

Product datasheet for RC207905L2V

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

alpha 2 Macroglobulin (A2M) (NM 000014) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: alpha 2 Macroglobulin (A2M) (NM_000014) Human Tagged ORF Clone Lentiviral Particle

Symbol: alpha 2 Macroglobulin

Synonyms: A2MD; CPAMD5; FWP007; S863-7

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_000014 **ORF Size:** 4422 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC207905).

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 000014.4

RefSeq Size: 4678 bp RefSeq ORF: 4425 bp

Locus ID: 2

 UniProt ID:
 P01023

 Cytogenetics:
 12p13.31

 Domains:
 A2M, A2M_N

Protein Families: Druggable Genome, Secreted Protein





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Protein Pathways: Complement and coagulation cascades

MW: 163.3 kDa

Gene Summary: The protein encoded by this gene is a protease inhibitor and cytokine transporter. It uses a

bait-and-trap mechanism to inhibit a broad spectrum of proteases, including trypsin, thrombin and collagenase. It can also inhibit inflammatory cytokines, and it thus disrupts inflammatory cascades. Mutations in this gene are a cause of alpha-2-macroglobulin deficiency. This gene is implicated in Alzheimer's disease (AD) due to its ability to mediate the clearance and degradation of A-beta, the major component of beta-amyloid deposits. A related pseudogene, which is also located on the p arm of chromosome 12, has been

identified. [provided by RefSeq, Nov 2016]