

Product datasheet for TP300223L

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

NDUFB9 (NM 005005) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant protein of human NADH dehydrogenase (ubiquinone) 1 beta subcomplex, 9,

22kDa (NDUFB9), 1 mg

Species: Human
Expression Host: HEK293T

Expression cDNA Clone >RC200223 protein sequence or AA Sequence: Red=Cloning site Green=Tags(s)

MAFLASGPYLTHQQKVLRLYKRALRHLESWCVQRDKYRYFACLMRARFEEHKNEKDMAKATQLLKEAEEE FWYRQHPQPYIFPDSPGGTSYERYDCYKVPEWCLDDWHPSEKAMYPDYFAKREQWKKLRRESWEREVKQL

QEETPPGGPLTEALPPARKEGDLPPLWWYIVTRPRERPM

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK
Predicted MW: 21.7 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.

RefSeg: NP 004996

Locus ID: 4715

UniProt ID: Q9Y6M9





RefSeq Size: 736

Cytogenetics: 8q24.13 RefSeq ORF: 537

Synonyms: B22; CI-B22; LYRM3; MC1DN24; UQOR22

Summary: The protein encoded by this gene is a subunit of the mitochondrial oxidative phosphorylation

complex I (nicotinamide adenine dinucleotide: ubiquinone oxidoreductase). Complex I is localized to the inner mitochondrial membrane and functions to dehydrogenate nicotinamide adenine dinucleotide and to shuttle electrons to coenzyme Q. Complex I deficiency is the most

common defect found in oxidative phosphorylation disorders and results in a range of conditions, including lethal neonatal disease, hypertrophic cardiomyopathy, liver disease, and

adult-onset neurodegenerative disorders. Pseudogenes of this gene are found on

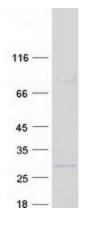
chromosomes five, seven and eight. Alternative splicing results in multiple transcript variants.

[provided by RefSeq, Jul 2015]

Protein Pathways: Alzheimer's disease, Huntington's disease, Metabolic pathways, Oxidative phosphorylation,

Parkinson's disease

Product images:



Coomassie blue staining of purified NDUFB9 protein (Cat# [TP300223]). The protein was produced from HEK293T cells transfected with NDUFB9 cDNA clone (Cat# [RC200223]) using MegaTran 2.0 (Cat# [TT210002]).

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2023 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US