40801</u> + Rat brain whole cell lysate.

Lane 3 (-): Rabbit monoclonal lgG (<u>ab172730</u>) instead of <u>ab40801</u> in Rat brain whole cell lysate

For western blotting, <u>**ab131366**</u> VeriBlot for IP (HRP) was used for detection at 1/1000 dilution.

Blocking buffer and concentration: 5% NFDM/TBST. Diluting buffer and concentration: 5% NFDM /TBST.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab40801</u>).

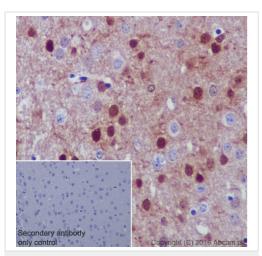


Immunocytochemistry/ Immunofluorescence - Anti-DARPP32 antibody [EP720Y] - BSA and Azide free (ab220808)

Immunocytochemistry/Immunofluorescence analysis of mouse brain tissue lysate labelling DARPP32 with purified <u>**ab40801**</u> at 1/100 (3.4 µg/mL). Cells were fixed with 4% Paraformaldehyde and permeabilized with 0.2% Triton X-100. Antigen retrieval was performed with Heated citrate solution (10mM citrate PH 6.0 + 0.05% Tween-20). <u>**ab150077**</u>, Alexa Fluor[®] 488-conjugated goat anti-rabbit IgG (1/1000, 2 µg/mL) was used as the secondary antibody.

Secondary Only Control: PBS was used instead of the primary antibody as the negative control and is shown in the inset.

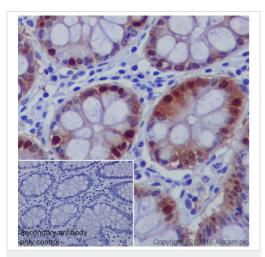
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab40801</u>).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DARPP32 antibody [EP720Y] - BSA and Azide free (ab220808)

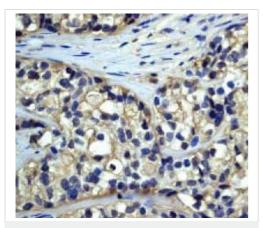
Immunohistochemical analysis of paraffin-embedded Mouse cerebral cortex sections labelling DARPP32 with purified <u>ab40801</u> at dilution of 1/50. The secondary antibody used was <u>ab97051</u>; a goat anti-rabbit IgG H&L (HRP) at dilution of 1/500. The sample was counterstained with hematoxylin. Antigen retrieval was performed using EDTA Buffer; pH 9.0. PBS was used instead of the primary antibody as the negative control and is shown in the inset.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab40801</u>).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DARPP32 antibody [EP720Y] - BSA and Azide free (ab220808)

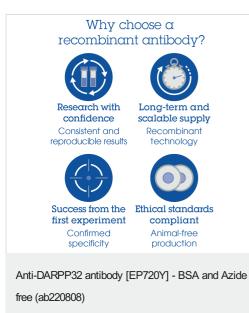
Immunohistochemical analysis of paraffin-embedded human colon sections labelling DARPP32 with purified **ab40801** at dilution of 1/50. The secondary antibody used was **ab97051**; a goat antirabbit IgG H&L (HRP) at dilution of 1/500. The sample was counterstained with hematoxylin. Antigen retrieval was performed using EDTA Buffer; pH 9.0. PBS was used instead of the primary antibody as the negative control and is shown in the inset. This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab40801**).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DARPP32 antibody [EP720Y] - BSA and Azide free (ab220808)

Immunohistochemical analysis of human breast adenocarcinoma sections labelling DARPP32 with unpurified **<u>ab40801</u>** at a dilution of 1/50.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab40801</u>).



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