

RBM10 Polyclonal Antibody

Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP57653

Product Information

Application IHC-P, IHC-F, IF, ICC, E

Primary Accession P98175

Reactivity Rat, Pig, Bovine

Host Rabbit Clonality Polyclonal Calculated MW 103533 **Physical State** Liquid

Immunogen KLH conjugated synthetic peptide derived from human RBM10

101-200/930 **Epitope Specificity**

Isotype IgG

affinity purified by Protein A **Purity**

Buffer 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

SUBCELLULAR LOCATION Nucleus. In the extranucleolar nucleoplasm constitutes hundreds of nuclear

domains, which dynamically change their structures in a reversible manner. Upon globally reducing RNA polymerase II transcription, the nuclear bodies enlarge and decrease in number. They occur closely adjacent to nuclear speckles or IGCs (interchromatin granule clusters) but coincide with TIDRs. Contains 1 C2H2-type zinc finger. Contains 1 G-patch domain. Contains 1

SIMILARITY

RanBP2-type zinc finger. Contains 2 RRM (RNA recognition motif) domains. Associates with the spliceosome. Component of a large chromatin remodeling

SUBUNIT

complex, at least composed of MYSM1, PCAF,RBM10 and KIF11/TRIP5.

Post-translational modifications

Defects in RBM10 are the cause of TARP syndrome (TARPS) [MIM:311900]. It is **DISEASE**

a disorder characterized by the Robin sequence (micrognathia, glossoptosis

and cleft palate), talipes equinovarus and cardiac defects.

Phosphorylated upon DNA damage, probably by ATM or ATR.

This product as supplied is intended for research use only, not for use in **Important Note**

human, therapeutic or diagnostic applications.

This gene encodes a nuclear protein that belongs to a family proteins that **Background Descriptions**

> 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]

Additional Information

Gene ID 8241

Other Names RNA-binding protein 10, G patch domain-containing protein 9, RNA-binding

motif protein 10, RNA-binding protein S1-1, S1-1, RBM10 (HGNC:9896)

Dilution IHC-P=1:100-500,IHC-F=1:100-500,ICC=1:100-500,IF=1:100-500,ELISA=1:5000-

10000

Format 0.01 M TBS(pH7.4) with 1% BSA, 0.09% (W/V) sodium azide and 50% Glyce

Storage Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When

reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody

is stable for at least two weeks at 2-4 °C.

Protein Information

Name RBM10 (<u>HGNC:9896</u>)

Function Binds to ssRNA containing the consensus sequence 5'-AGGUAA-3'

(PubMed:<u>21256132</u>). May be involved in post-transcriptional processing, most probably in mRNA splicing (PubMed:<u>18315527</u>). Binds to RNA homopolymers, with a preference for poly(G) and poly(U) and little for poly(A) (By similarity).

May bind to specific miRNA hairpins (PubMed: 28431233).

Cellular Location Nucleus. Note=In the extranucleolar nucleoplasm constitutes hundreds of

nuclear domains, which dynamically change their structures in a reversible manner. Upon globally reducing RNA polymerase II transcription, the nuclear bodies enlarge and decrease in number. They occur closely adjacent to nuclear speckles or IGCs (interchromatin granule clusters) but coincide with

TIDRs (transcription-inactivation-dependent RNA domains)

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