

## Product datasheet for RC215901L1V

## OriGene Technologies, Inc.

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## ADAMTS10 (NM\_030957) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** ADAMTS10 (NM\_030957) Human Tagged ORF Clone Lentiviral Particle

Symbol: ADAMTS10

Synonyms: ADAM-TS10; ADAMTS-10; WMS; WMS1

Mammalian Cell

Selection:

None

**Vector:** pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK

**ACCN:** NM\_030957 **ORF Size:** 3309 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC215901).

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 030957.2

 RefSeq Size:
 4237 bp

 RefSeq ORF:
 3312 bp

 Locus ID:
 81794

 UniProt ID:
 Q9H324

Cytogenetics: 19p13.2

Domains: tsp\_1, Reprolysin, Pep\_M12B\_propep

Protein Families: Druggable Genome, Secreted Protein







MW: 120.7 kDa

Gene Summary:

This gene belongs to the ADAMTS (a disintegrin and metalloproteinase domain with thrombospondin type-1 motifs) family of zinc-dependent proteases. ADAMTS proteases are complex secreted enzymes containing a prometalloprotease domain of the reprolysin type attached to an ancillary domain with a highly conserved structure that includes at least one thrombospondin type 1 repeat. They have been demonstrated to have important roles in connective tissue organization, coagulation, inflammation, arthritis, angiogenesis and cell migration. The product of this gene plays a major role in growth and in skin, lens, and heart development. It is also a candidate gene for autosomal recessive Weill-Marchesani syndrome. [provided by RefSeq, Jul 2008]