

Product datasheet for RC203279L3V

OriGene Technologies, Inc.

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NDUFS7 (NM_024407) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: NDUFS7 (NM 024407) Human Tagged ORF Clone Lentiviral Particle

Symbol: NDUFS7

Synonyms: CI-20; CI-20KD; MC1DN3; MY017; PSST

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK
ACCN: NM 024407

ORF Size: 639 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC203279).

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 024407.3, NP 077718.2

 RefSeq Size:
 799 bp

 RefSeq ORF:
 642 bp

 Locus ID:
 374291

 UniProt ID:
 075251

 Cytogenetics:
 19p13.3

Domains: oxidored_q6





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

Parkinson's disease

MW: 23.6 kDa

Gene Summary: This gene encodes a protein that is a subunit of one of the complexes that forms the

mitochondrial respiratory chain. This protein is one of over 40 subunits found in complex I, the nicotinamide adenine dinucleotide (NADH):ubiquinone oxidoreductase. This complex functions in the transfer of electrons from NADH to the respiratory chain, and ubiquinone is believed to be the immediate electron acceptor for the enzyme. Mutations in this gene cause Leigh syndrome due to mitochondrial complex I deficiency, a severe neurological disorder that results in bilaterally symmetrical necrotic lesions in subcortical brain regions. [provided

by RefSeq, Jul 2008]