bs-3833R

- DATASHEET -

[Primary Antibody]

PABP Rabbit pAb



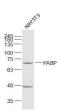
www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn 400-901-9800

DATAGHEET		
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		IHC-P (1:100-500)
GenelD: 26986	SWISS: P11940	IHC-F (1:100-500) IF (1:100-500)
Target: PABP		Flow-Cyt (1ug/Test) ICC/IF (1:50)
Immunogen: KLH conjugated synthetic peptide derived from human PABP: 101-200/636.		Reactivity: Human, Mouse (predicted: Rat, Rabbit, Pig, Horse)
Purification: affinity purified by Protein A		
Concentration: 1mg/ml		
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Predicted 70 kDa MW.: ^{70 kDa} Subcellular Location: ^{Cytoplasm} ,Nucleus
Background: This gene encodes a poly(A) binding protein. The protein shuttles between the nucleus and cytoplasm and binds to the 3' poly(A) tail of eukaryotic messenger RNAs via RNA-recognition motifs. The binding of this protein to poly(A) promotes ribosome recruitment and translation initiation; it is also required for poly(A) shortening which is the first step in mRNA decay. The gene is part of a small		

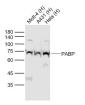
gene family including three protein-coding genes and several

pseudogenes.[provided by RefSeq, Aug 2010].

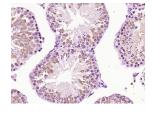
- VALIDATION IMAGES -



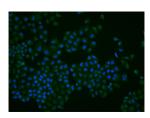
Sample: NIH/3T3(Mouse) Cell Lysate at 30 ug Primary: Anti-PABP (bs-3833R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 70 kD Observed band size: 70 kD



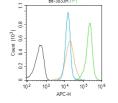
Sample: Lane 1: Molt-4 (Human) Cell Lysate at 30 ug Lane 2: A431 (Human) Cell Lysate at 30 ug Lane 3: Hela (Human) Cell Lysate at 30 ug Primary: Anti-PABP (bs-3833R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 70 kD Observed band size: 72 kD



Paraformaldehyde-fixed, paraffin embedded (Mouse testis); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (PABP) Polyclonal Antibody, Unconjugated (bs-3833R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Hela cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (PABP) polyclonal Antibody, Unconjugated (bs-3833R) 1:50, 90 minutes at 37°C; followed by a conjugated Goat



Blank control (Black line):Molt4 (Black). Primary Antibody (green line): Rabbit Anti-PABP antibody (bs-3833R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat antirabbit IgG-AF647 Dilution: 1µg /test. Protocol Anti-Rabbit IgG antibody at 37° C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.

The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at room temperature. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.