

SLC2A12 Antibody (C-term) Blocking peptide
Synthetic peptide
Catalog # BP10200b**Specification**

SLC2A12 Antibody (C-term) Blocking peptide - Product Information

Primary Accession [Q8TD20](#)
Other Accession [NP_660159.1](#)

SLC2A12 Antibody (C-term) Blocking peptide - Additional Information

Gene ID 154091

Other Names

Solute carrier family 2, facilitated glucose transporter member 12, Glucose transporter type 12, GLUT-12, SLC2A12, GLUT12, GLUT8

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.

SLC2A12 Antibody (C-term) Blocking peptide - Protein Information

Name SLC2A12 ([HGNC:18067](#))

Function

Insulin-independent facilitative glucose transporter.

Cellular Location

Cell membrane {ECO:0000250|UniProtKB:Q8BFW9}; Multi-pass membrane protein.
Endomembrane system {ECO:0000250|UniProtKB:Q5J316}; Multi-pass membrane protein.
Cytoplasm, perinuclear region. Note=Localizes primarily perinuclear region in the absence of insulin.

Tissue Location

Predominantly expressed in skeletal muscle, heart and prostate, with lower levels in brain, placenta and kidney

SLC2A12 Antibody (C-term) Blocking peptide - Protocols

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

- [Blocking Peptides](#)

SLC2A12 Antibody (C-term) Blocking peptide - Images

SLC2A12 Antibody (C-term) Blocking peptide - Background

SLC2A12 belongs to a family of transporters that catalyze the uptake of sugars through facilitated diffusion (Rogers et al., 2002). This family of transporters show conservation of 12 transmembrane helices as well as functionally significant amino acid residues (Joost and Thorens, 2001 [PubMed 11780753]). [supplied by OMIM].

SLC2A12 Antibody (C-term) Blocking peptide - References

Stuart, C.A., et al. J. Clin. Endocrinol. Metab. 94(9):3535-3542(2009) Vieira, A.R., et al. Genet. Med. 10(9):668-674(2008) Stuart, C.A., et al. Am. J. Physiol. Endocrinol. Metab. 291 (5), E1067-E1073 (2006) :Linden, K.C., et al. Am. J. Physiol. Renal Physiol. 290 (1), F205-F213 (2006) :Rogers, S., et al. Biochem. Biophys. Res. Commun. 308(3):422-426(2003)