

Product datasheet for SC332503

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NDUFB3 (NM_001257102) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: NDUFB3 (NM_001257102) Human Untagged Clone

Tag: Tag Free Symbol: NDUFB3

 Synonyms:
 B12; CI-B12; MC1DN25

 Vector:
 pCMV6-Entry (PS100001)

Fully Sequenced ORF: >SC332503 representing NM_001257102.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

AAAGATAAGAAGCATCACTGA

Restriction Sites: Sgfl-Mlul

ACCN: NM_001257102

Insert Size: 297 bp

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).

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: NM 001257102.1





NDUFB3 (NM_001257102) Human Untagged Clone - SC332503

RefSeq Size: 793 bp

 RefSeq ORF:
 297 bp

 Locus ID:
 4709

 UniProt ID:
 043676

Cytogenetics: 2q33.1

Protein Families: Transmembrane

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

Parkinson's disease

MW: 11.4 kDa

Gene Summary: This gene encodes an accessory subunit of the mitochondrial membrane respiratory chain

NADH dehydrogenase (Complex I) which is the first enzyme in the electron transport chain of mitochondria. This protein localizes to the inner membrane of the mitochondrion as a single-pass membrane protein. Mutations in this gene contribute to mitochondrial complex 1 deficiency. Alternative splicing results in multiple transcript variants encoding the same protein. Humans have multiple pseudogenes of this gene. [provided by RefSeq, Mar 2012] Transcript Variant: This variant (2) represents the longer transcript. Variants 1 and 2 encode

the same protein.