

Product datasheet for RC224428L2V

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

Hsp60 (HSPD1) (NM 002156) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Hsp60 (HSPD1) (NM_002156) Human Tagged ORF Clone Lentiviral Particle

Symbol: Hsp60

Synonyms: CPN60; GROEL; HLD4; HSP-60; HSP60; HSP65; HuCHA60; SPG13

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_002156 **ORF Size:** 1719 bp

ORF Nucleotide

OTI Disclaimer:

The ORF insert of this clone is exactly the same as(RC224428).

Sequence:

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.

RefSeg: NM 002156.4

 RefSeq Size:
 2339 bp

 RefSeq ORF:
 1722 bp

 Locus ID:
 3329

 UniProt ID:
 P10809

 Cytogenetics:
 2q33.1

Domains: cpn60_TCP1

Protein Families: Druggable Genome, Stem cell - Pluripotency





Protein Pathways: RNA degradation, Type I diabetes mellitus

MW: 60.9 kDa

ORIGENE

Gene Summary: This gene encodes a member of the chaperonin family. The encoded mitochondrial protein

may function as a signaling molecule in the innate immune system. This protein is essential for the folding and assembly of newly imported proteins in the mitochondria. This gene is adjacent to a related family member and the region between the 2 genes functions as a bidirectional promoter. Several pseudogenes have been associated with this gene. Two transcript variants encoding the same protein have been identified for this gene. Mutations associated with this gene cause autosomal recessive spastic paraplegia 13. [provided by

RefSeq, Jun 2010]