

ATP2B3 polyclonal antibody

Catalog: BS32358

Host: Rabbit

Reactivity: Human

BackGround:

Plasma membrane calcium-transporting ATPase 3: ATP-driven Ca^{2+} ion pump involved in the maintenance of basal intracellular Ca^{2+} levels at the presynaptic terminals. Uses ATP as an energy source to transport cytosolic Ca^{2+} ions across the plasma membrane to the extracellular compartment. May counter-transport protons, but the mechanism and the stoichiometry of this $\text{Ca}^{2+}/\text{H}^{+}$ exchange remains to be established (By similarity)

Product:

1mg/ml in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.

Molecular Weight:

~134 kDa

Swiss-Prot:

Q16720

Purification&Purity:

The antibody was purified by immunogen affinity chromatography.

Applications:

WB (1/500 - 1/1000), IF/ICC (1/10 - 1/50)

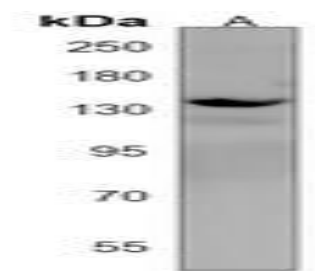
Storage&Stability:

Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

Specificity:

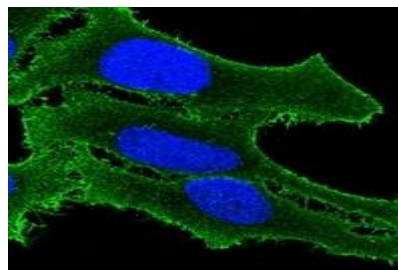
Recognizes endogenous levels of ATP2B3 protein.

DATA:



Western blot analysis of Cathepsin V expression in mouse lung (A), mouse testis (B), rat lung (C), rat testis (D) whole cell lysates.

Immunohistochemical analysis of Cathepsin V staining in rat heart formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of Cathepsin V staining in NIH3T3 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

Note:

For research use only, not for use in diagnostic procedure.

Bioworld Technology, Inc.

Add: 1660 South Highway 100, Suite 500 St. Louis Park, MN 55416, USA.

Email: info@bioworld.com

Tel: 6123263284

Fax: 6122933841

Bioworld technology, co. Ltd.

Add: No 9, weidi road Qixia District Nanjing, 210046, P. R. China.

Email: info@biogot.com

Tel: 0086-025-68037686

Fax: 0086-025-68035151