

Product datasheet for RC200957L1V

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Cytochrome C Oxidase subunit VIb (COX6B1) (NM_001863) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Cytochrome C Oxidase subunit VIb (COX6B1) (NM_001863) Human Tagged ORF Clone

Lentiviral Particle

Symbol: Cytochrome C Oxidase subunit VIb
Synonyms: COX6B; COXG; COXVIb1; MC4DN7

Mammalian Cell

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK
ACCN: NM_001863

ORF Size: 258 bp

ORF Nucleotide

Sequence:

The ORF insert of this clone is exactly the same as(RC200957).

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.

RefSeq: <u>NM 001863.3</u>

 RefSeq Size:
 590 bp

 RefSeq ORF:
 261 bp

 Locus ID:
 1340

 UniProt ID:
 P14854

 Cytogenetics:
 19q13.12

 Domains:
 COX6B





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Protein Pathways: Alzheimer's disease, Cardiac muscle contraction, Huntington's disease, Metabolic pathways,

Oxidative phosphorylation, Parkinson's disease

MW: 10.2 kDa

Gene Summary: Cytochrome c oxidase (COX), the terminal enzyme of the mitochondrial respiratory chain,

catalyzes the electron transfer from reduced cytochrome c to oxygen. It is a heteromeric complex consisting of 3 catalytic subunits encoded by mitochondrial genes and multiple structural subunits encoded by nuclear genes. The mitochondrially-encoded subunits function in electron transfer, and the nuclear-encoded subunits may be involved in the regulation and assembly of the complex. This nuclear gene encodes subunit VIb. Mutations in this gene are associated with severe infantile encephalomyopathy. Three pseudogenes COX6BP-1, COX6BP-2 and COX6BP-3 have been found on chromosomes 7, 17 and 22q13.1-

13.2, respectively. [provided by RefSeq, Jan 2010]