

Product datasheet for RC223512L4V

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Caspase 1 (CASP1) (NM_001223) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Caspase 1 (CASP1) (NM_001223) Human Tagged ORF Clone Lentiviral Particle

Symbol: Caspase 1

Synonyms: ICE; IL1BC; P45

Mammalian Cell

Selection:

CII

Puromycin

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM_001223 **ORF Size:** 1149 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC223512).

Sequence:

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.

RefSeg: NM 001223.3

RefSeq Size: 1301 bp
RefSeq ORF: 1152 bp
Locus ID: 834
UniProt ID: P29466

Cytogenetics: 11q22.3

Domains: CARD, CASc, ICE_p10, ICE_p20

Protein Families: Druggable Genome, Protease





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Protein Pathways: Amyotrophic lateral sclerosis (ALS), Cytosolic DNA-sensing pathway, NOD-like receptor

signaling pathway

MW: 42.7 kDa

Gene Summary: This gene encodes a protein which is a member of the cysteine-aspartic acid protease

(caspase) family. Sequential activation of caspases plays a central role in the execution-phase of cell apoptosis. Caspases exist as inactive proenzymes which undergo proteolytic processing at conserved aspartic residues to produce 2 subunits, large and small, that dimerize to form the active enzyme. This gene was identified by its ability to proteolytically cleave and activate the inactive precursor of interleukin-1, a cytokine involved in the

processes such as inflammation, septic shock, and wound healing. This gene has been shown to induce cell apoptosis and may function in various developmental stages. Studies of a similar gene in mouse suggest a role in the pathogenesis of Huntington disease. Alternative splicing results in transcript variants encoding distinct isoforms. [provided by RefSeq, Mar

2012]