

FVT1 Antibody (N-term) Blocking Peptide

Synthetic peptide Catalog # BP7389a

Specification

FVT1 Antibody (N-term) Blocking Peptide - Product Information

Primary Accession

<u>Q06136</u>

FVT1 Antibody (N-term) Blocking Peptide - Additional Information

Gene ID 2531

Other Names

3-ketodihydrosphingosine reductase, KDS reductase, 3-dehydrosphinganine reductase, Follicular variant translocation protein 1, FVT-1, KDSR, FVT1

Target/Specificity

The synthetic peptide sequence used to generate the antibody AP7389a was selected from the N-term region of human FVT1. A 10 to 100 fold molar excess to antibody is recommended. Precise conditions should be optimized for a particular assay.

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.

FVT1 Antibody (N-term) Blocking Peptide - Protein Information

Name KDSR

Synonyms FVT1, SDR35C1

Function

Catalyzes the reduction of 3-ketodihydrosphingosine (KDS) to dihydrosphingosine (DHS).

Cellular Location

Endoplasmic reticulum membrane; Multi-pass membrane protein

Tissue Location

Expressed in all tissues examined. Highest expression in placenta. High expression in lung, kidney, stomach and small intestine, low expression in heart, spleen and skeletal muscle Weakly expressed in normal hematopoietic tissues. Higher expression in some T-cell malignancies and



PHA-stimulated lymphocytes

FVT1 Antibody (N-term) Blocking Peptide - Protocols

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

• Blocking Peptides

FVT1 Antibody (N-term) Blocking Peptide - Images

FVT1 Antibody (N-term) Blocking Peptide - Background

FVT1 catalyzes the reduction of 3-ketodihydrosphingosine to dihydrosphingosine. The putative active site residues of the protein are found on the cytosolic side of the endoplasmic reticulum membrane. A chromosomal rearrangement involving its gene is a cause of follicular lymphoma, also known as type II chronic lymphatic leukemia. The mutation of a conserved residue in the bovine ortholog causes spinal muscular atrophy.

FVT1 Antibody (N-term) Blocking Peptide - References

Krebs, S., Proc. Natl. Acad. Sci. U.S.A. 104 (16), 6746-6751 (2007) Kihara, A., J. Biol. Chem. 279 (47), 49243-49250 (2004)