

Product datasheet for SC331067

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SDHD (NM_001276504) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: SDHD (NM_001276504) Human Untagged Clone

Tag: Tag Free Symbol: SDHD

Synonyms: CBT1; CII-4; CWS3; cybS; MC2DN3; PGL; PGL1; QPs3; SDH4

Vector: pCMV6-Entry (PS100001)

Fully Sequenced ORF: >SC331067 representing NM_001276504.

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

ATGCTGTGGAAGCTCTGA

Restriction Sites: Sgfl-Mlul

ACCN: NM_001276504

Insert Size: 363 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.



SDHD (NM_001276504) Human Untagged Clone - SC331067

RefSeq: <u>NM 001276504.1</u>

 RefSeq Size:
 1278 bp

 RefSeq ORF:
 363 bp

 Locus ID:
 6392

 UniProt ID:
 014521

 Cytogenetics:
 11q23.1

Protein Pathways: Alzheimer's disease, Citrate cycle (TCA cycle), Huntington's disease, Metabolic pathways,

Oxidative phosphorylation, Parkinson's disease

MW: 12.6 kDa

Gene Summary: This gene encodes a member of complex II of the respiratory chain, which is responsible for

the oxidation of succinate. The encoded protein is one of two integral membrane proteins anchoring the complex to the matrix side of the mitochondrial inner membrane. Mutations in this gene are associated with the formation of tumors, including hereditary paraganglioma. Transmission of disease occurs almost exclusively through the paternal allele, suggesting that

this locus may be maternally imprinted. There are pseudogenes for this gene on

chromosomes 1, 2, 3, 7, and 18. Alternative splicing results in multiple transcript variants.

[provided by RefSeq, Feb 2013]

Transcript Variant: This variant (3) lacks an alternate coding exon, but retains the reading

frame, compared to variant 1. The encoded isoform (c) is shorter than isoform a.