

## Product datasheet for SC328718

## OriGene Technologies, Inc.

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## Repulsive Guidance Molecule A (RGMA) (NM 001166288) Human Untagged Clone

**Product data:** 

**Product Type: Expression Plasmids** 

**Product Name:** Repulsive Guidance Molecule A (RGMA) (NM\_001166288) Human Untagged Clone

Tag: Tag Free

Symbol: Repulsive Guidance Molecule A

Synonyms: **RGM Mammalian Cell** 

Selection:

None

Vector: pCMV6-XL5

E. coli Selection: Ampicillin (100 ug/mL)

>NCBI ORF sequence for NM\_001166288, the custom clone sequence may differ by one or **Fully Sequenced ORF:** 

more nucleotides

ATGGGTATGGGGAGAGGGCAGGACGTTCAGCCCTGGGATTCTGGCCGACCCTCGCCTTC CTTCTCTGCAGCTTCCCCGCAGCCACCTCCCCGTGCAAGATCCTCAAGTGCAACTCTGAG TTCTGGAGCGCCACGTCGGGCAGCCACGCCCCAGCCTCAGACGACACCCCCGAGTTCTGT GCAGCCTTGCGCAGCTACGCCCTGTGCACGCGGCGGACGGCCCGCACCTGCCGGGGTGAC CTGGCCTACCACTCGGCCGTCCATGGCATAGAGGACCTCATGAGCCAGCACAACTGCTCC GAGCGCTCGGACAGCCCCGAGATCTGCCATTACGAGAAGAGCTTTCACAAGCACTCGGCC ACCCCAACTACACGCACTGTGGCCTCTTCGGGGACCCACACCTCAGGACTTTCACCGAC CGCTTCCAGACCTGCAAGGTGCAGGGCGCCTGGCCGCTCATCGACAATAATTACCTGAAC GTGCAGGTCACCAACACGCCTGTGCTGCCCGGCTCAGCGGCCACTGCCACCAGCAAGCTC ACCATCATCTTCAAGAACTTCCAGGAGTGTGTGGACCAGAAGGTGTACCAGGCTGAGATG GACGAGCTCCCGGCCGCCTTCGTGGATGGCTCTAAGAACGGTGGGGACAAGCACGGGGCC AACAGCCTGAAGATCACTGAGAAGGTGTCAGGCCAGCACGTGGAGATCCAGGCCAAGTAC ATCGGCACCACCATCGTGGTGCGCCAGGTGGGCCGCTACCTGACCTTTGCCGTCCGCATG CCAGAGGAAGTGGTCAATGCTGTGGAGGACTGGGACAGCCAGGGTCTCTACCTCTGCCTG CGGGGCTGCCCCTCAACCAGCAGATCGACTTCCAGGCCTTCCACACCAATGCTGAGGGC ACCGGTGCCCGCAGGCTGCCAGCCCTGCACCCACAGCCCCCGAGACCTTCCCA TACGAGACAGCCGTGGCCAAGTGCAAGGAGAAGCTGCCGGTGGAGGACCTGTACTACCAG GCCTGCGTCTTCGACCTCCTCACCACGGGCGACGTGAACTTCACACTGGCCGCCTACTAC GCGTTGGAGGATGTCAAGATGCTCCACTCCAACAAAGACAAACTGCACCTGTATGAGAGG GGCGCCCTCGTCCCGCTCCTGGCCCTGCTCCCTGTGTTCTGCTAG

**Restriction Sites:** Please inquire



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**ACCN:** NM\_001166288

**OTI Disclaimer:** Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

**OTI Annotation:** This TrueClone is provided through our Custom Cloning Process that includes sub-cloning

into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

**Components:** The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

**Reconstitution Method:** 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

**RefSeq:** <u>NM 001166288.1</u>, <u>NP 001159760.1</u>

 RefSeq Size:
 3058 bp

 RefSeq ORF:
 1305 bp

 Locus ID:
 56963

 UniProt ID:
 Q96B86

 Cytogenetics:
 15q26.1

**Gene Summary:** This gene encodes a member of the repulsive guidance molecule family. The encoded protein

is a glycosylphosphatidylinositol-anchored glycoprotein that functions as an axon guidance protein in the developing and adult central nervous system. This protein may also function as a tumor suppressor in some cancers. Alternate splicing results in multiple transcript variants.

[provided by RefSeq, Oct 2009]

Transcript Variant: This variant (5) differs in the 5' UTR, lacks a portion of the 5' coding region, and initiates translation at a downstream start codon, compared to variant 1. The encoded isoform (2) has a shorter N-terminus, compared to isoform 1. Variants 2, 3, 5 and 6 encode the same isoform (2). Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript

alignments.