# abcam

### Product datasheet

## Anti-MAP2 antibody [EPR22036-127] ab221693

Recombinant RabMAb

#### 1 References 7 Images

#### Overview

**Product name** Anti-MAP2 antibody [EPR22036-127]

**Description** Rabbit monoclonal [EPR22036-127] to MAP2

**Host species** Rabbit

**Tested applications** Suitable for: IHC-Fr, ICC/IF, Flow Cyt (Intra), IP

Species reactivity Reacts with: Mouse, Human

**Immunogen** Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control Flow Cyt: Mouse brain cell. IP: SK-N-BE cell.

**General notes** 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**.

#### **Properties**

**Form** Liquid

Storage instructions 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.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

**Purity** Protein A purified

Clonality Monoclonal

Clone number EPR22036-127

Isotype lgG

#### **Applications**

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab221693 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
IHC-Fr		1/1000.
ICC/IF		1/1000.
Flow Cyt (Intra)		1/500.
IP		1/30.

#### **Target**

Function	The exact function of MAP2 is unknown but MAPs ma	y stabilize the microtubules against
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depolymerization. They also seem to have a stiffening effect on microtubules.

**Sequence similarities**Contains 3 Tau/MAP repeats.

Post-translational modifications

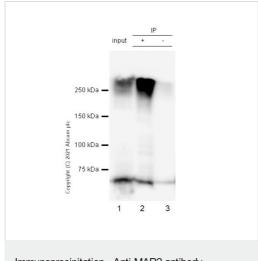
Phosphorylated at serine residues in K-X-G-S motifs by MAP/microtubule affinity-regulating kinase (MARK1 or MARK2), causing detachment from microtubules, and their disassembly (By similarity). Isoform 2 is probably phosphorylated by PKA at Ser-323, Ser-354 and Ser-386 and by

FYN at Tyr-67.

Cellular localization

Cytoplasm, cytoskeleton.

#### **Images**



Immunoprecipitation - Anti-MAP2 antibody [EPR22036-127] (ab221693)

MAP2 was immunoprecipitated from 0.35 mg SK-N-BE(2) (Human neuroblastoma neuroblast) whole cell lysate 10 ug with ab221693 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab221693 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP)(ab131366) was used at 1/5000 dilution.

Lane 1: SK-N-BE(2) (Human neuroblastoma neuroblast) whole cell lysate 10 ug

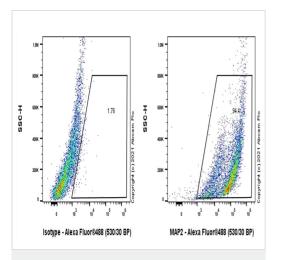
Lane 2: ab221693 IP in SK-N-BE(2) whole cell lysate

Lane 3: Rabbit monoclonal  $\lg G (\underline{ab172730})$  instead of ab221693 in SK-N-BE(2) whole cell lysate

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

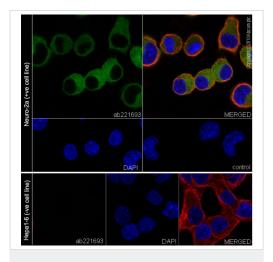
Exposure time: 8 seconds

Fresh made lysate is required to minimize protein degradation.



Flow Cytometry (Intracellular) - Anti-MAP2 antibody [EPR22036-127] (ab221693)

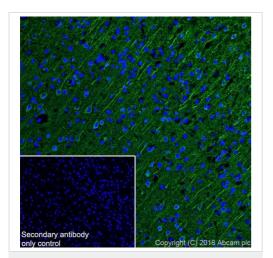
Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized Mouse brain cells cells labelling MAP2 with ab221693 at 1/500 dilution (0.1ug)/ Right compared with a Rabbit monoclonal lgG (ab172730) / Left isotype control. A Goat anti rabbit lgG (Alexa Fluor® 488, ab150077) at 1/2000 dilution was used as the secondary antibody.



Immunocytochemistry/ Immunofluorescence - Anti-MAP2 antibody [EPR22036-127] (ab221693)

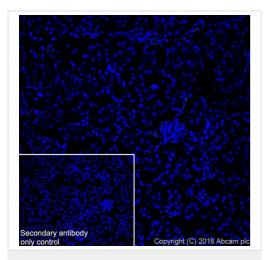
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized Neuro-2a (Mouse neuroblastoma neuroblast) cells labeling MAP2 with ab221693 at 1/1000 dilution, followed by AlexaFluor<sup>®</sup>488 Goat anti-Rabbit (ab150077) secondary antibody at 1/1000 dilution (Green). Confocal image showing cytoplasmic staining in Neuro-2a cells is observed. Antialpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor<sup>®</sup> 594) (ab195889) was used as a counterstain at 1/200 dilution (Red). The nuclear counterstain was DAPI (Blue).

Negative control: Hepa1-6 cells (PMID: 7001466)



Immunohistochemistry (Frozen sections) - Anti-MAP2 antibody [EPR22036-127] (ab221693)

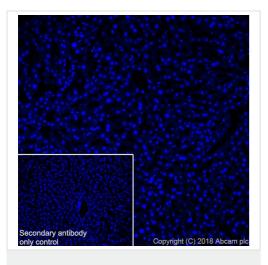
Immunohistochemical analysis of frozen 4% paraformaldehyde-fixed 0.2% Triton X-100 permeabilized mouse cerebrum tissue labeling MAP2 (Green) using ab221693 at 1/1000 dilution, followed by an AlexaFluor<sup>®</sup>488 Goat anti-Rabbit secondary antibody at 1/1000 dilution. Cytoplasmic staining on mouse cerebrum (PMID: 22479190) is observed. The nuclear counterstain is DAPI (Blue). Heat mediated antigen retrieval using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).



Immunohistochemistry (Frozen sections) - Anti-MAP2 antibody [EPR22036-127] (ab221693)

Immunohistochemical analysis of frozen 4% paraformaldehydefixed 0.2% Triton X-100 permeabilized mouse kidney tissue labeling MAP2 using ab221693 at 1/1000 dilution, followed by an AlexaFluor®488 Goat anti-Rabbit secondary antibody at 1/1000 dilution. Negative control: No staining on mouse kidney (PMID: 7001466, PMID: 2423532) is observed. The nuclear counterstain is DAPI (Blue).

Heat mediated antigen retrieval using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).



Immunohistochemistry (Frozen sections) - Anti-MAP2 antibody [EPR22036-127] (ab221693)

Immunohistochemical analysis of frozen 4% paraformaldehydefixed 0.2% Triton X-100 permeabilized mouse liver tissue labeling MAP2 using ab221693 at 1/1000 dilution, followed by an AlexaFluor®488 Goat anti-Rabbit secondary antibody at 1/1000 dilution. Negative control: No staining on mouse liver (PMID: 7001466, PMID: 2423532). is observed. The nuclear counterstain is DAPI (Blue).

Heat mediated antigen retrieval using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).





Research with confidence Consistent and reproducible results





Success from the Ethical standards first experiment Confirmed specificity



compliant Animal-free production

Anti-MAP2 antibody [EPR22036-127] (ab221693)

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