abcam

Product datasheet

Recombinant Human Mimitin protein ab109967

1 图像

描述

产品名称 重组人Mimitin蛋白

纯**度** > 85 % SDS-PAGE.

ab109967 was purified using conventional chromatography techniques.

表达系统 Escherichia coli

Accession Q8N183

蛋白长度 Full length protein

无动物成分 No

性质 Recombinant

种属 Human

序列 MGSSHHHHHHSSGLVPRGSHMGWSQDLFRALWRSLSREVK

EHVGTDQFGN

KYYYIPQYKNWRGQTIREKRIVEAANKKEVDYEAGDIPTEWE

AWIRRTRK

TPPTMEEILKNEKHREEIKIKSQDFYEKEKLLSKETSEELLP

PPVQTQIK

 ${\tt GHASAPYFGKEEPSVAPSSTGKTFQPGSWMPRDGKSHNQ}$

预**测分子量** 22 kDa including tags

氨基酸 1 to 169

标签 His tag N-Terminus

技术指标

Our <u>Abpromise guarantee</u> covers the use of ab109967 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. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.

pH: 8.00

 $Constituents: 0.0154\%\ DTT, 0.316\%\ Tris\ HCl, 20\%\ Glycerol\ (glycerin, glycerine), 0.116\%\ Sodium$

chloride

常规信息

功能 Acts as a molecular chaperone for mitochondrial complex I assembly.

组织特异性 Highly expressed in ESCC cells. Also expressed in heart, skeletal muscle, liver, and in fibroblasts.

疾病相关 Defects in NDUFAF2 are a cause of mitochondrial complex I deficiency (MT-C1D) [MIM:252010].

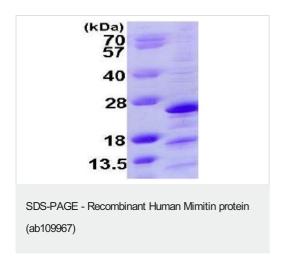
A disorder of the mitochondrial respiratory chain that causes a wide range of clinical disorders, from lethal neonatal disease to adult-onset neurodegenerative disorders. Phenotypes include macrocephaly with progressive leukodystrophy, non-specific encephalopathy, cardiomyopathy, myopathy, liver disease, Leigh syndrome, Leber hereditary optic neuropathy, and some forms of

Parkinson disease.

序列相似性 Belongs to the complex I NDUFA12 subunit family.

细胞定位 Mitochondrion.

图片



15% SDS-PAGE analysis of 3 µg ab109967.

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