

## Product datasheet for AR09930PU-L

## OriGene Technologies, Inc.

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## RBM8A / RBM8 (1-174, His-tag) Human Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** RBM8A / RBM8 (1-174, His-tag) human recombinant protein, 0.25 mg

Species: Human
Expression Host: E. coli

**Expression cDNA Clone** MADVLDLHEA GGEDFAMDED GDESIHKLKE KAKKRKGRGF GSEEGSRARM REDYDSVEQD

or AA Sequence: GDEPGPQRSV EGWILFVTGV HEEATEEDIH DKFAEYGEIK NIHLNLDRRT GYLKGYTLVE

YETYKEAQAA MEGLNGQDLM GQPISVDWCF VRGPPKGKRR GGRRRSRSPD RRRRLEHHHH HH

Tag: His-tag

Predicted MW: 20.9 kDa

Concentration: lot specific

Purity: >90%

**Buffer:** Presentation State: Purified

State: Liquid purified protein

Buffer System: 20 mM Tris-HCl buffer (pH 8.0) containing 10% glycerol, 0.1M NaCl, 1 mM DTT

**Preparation:** Liquid purified protein

**Protein Description:** Recombinant human RBM8A protein, fused to His-tag at C-terminus, was expressed in E.coli

and purified by using conventional chromatography.

Storage: Store undiluted at 2-8°C for up to two weeks or (in aliquots) at -20°C or -70°C for longer.

Avoid repeated freezing and thawing.

**Stability:** Shelf life: one year from despatch.

**RefSeq:** NP 005096

 Locus ID:
 9939

 UniProt ID:
 Q9Y5S9

 Cytogenetics:
 1q21.1

Synonyms: RNA-binding protein 8A, BOV-1, HSPC114, Y14, MDS014





**Summary:** 

This gene encodes a protein with a conserved RNA-binding motif. The protein is found predominantly in the nucleus, although it is also present in the cytoplasm. It is preferentially associated with mRNAs produced by splicing, including both nuclear mRNAs and newly exported cytoplasmic mRNAs. It is thought that the protein remains associated with spliced mRNAs as a tag to indicate where introns had been present, thus coupling pre- and post-mRNA splicing events. Previously, it was thought that two genes encode this protein, RBM8A and RBM8B; it is now thought that the RBM8B locus is a pseudogene. There are two alternate translation start codons with this gene, which result in two forms of the protein. An allele mutation and a low-frequency noncoding single-nucleotide polymorphism (SNP) in this gene cause thrombocytopenia-absent radius (TAR) syndrome. [provided by RefSeq, Jul 2013]

**Protein Families:** Druggable Genome

**Protein Pathways:** Spliceosome

## **Product images:**

