

Product datasheet for RC209910L3

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DNAJC19 (NM_145261) Human Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: DNAJC19 (NM_145261) Human Tagged Lenti ORF Clone

Tag:Myc-DDKSymbol:DNAJC19

Synonyms: PAM18; TIM14; TIMM14

Mammalian Cell Puromycin

Selection:

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

E. coli Selection: Chloramphenicol (34 ug/mL)

ORF Nucleotide The ORF insert of this clone is exactly the same as(RC209910).

Sequence:

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





^{*} The last codon before the Stop codon of the ORF.

ACCN: NM_145261

ORF Size: 348 bp





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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.

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.

RefSeq: <u>NM 145261.2</u>

RefSeq Size: 1476 bp

RefSeq ORF: 351 bp

Locus ID: 131118

UniProt ID: Q96DA6

Cytogenetics: 3q26.33

Domains: Dnal

Protein Families: Transmembrane

MW: 12.5 kDa

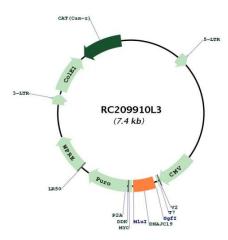
Gene Summary: The protein encoded by this gene is thought to be part of a complex involved in the ATP-

dependent transport of transit peptide-containing proteins from the inner cell membrane to the mitochondrial matrix. Defects in this gene are a cause of 3-methylglutaconic aciduria type 5 (MGA5), also known as dilated cardiomyopathy with ataxia (DCMA). Alternative splicing of this gene results in multiple transcript variants. Related pseudogenes have been identified on

chromosomes 1, 2, 6, 10, 14 and 19. [provided by RefSeq, Jan 2012]



Product images:



Circular map for RC209910L3