

Product datasheet for RC200201L4V

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

RBM10 (NM_152856) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: RBM10 (NM_152856) Human Tagged ORF Clone Lentiviral Particle

Symbol: RBM10

Synonyms: DXS8237E; GPATC9; GPATCH9; S1-1; TARPS; ZRANB5

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

ACCN: NM_152856 **ORF Size:** 2556 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC200201).

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 152856.1

RefSeq Size: 3178 bp
RefSeq ORF: 2559 bp
Locus ID: 8241
UniProt ID: P98175
Cytogenetics: Xp11.3

Domains: G-patch, RRM, zf-RanBP, zf-C2H2

Protein Families: Druggable Genome





ORIGENE

MW: 94.4 kDa

Gene Summary: This gene encodes a nuclear protein that belongs to a family proteins that contain an RNA-

binding motif. The encoded protein associates with hnRNP proteins and may be involved in regulating alternative splicing. Defects in this gene are the cause of the X-linked recessive disorder, TARP syndrome. Alternate splicing results in multiple transcript variants.[provided

by RefSeq, Mar 2011]