

Product datasheet for SC206500

OriGene Technologies, Inc.

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Hsp60 (HSPD1) (NM_002156) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: Hsp60 (HSPD1) (NM 002156) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

Symbol: HSPD1

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

ACCN: NM_002156

Insert Size: 492 bp

The sequence shown below is from the reference sequence of NM_002156. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

TAAAGTTAA

 ${\tt CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG}$

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.

RefSeg: NM 002156.5





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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]

Locus ID: 3329

MW: 18.3