

Product datasheet for RC203485L3V

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NDUFS2 (NM_004550) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: NDUFS2 (NM_004550) Human Tagged ORF Clone Lentiviral Particle

Symbol: NDUFS2

Synonyms: CI-49; MC1DN6

Mammalian Cell

Puromycin

NM 004550

Selection:

Vector:

ACCN:

pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK

ORF Size: 1389 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC203485).

Sequence:

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

RefSeg: NM 004550.3

 RefSeq Size:
 2059 bp

 RefSeq ORF:
 1392 bp

 Locus ID:
 4720

 UniProt ID:
 075306

 Cytogenetics:
 1q23.3

Domains: complex1_49Kd



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

Parkinson's disease

MW: 52.5 kDa

Gene Summary: The protein encoded by this gene is a core subunit of the mitochondrial membrane

respiratory chain NADH dehydrogenase (complex I). Mammalian mitochondrial complex I is composed of at least 43 different subunits, 7 of which are encoded by the mitochondrial genome, and the rest are the products of nuclear genes. The iron-sulfur protein fraction of complex I is made up of 7 subunits, including this gene product. Complex I catalyzes the NADH oxidation with concomitant ubiquinone reduction and proton ejection out of the mitochondria. Mutations in this gene are associated with mitochondrial complex I deficiency. Alternatively spliced transcript variants encoding different isoforms have been found for this

gene.[provided by RefSeq, Oct 2009]