

VPS13A polyclonal antibody

Catalog: BS31364

Host: Rabbit

Reactivity: Human, Mouse

BackGround:

Intermembrane lipid transfer protein VPS13A: Mediates the transfer of lipids between membranes at organelle contact sites (By similarity). Binds phospholipids . Required for the formation or stabilization of ER-mitochondria contact sites which enable transfer of lipids between the ER and mitochondria . Negatively regulates lipid droplet size and motility . Required for efficient lysosomal protein degradation

Product:

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

Molecular Weight:

~360 kDa

Swiss-Prot:

Q96RL7

Purification&Purity:

The antibody was purified by immunogen affinity chromatography.

Applications:

WB (1/500 - 1/2000)

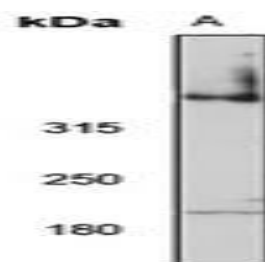
Storage&Stability:

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

Specificity:

Recognizes endogenous levels of VPS13A protein

DATA:



Western blot analysis of IGFBP1 expression in U2OS (A) whole cell lysates.

Immunohistochemical analysis of IGFBP1 staining in human liver 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 IGFBP1 staining in HepG2 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.

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