

Product datasheet for TP307108

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

NDUFS6 (NM_004553) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant protein of human NADH dehydrogenase (ubiquinone) Fe-S protein 6, 13kDa

(NADH-coenzyme Q reductase) (NDUFS6), 20 µg

Species: Human
Expression Host: HEK293T

Expression cDNA Clone >RC207108 protein sequence or AA Sequence: Red=Cloning site Green=Tags(s)

MAAAMTFCRLLNRCGEAARSLPLGARCFGVRVSPTGEKVTHTGQVYDDKDYRRIRFVGRQKEVNENFAID

LIAEQPVSEVETRVIACDGGGGALGHPKVYINLDKETKTGTCGYCGLQFRQHHH

TRTRPLEQKLISEEDLAANDILDYKDDDDK**V**

Tag: C-Myc/DDK

Predicted MW: 10.8 kDa

Concentration: $>0.05 \mu g/\mu L$ 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 004544

Locus ID: 4726

UniProt ID: <u>075380</u>, <u>Q6IBC4</u>

RefSeq Size: 554



NDUFS6 (NM_004553) Human Recombinant Protein - TP307108

Cytogenetics: 5p15.33

RefSeq ORF: 372

Synonyms: CI-13kA; CI-13kD-A; CI13KDA; MC1DN9

Summary: This gene encodes a subunit of the NADH:ubiquinone oxidoreductase (complex I), which is the

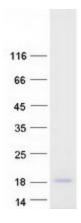
first enzyme complex in the electron transport chain of mitochondria. This complex functions in the transfer of electrons from NADH to the respiratory chain. The subunit encoded by this gene is one of seven subunits in the iron-sulfur protein fraction. Mutations in this gene cause mitochondrial complex I deficiency, a disease that causes a wide variety of clinical disorders, including neonatal disease and adult-onset neurodegenerative disorders.[provided by RefSeq,

Oct 20091

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

Parkinson's disease

Product images:



Coomassie blue staining of purified NDUFS6 protein (Cat# TP307108). The protein was produced from HEK293T cells transfected with NDUFS6 cDNA clone (Cat# [RC207108]) using MegaTran 2.0 (Cat# [TT210002]).