

Product datasheet for AP39032PU-N

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OriGene Technologies, Inc.

SLC36A1 (1-30) Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

Applications: FC, IF, IHC, WB

Recommended Dilution: Flow Cytometry: 1/10 - 1/50.

Immunofluorescence: 1/10 - 1/50.

Immunohistochemistry on Paraffin Sections: 5 µg/ml.

Western Blot: 1/1000.

Reactivity: Human
Host: Rabbit
Isotype: IgG

Clonality: Polyclonal

Immunogen: KLH-conjugated synthetic peptide from human SLC36A1, N-terminal epitope (aa 1-30)

Specificity: The antibody detects human SLC36A1.

Other species not tested.

Formulation: PBS, containing 0.09% sodium azide

State: Aff - Purified State: Liquid Ig fraction

Purification: Immunoaffinity Chromatography

Conjugation: Unconjugated

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

Gene Name: solute carrier family 36 member 1

Database Link: Entrez Gene 206358 Human

Q7Z2H8

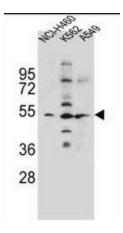
Background: Proton-coupled amino acid transporter 1 has transport activity for small amino acids.

Synonyms: hPAT1; LYAAT-1; LYAAT1; PAT1; TRAMD3

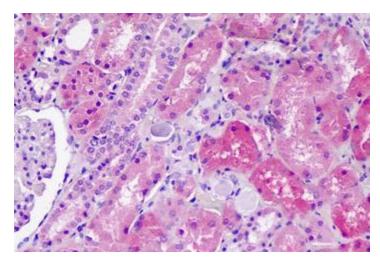




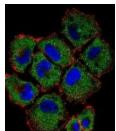
Product images:



Western blot detection of SLC36A1 protein (indicated by arrow) in NCI-H460, K562, and A549 cell line lysates (35 ug/lane), using anti-SLC36A1 / PAT1 antibody cat- no. AP39032PU-N.

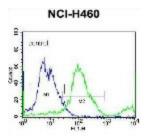


Immunohistochemical staining of Formalin-Fixed, Paraffin-Embedded human Kidney using anti-SLC36A1 antibody cat. no. AP39032PU-N



Confocal immunofluorescence using anti-SLC36A1 antibody cat. no. AP39032PU-N on NCI-H460 cells, followed by Alexa Fluor 488-conjugated goat anti-rabbit IgG (green). Actin filaments have been labeled with Alexa Fluor 555 phalloidin (red).DAPI was used to stain the cell nuclear (blue).





Flow cytometry of NCI-H460 cells (right histogram) compared to a negative control cell (left histogram) using anti-SLC36A1 antibody cat. no. AP39032PU-N. FITC-conjugated donkey-antirabbit secondary antibodies were used for the analysis.