

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	<u>Q8N183</u>
蛋白长度	Full length protein
无动物成分	No
性质	Recombinant
种属	Human
序列	MGSSHHHHHHSSGLVPRGSHMGWSQDLFRALWRSLSREVK EHVGTDQFGN KYYYIPQYKNWRGQTIREKRIVEAANKKEVDYEAGDIPTWE AWIRRTRK TPPTMEEILKNEKHREEIKIKSQDFYEKEKLLSKETSEELLP PPVQTQIK 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 <b>Abpromise guarantee</b> covers the use of <b>ab109967</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. 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.

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

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