

Product datasheet for TP300029M

OriGene Technologies, Inc.

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NDUFAF1 (NM_016013) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

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

assembly factor 1 (NDUFAF1), 100 µg

Species: Human
Expression Host: HEK293T

Expression cDNA Clone >RC200029 protein sequence

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

MALVHKLLRGTYFLRKFSKPTSALYPFLGIRFAEYSSSLQKPVASPGKASSQRKTEGDLQGDHQKEVALD ITSSEEKPDVSFDKAIRDEAIYHFRLLKDEIVDHWRGPEGHPLHEVLLEQAKVVWQFRGKEDLDKWTVTS DKTIGGRSEVFLKMGKNNQSALLYGTLSSEAPQDGESTRSGYCAMISRIPRGAFERKMSYDWSQFNTLYL RVRGDGRPWMVNIKEDTDFFQRTNQMYSYFMFTRGGPYWQEVKIPFSKFFFSNRGRIRDVQHELPLDKI

S

SIGFTLADKVDGPFFLEIDFIGVFTDPAHTEEFAYENSPELNPRLFK

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK
Predicted MW: 37.6 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 057097



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Locus ID: 51103

UniProt ID: Q9Y375

RefSeq Size: 1488

Cytogenetics: 15q15.1

RefSeq ORF: 981

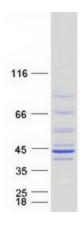
Synonyms: CGI-65; CGI65; CIA30; MC1DN11

Summary: This gene encodes a complex I assembly factor protein. Complex I (NADH-ubiquinone

oxidoreductase) 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. The encoded protein is required for assembly of complex I, and mutations in this gene are a cause of mitochondrial complex I deficiency. Alternatively spliced transcript variants have been observed for this gene, and a pseudogene of this gene is located on the long arm of chromosome 19. [provided by RefSeq,

Dec 2011]

Product images:



Coomassie blue staining of purified NDUFAF1 protein (Cat# [TP300029]). The protein was produced from HEK293T cells transfected with NDUFAF1 cDNA clone (Cat# [RC200029]) using MegaTran 2.0 (Cat# [TT210002]).