

Product datasheet for SC204356

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

NDUF3 (NDUFAF3) (NM_199069) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: NDUF3 (NDUFAF3) (NM_199069) Human 3' UTR Clone

Symbol: NDUF3

Synonyms: 2P1; C3orf60; E3-3; MC1DN18

Mammalian Cell

Selection:

Neomycin

Vector: pMirTarget (PS100062)

ACCN: NM_199069

Insert Size: 353 bp

Insert Sequence: >SC204356 3'UTR clone of NM_199069

The sequence shown below is from the reference sequence of NM_199069. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CTTACATCTTTGGGCCAAGCTGCTCAATGAACCGCCAGGAACTGACCTGCTGACTGCCAGGC
TTCCCAATGCTTTCACTCTTATCTACCCTTTTGGCACTTATCTTGCTTATCAACATAATATTATACAC
TTCTCCCATTTTGTATCAGGTGTGTTGCTGGCCAGGAGCTGATGGCTCACTGGGCTCTTGGAGGGGAAT
GTGAAGAAACCAAGGAGTCACTTTTTCATCTAGATTACTTAGGATTCCTTGACTTTTCAGAAGTCGGGA
AGCAGTATGTTTGCCTGTTTGTAGACCTACTTGCTCACATGCAGATTTGAGAGGACCTCAACGGCTTTTC

TCACAAAA

ACGCGTAAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul

OTI Disclaimer: Our molecular clone sequence data has been matched to the sequence identifier above as a

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).

Components: The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.





NDUF3 (NDUFAF3) (NM_199069) Human 3' UTR Clone - SC204356

RefSeq: <u>NM 199069.2</u>

Summary: This gene encodes a mitochondrial complex I assembly protein that interacts with complex I

subunits. Mutations in this gene cause mitochondrial complex I deficiency, a fatal neonatal disorder of the oxidative phosphorylation system. Alternatively spliced transcript variants

encoding different isoforms have been identified. [provided by RefSeq, Jul 2009]

Locus ID: 25915

MW: 12.9