Product datasheet
Anti-RBM41 Antibody
Catalog Number: A14311-1



Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

Web: www.boster.com Phone: 027-67845390/1/2 Email: boster@boster.com

<b>Basic Information</b>	
Product Name	Anti-RBM41 Antibody
Gene Name	RBM41
Source	Rabbit
Clonality	Polyclonal
Isotype	IgG
Species Reactivity	human, mouse, rat
Tested Application	WB, FCM, ELISA
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.
Immunogen	E.coli-derived human RBM41 recombinant protein (Position: E130-F384).
Concentration	500 ug/ml
Purification	Immunogen affinity purified.
Observed MW	47 kDa
Dilution Ratios	Western blot (WB): 1:500-2000 Flow Cytometry (Fixed): 1:50-200 Enzyme linked immunosorbent assay (ELISA):1:100-1000

## **Storage**

12 months from date of receipt, -20°C as supplied.

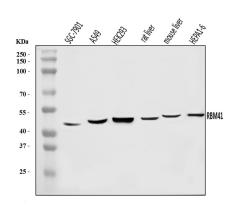
## **Background Information**

RBM41 is an RNA-binding protein of unknown specific function. The human gene is mapped to Xq22.3.

## **Selected Validation Data**

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Western blot analysis of RBM41 using anti-RBM41 antibody (A14311-1). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: SGC-7901 whole cell lysates,

Lane 2: A549 whole cell lysates,

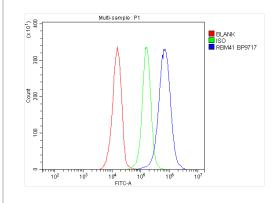
Lane 3: HEK293 whole cell lysates,

Lane 4: rat liver tissue lysates,

Lane 5: mouse liver tissue lysates,

Lane 6: HEPA1-6 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-RBM41 antigen affinity purified polyclonal antibody (A14311-1) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for RBM41 at approximately 47 kDa. The expected band size for RBM41 is at 47 kDa.



Flow Cytometry analysis of Jurkat cells using anti-RBM41 antibody (A14311-1).

Overlay histogram showing Jurkat cells stained with A14311-1 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-RBM41 Antibody (A14311-1) at 1:100 dilution for 30 min at 20°C. Fluoro488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.