

SLC22A8 Antibody (C-term) Blocking peptide
Synthetic peptide
Catalog # BP12434b**Specification**

SLC22A8 Antibody (C-term) Blocking peptide - Product InformationPrimary Accession [Q8TCC7](#)**SLC22A8 Antibody (C-term) Blocking peptide - Additional Information**

Gene ID 9376

Other Names

Solute carrier family 22 member 8, Organic anion transporter 3, hOAT3, SLC22A8, OAT3

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

Precautions

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

SLC22A8 Antibody (C-term) Blocking peptide - Protein InformationName SLC22A8 ([HGNC:10972](#))

Synonyms OAT3

Function

Functions as an organic anion/dicarboxylate exchanger that couples organic anion uptake indirectly to the sodium gradient (PubMed: [14586168](http://www.uniprot.org/citations/14586168), PubMed: [15644426](http://www.uniprot.org/citations/15644426), PubMed: [15846473](http://www.uniprot.org/citations/15846473), PubMed: [16455804](http://www.uniprot.org/citations/16455804), PubMed: [31553721](http://www.uniprot.org/citations/31553721)). Transports organic anions such as estrone 3-sulfate (E1S) and urate in exchange for dicarboxylates such as glutarate or ketoglutarate (2-oxoglutarate) (PubMed: [14586168](http://www.uniprot.org/citations/14586168), PubMed: [15846473](http://www.uniprot.org/citations/15846473), PubMed: [15864504](http://www.uniprot.org/citations/15864504), PubMed: [22108572](http://www.uniprot.org/citations/22108572), PubMed: [23832370](http://www.uniprot.org/citations/23832370)). Plays an important role in the excretion of endogenous and exogenous organic anions, especially from the kidney and the brain (PubMed: [14586168](http://www.uniprot.org/citations/14586168), PubMed: [15846473](http://www.uniprot.org/citations/15846473)).

target="_blank">15846473, PubMed:11306713). E1S transport is pH- and chloride- dependent and may also involve E1S/cGMP exchange (PubMed:26377792). Responsible for the transport of prostaglandin E2 (PGE2) and prostaglandin F2(alpha) (PGF2(alpha)) in the basolateral side of the renal tubule (PubMed:11907186). Involved in the transport of neuroactive tryptophan metabolites kynurenate and xanthurenate (PubMed:22108572, PubMed:23832370). Functions as a biopterin transporters involved in the uptake and the secretion of coenzymes tetrahydrobiopterin (BH4), dihydrobiopterin (BH2) and sepiapterin to urine, thereby determining baseline levels of blood biopterins (PubMed:28534121). May be involved in the basolateral transport of steviol, a metabolite of the popular sugar substitute stevioside (PubMed:15644426). May participate in the detoxification/ renal excretion of drugs and xenobiotics, such as the histamine H(2)-receptor antagonists fexofenadine and cimetidine, the antibiotic benzylpenicillin (PCG), the anionic herbicide 2,4-dichloro- phenoxyacetate (2,4-D), the diagnostic agent p-aminohippurate (PAH), the antiviral acyclovir (ACV), and the mycotoxin ochratoxin (OTA), by transporting these exogenous organic anions across the cell membrane in exchange for dicarboxylates such as 2-oxoglutarate (PubMed:15846473, PubMed:16455804, PubMed:11669456). Contributes to the renal uptake of potent uremic toxins (indoxyl sulfate (IS), indole acetate (IA), hippurate/N-benzoylglycine (HA) and 3-carboxy-4-methyl-5-propyl-2- furanpropionate (CMPF)), pravastatin, PCG, E1S and dehydroepiandrosterone sulfate (DHEAS), and is partly involved in the renal uptake of temocaprilat (an angiotensin-converting enzyme (ACE) inhibitor) (PubMed:14675047). May contribute to the release of cortisol in the adrenals (PubMed:15864504). Involved in one of the detoxification systems on the choroid plexus (CP), removes substrates such as E1S or taurocholate (TC), PCG, 2,4-D and PAH, from the cerebrospinal fluid (CSF) to the blood for eventual excretion in urine and bile (By similarity). Also contributes to the uptake of several other organic compounds such as the prostanoids prostaglandin E(2) and prostaglandin F(2-alpha), L-carnitine, and the therapeutic drugs allopurinol, 6-mercaptopurine (6-MP) and 5-fluorouracil (5-FU) (By similarity). Mediates the transport of PAH, PCG, and the statins pravastatin and pitavastatin, from the cerebrum into the blood circulation across the blood-brain barrier (BBB). In summary, plays a role in the efflux of drugs and xenobiotics, helping reduce their undesired toxicological effects on the body (By similarity).

Cellular Location

Basolateral cell membrane; Multi-pass membrane protein. Note=Localizes on the brush border membrane of the choroid epithelial cells (By similarity). Localizes to the basolateral membrane of the proximal tubular cells (PubMed:11306713). Localizes on the abluminal and possibly, luminal membrane of the brain capillary endothelial cells (BCEC) (By similarity). {ECO:0000250, ECO:0000250|UniProtKB:Q9R1U7, ECO:0000269|PubMed:11306713}

Tissue Location

Strongly expressed in kidney (PubMed:11306713, PubMed:11912245). Weaker expression in brain and skeletal muscle (PubMed:11306713). Expressed in adrenal glands (PubMed:15864504)

SLC22A8 Antibody (C-term) Blocking peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

SLC22A8 Antibody (C-term) Blocking peptide - Images

SLC22A8 Antibody (C-term) Blocking peptide - Background

This gene encodes a protein involved in the sodium-independent transport and excretion of organic anions, some of which are potentially toxic. The encoded protein is an integral membrane protein and appears to be localized to the basolateral membrane of the kidney. Multiple alternatively spliced transcript variants that encode different protein isoforms have been described for this gene.

SLC22A8 Antibody (C-term) Blocking peptide - References

Duan, P., et al. Eur. J. Pharmacol. 627 (1-3), 49-55 (2010) :Rodiger, M., et al. Can. J. Physiol. Pharmacol. 88(2):141-146(2010) Saito, A., et al. J. Hum. Genet. 54(6):317-323(2009) Holmes, M.V., et al. PLoS ONE 4 (12), E7960 (2009) :Torres, A.M. World J. Gastroenterol. 14(43):6616-6621(2008)