

Recombinant Human Mimitin protein ab109967

画像数 1

製品の詳細

製品名	Recombinant Human Mimitin protein
精製度	> 85 % SDS-PAGE. ab109967 was purified using conventional chromatography techniques.
発現系	Escherichia coli
アクセッション番号	<u>Q8N183</u>
タンパク質長	Full length protein
Animal free	No
由来	Recombinant
生物種	Human
配列	<p>MGSSHHHHHHSSGLVPRGSHMGWSQDLFRALWRSLSREVK EHVGTDQFGN KYYYIPQYKNWRGQTIREKRIVEAANKKEVDYEAGDIPTWE AWIRRTRK TPPTMEEILKNEKHREEIKIKSQDFYEKEKLLSKETSEELL PPVQTQIK GHASAPYFGKEEPSVAPSSSTGKTFQPGSWMPRDGKSHNQ</p>
予測される分子量	22 kDa including tags
領域	1 to 169
タグ	His tag N-Terminus

特性

Our **Abpromise guarantee** 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.

アプリケーション	SDS-PAGE
	Mass Spectrometry
質量分析	MALDI-TOF
製品の状態	Liquid

前処理および保存

保存方法および安定性

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

関連情報

機能

Acts as a molecular chaperone for mitochondrial complex I assembly.

組織特異性

Highly expressed in ESCC cells. Also expressed in heart, skeletal muscle, liver, and in fibroblasts.

関連疾患

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.

配列類似性

Belongs to the complex I NDUFA12 subunit family.

細胞内局在

Mitochondrion.

画像



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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