

Product datasheet for SC320828

NDUFS4 (NM 002495) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: NDUFS4 (NM_002495) Human Untagged Clone

Tag: Tag Free Symbol: NDUFS4

Synonyms: AQDQ; CI-18; CI-18 kDa; CI-AQDQ; MC1DN1

Mammalian Cell

Selection:

Neomycin

Vector:pCMV6-AC (PS100020)E. coli Selection:Ampicillin (100 ug/mL)

Fully Sequenced ORF: >OriGene sequence for NM_002495.1

CAAAAAAAAAAAAAAAAAAAAAAAAAAA

Restriction Sites: Please inquire ACCN: NM 002495

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

OTI Annotation: This TrueClone is provided through our Custom Cloning Process that includes sub-cloning

into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.



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NDUFS4 (NM_002495) Human Untagged Clone - SC320828

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: <u>NM 002495.1</u>, <u>NP 002486.1</u>

 RefSeq Size:
 668 bp

 RefSeq ORF:
 528 bp

 Locus ID:
 4724

 UniProt ID:
 043181

 Cytogenetics:
 5q11.2

 Domains:
 ETC_CI_21

Protein Families: Druggable Genome

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

Parkinson's disease

Gene Summary: This gene encodes an nuclear-encoded accessory subunit of the mitochondrial membrane

respiratory chain NADH dehydrogenase (complex I, or NADH:ubiquinone oxidoreductase).

Complex I removes electrons from NADH and passes them to the electron acceptor ubiquinone. Mutations in this gene can cause mitochondrial complex I deficiencies such as

Leigh syndrome. Alternative splicing results in multiple transcript variants. [provided by

RefSeq, Dec 2015]

Transcript Variant: This variant (1) encodes the longer isoform (1).