

Product datasheet for SC208496

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OriGene Technologies, Inc.

ADAMTS10 (NM_030957) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: ADAMTS10 (NM_030957) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

Symbol: ADAMTS10

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

ACCN: NM_030957

Insert Size: 683 bp

Insert Sequence: >SC208496 3'UTR clone of NM_030957

The sequence shown below is from the reference sequence of NM_030957. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

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.





ADAMTS10 (NM_030957) Human 3' UTR Clone - SC208496

RefSeq: <u>NM 030957.4</u>

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]

Locus ID: 81794 **MW:** 23.6