abcam

Product datasheet

Recombinant Human Mannose Phosphate Isomerase protein (denatured) ab111629

1 图像

描述

产品名称 重组人Mannose Phosphate Isomerase蛋白(denatured)

纯**度** > 90 % SDS-PAGE.

表达系统 Escherichia coli

Accession P34949

蛋白长度 Full length protein

无动物成分 No

性质 Recombinant

种属 Human

序列 MGSSHHHHHHSSGLVPRGSHMAAPRVFPLSCAVQQYAWGKMG

SNSEVARL

 ${\tt LASSDPLAQIAEDKPYAELWMGTHPRGDAKILDNRISQKTLS}$

QWIAENQD

 ${\tt SLGSKVKDTFNGNLPFLFKVLSVETPLSIQAHPNKELAEKLH}$

LQAPQHYP

DANHKPEMAIALTPFQGLCGFRPVEEIVTFLKTAAGNNMEDI

FGELLLQL

HQQYPGDIGCFAIYFLNLLTLKPGEAMFLEANVPHAYLKGDC

VECMACSD

NTVRAGLTPKFIDVPTLCEMLSYTPSSSKDRLFLPTRSQEDP

YLSIYDPP

VPDFTIMKTEVPGSVTEYKVLALDSASILLMVQGTVIASTPT TQTPIPLQ RGGVLFIGANESVSLKLTEPKDLLIFRACCLL

预**测分子量** 42 kDa including tags

氨基酸 1 to 362

标签 His tag N-Terminus

描述 重组人Mannose Phosphate Isomerase蛋白

技术指标

Our **Abpromise guarantee** covers the use of **ab111629** in the following tested applications.

1

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

应用 SDS-PAGE

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: 2.4% Urea, 0.32% Tris HCI

常规信息

功能 Involved in the synthesis of the GDP-mannose and dolichol-phosphate-mannose required for a

number of critical mannosyl transfer reactions.

组织特异性 Expressed in all tissues, but more abundant in heart, brain and skeletal muscle.

通路 Nucleotide-sugar biosynthesis; GDP-alpha-D-mannose biosynthesis; alpha-D-mannose 1-

phosphate from D-fructose 6-phosphate: step 1/2.

疾病相关 Defects in MPI are the cause of congenital disorder of glycosylation type 1B (CDG1B)

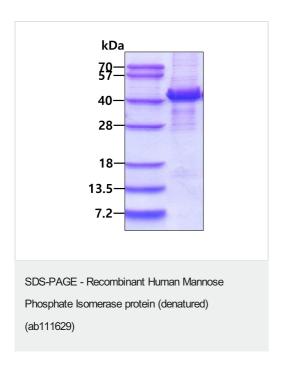
[MIM:602579]; also known as carbohydrate-deficient glycoprotein syndrome type lb (CDGS1B). Congenital disorders of glycosylation are metabolic deficiencies in glycoprotein biosynthesis that usually cause severe mental and psychomotor retardation. They are characterized by underglycosylated serum glycoproteins. CDG1B is clinically characterized by protein-losing

enteropathy.

序列相似性 Belongs to the mannose-6-phosphate isomerase type 1 family.

细胞定位 Cytoplasm.

图片



15% SDS-PAGE showing ab111629 at approximately 41.9kDa (3μg).

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