

SLC22A1 Antibody

Purified Mouse Monoclonal Antibody Catalog # AO1492a

Product Information

ApplicationWB, FC, EPrimary AccessionO15245ReactivityHumanHostMouseClonalityMonoclonal

Clone Names2C5IsotypeIgG1Calculated MW61154

Description Polyspecific organic cation transporters in the liver, kidney, intestine, and

other organs are critical for elimination of many endogenous small organic cations as well as a wide array of drugs and environmental toxins. This gene is

one of three similar cation transporter genes located in a cluster on chromosome 6. The encoded protein contains twelve putative

transmembrane domains and is a plasma integral membrane protein. Tissue

specificity: Widely expressed with high level in liver.

Immunogen Purified recombinant fragment of human SLC22A1 expressed in E. Coli.

Formulation Ascitic fluid containing 0.03% sodium azide.

Additional Information

Gene ID 6580

Other Names Solute carrier family 22 member 1, Organic cation transporter 1, hOCT1,

SLC22A1, OCT1

Dilution WB~~1/500 - 1/2000 FC~~1/200 - 1/400 E~~N/A

Storage Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions SLC22A1 Antibody is for research use only and not for use in diagnostic or

therapeutic procedures.

Protein Information

Name SLC22A1 (HGNC:10963)

Synonyms OCT1

Function

Electrogenic voltage-dependent transporter that mediates the transport of a variety of organic cations such as endogenous bioactive amines, cationic drugs and xenobiotics (PubMed: 11388889, PubMed: 11408531, PubMed:12439218, PubMed:12719534, PubMed:15389554, PubMed: 16263091, PubMed: 16272756, PubMed: 16581093, PubMed:19536068, PubMed:21128598, PubMed:23680637, PubMed:24961373, PubMed:34040533, PubMed:9187257, PubMed:9260930, PubMed: 9655880). Functions as a pH- and Na(+)-independent, bidirectional transporter (By similarity). Cation cellular uptake or release is driven by the electrochemical potential (i.e. membrane potential and concentration gradient) and substrate selectivity (By similarity). Hydrophobicity is a major requirement for recognition in polyvalent substrates and inhibitors (By similarity). Primarily expressed at the basolateral membrane of hepatocytes and proximal tubules and involved in the uptake and disposition of cationic compounds by hepatic and renal clearance from the blood flow (By similarity). Most likely functions as an uptake carrier in enterocytes contributing to the intestinal elimination of organic cations from the systemic circulation (PubMed: 16263091). Transports endogenous monoamines such as N-1-methylnicotinamide (NMN), guanidine, histamine, neurotransmitters dopamine, serotonin and adrenaline (PubMed: 12439218, PubMed: 24961373, PubMed:35469921, PubMed:9260930). Also transports natural polyamines such as spermidine, agmatine and putrescine at low affinity, but relatively high turnover (PubMed:<u>21128598</u>). Involved in the hepatic uptake of vitamin B1/thiamine, hence regulating hepatic lipid and energy metabolism (PubMed:<u>24961373</u>). Mediates the bidirectional transport of acetylcholine (ACh) at the apical membrane of ciliated cell in airway epithelium, thereby playing a role in luminal release of ACh from bronchial epithelium (PubMed: 15817714). Transports dopaminergic neuromodulators cyclo(his-pro) and salsolinol with lower efficency (PubMed: 17460754). Also capable of transporting non-amine endogenous compounds such as prostaglandin E2 (PGE2) and prostaglandin F2-alpha (PGF2-alpha) (PubMed: 11907186). May contribute to the transport of cationic compounds in testes across the blood-testis-barrier (Probable). Also involved in the uptake of xenobiotics tributylmethylammonium (TBuMA), quinidine, N-methyl-quinine (NMQ), N- methyl-quinidine (NMQD) N-(4,4-azo-n-pentyl)-quinuclidine (APQ), azidoprocainamide methoiodide (AMP), N-(4,4-azo-n-pentyl)-21- deoxyajmalinium (APDA) and 4-(4-(dimethylamino)styryl)-N- methylpyridinium (ASP) (PubMed: 11408531, PubMed: 15389554, PubMed: 35469921, PubMed: 9260930).

Cellular Location

Basolateral cell membrane; Multi-pass membrane protein. Apical cell membrane; Multi-pass membrane protein. Lateral cell membrane; Multi-pass membrane protein. Basal cell membrane; Multi-pass membrane protein. Cell membrane; Multi-pass membrane protein. Note=Localized to the sinusoidal/basolateral membrane of hepatocytes (By similarity). Mainly localized to the basolateral membrane of renal proximal tubular cells (By similarity). However, also identified at the apical side of proximal tubular cells (PubMed:19536068). Mainly expressed at the lateral membrane of enterocytes (PubMed:23680637). Also observed at the apical side of enterocytes (PubMed:23680637). Localized to the luminal/apical membrane of ciliated epithelial cells in bronchi (PubMed:15817714). Localized to the basal membrane of Sertoli cells (PubMed:35307651) {ECO:0000250 | UniProtKB:Q63089, ECO:0000269 | PubMed:15817714, ECO:0000269 | PubMed:16263091, ECO:0000269 | PubMed:19536068, ECO:0000269 | PubMed:23680637, ECO:0000269 | PubMed:35307651}

Tissue Location

Widely expressed with high level in liver (PubMed:11388889, PubMed:23680637, PubMed:9187257, PubMed:9260930). In liver, expressed around the central vein (PubMed:16263091). Expressed in kidney (PubMed:9187257, PubMed:9260930). Expressed in small intestine

enterocytes (PubMed:16263091, PubMed:23680637). Localized to peritubular myoid cells, Leydig cells and moderately to the basal membrane of Sertoli cells in testes (PubMed:35307651). Expressed in tracheal and bronchial ciliated epithelium in the respiratory tract (PubMed:15817714). Also expressed in skeletal muscle, stomach, spleen, heart, placentacolon, brain, granulycytes and lympohocytes (PubMed:9187257, PubMed:9260930). [Isoform 2]: Expressed in liver and in glial cell lines. [Isoform 4]: Expressed in glial cell lines. Not expressed in liver.

References

1. Leuk Lymphoma. 2008 Nov;49(11):2222-3. 2. Blood. 2008 Oct 15;112(8):3348-54. 3. Pharm Res. 2008 Apr;25(4):827-35.

Images

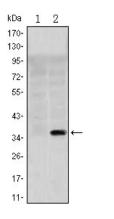


Figure 1: Western blot analysis using SLC22A1 mAb against HEK293 (1) and SLC22A1(AA: 284-347)-hIgGFc transfected HEK293 (2) cell lysate.

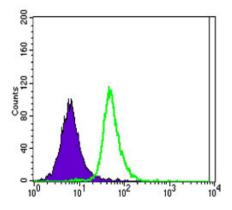


Figure 2: Flow cytometric analysis of Jurkat cells using SLC22A1 mouse mAb (green) and negative control (purple).

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