

## **Product datasheet for TP307387M**

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## Mimitin (NDUFAF2) (NM\_174889) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human NADH dehydrogenase (ubiquinone) 1 alpha subcomplex,

assembly factor 2 (NDUFAF2), nuclear gene encoding mitochondrial protein, 100 µg

Species: Human
Expression Host: HEK293T

Expression cDNA Clone

>RC207387 representing NM\_174889

or AA Sequence: Red=Cloning site Green=Tags(s)

MGWSQDLFRALWRSLSREVKEHVGTDQFGNKYYYIPQYKNWRGQTIREKRIVEAANKKEVDYEAGDIPTE WEAWIRRTRKTPPTMEEILKNEKHREEIKIKSQDFYEKEKLLSKETSEELLPPPVQTQIKGHASAPYFGK

EEPSVAPSSTGKTFQPGSWMPRDGKSHNQ

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK

Predicted MW: 19.7 kDa

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

RefSeq: NP 777549

**Locus ID:** 91942

UniProt ID: Q8N183





RefSeq Size: 650

Cytogenetics: 5q12.1 RefSeq ORF: 507

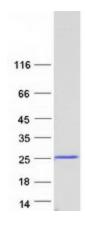
Synonyms: B17.2L; MC1DN10; mimitin; MMTN; NDUFA12L

Summary: NADH:ubiquinone oxidoreductase (complex I) catalyzes the transfer of electrons from NADH

to ubiquinone (coenzyme Q) in the first step of the mitochondrial respiratory chain, resulting in the translocation of protons across the inner mitochondrial membrane. This gene encodes a complex I assembly factor. Mutations in this gene cause progressive encephalopathy

resulting from mitochondrial complex I deficiency. [provided by RefSeq, Jul 2008]

## **Product images:**



Coomassie blue staining of purified NDUFAF2 protein (Cat# [TP307387]). The protein was produced from HEK293T cells transfected with NDUFAF2 cDNA clone (Cat# [RC207387]) using MegaTran 2.0 (Cat# [TT210002]).