

Product datasheet for RC202958L3

OriGene Technologies, Inc.

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DNAJC30 (NM_032317) Human Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: DNAJC30 (NM_032317) Human Tagged Lenti ORF Clone

Tag: Myc-DDK
Symbol: DNAJC30

Synonyms: LHONAR; MC1DN38; WBSCR18

Mammalian Cell Puromycin

Selection:

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

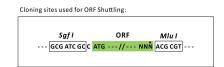
E. coli Selection: Chloramphenicol (34 ug/mL)

ORF Nucleotide The ORF insert of this clone is exactly the same as(RC202958).

Sequence:

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





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

ACCN: NM_032317

ORF Size: 678 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: NM 032317.2

RefSeq Size: 2534 bp
RefSeq ORF: 681 bp
Locus ID: 84277
UniProt ID: Q96LL9
Cytogenetics: 7q11.23

Domains: Dnal

Protein Families: Transmembrane

MW: 26.1 kDa

Gene Summary: This intronless gene encodes a member of the DNAJ molecular chaperone homology domain-

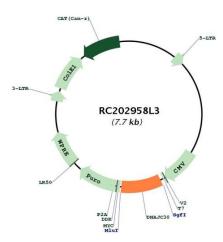
containing protein family. This gene is deleted in Williams syndrome, a multisystem

developmental disorder caused by the deletion of contiguous genes at 7q11.23. [provided by

RefSeq, Jul 2008]



Product images:



Circular map for RC202958L3