

Recombinant Human FVT1 protein ab104822

1 图像

描述	
产品名称	重组人FVT1蛋白
纯度	> 90 % SDS-PAGE. ab104822 was purified using conventional chromatography techniques.
表达系统	Escherichia coli
Accession	<u>Q06136</u>
蛋白长度	Protein fragment
无动物成分	No
性质	Recombinant
种属	Human
序列	MGSSHHHHHSSGLVPRGSHMKPLALPGAHVVTGGSSGIGK CIAIECYK QGAFITLVARNEDKLLQAKKEIEMHSINDKQVVLCSVDVSQ DYNQVENV IKQAQEKLGPDMLVNCAGMAVSGKFEDLEVSTFERLMSINY LGSVYPSR AVITTMKERRVGRIVFVSSQAGQLGLFGFTAYSASKFAIRGL AEALQMEV KPYNVYITVAYPPDTPGFAEENRTKPLETRLISETTSVCK PEQVAKQI VKDAIQGNFNSSLGSD
预测分子量	29 kDa including tags
氨基酸	26 to 270
标签	His tag N-Terminus

技术指标	
Our <b>Abpromise guarantee</b> covers the use of <b>ab104822</b> in the following tested applications.	
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.	
应用	SDS-PAGE Mass Spectrometry
质谱法	MALDI-TOF
形式	Liquid

制备和贮存

稳定性和存储

Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw cycles.

pH: 8.00

Constituents: 0.00174% PMSF, 0.0154% DTT, 0.316% Tris HCl, 10% Glycerol (glycerin, glycerine), 0.58% Sodium chloride

常规信息

功能

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

组织特异性

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.

通路

Lipid metabolism; sphingolipid metabolism.

疾病相关

A chromosomal aberration involving KDSR is a cause of follicular lymphoma; also known as type II chronic lymphatic leukemia. Translocation t(2;18)(p11;q21) with a lg J kappa chain region (PubMed:8417785).

序列相似性

Belongs to the short-chain dehydrogenases/reductases (SDR) family.

细胞定位

Endoplasmic reticulum membrane.

图片



SDS-PAGE analysis of ab104822 (3µg).

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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