## **PABPN1 Recombinant Antibody**

## RAC00333



Product Information	
Size:	Protein Background:
50ul	Involved in the 3'-end formation of mRNA precursors (pre-mRNA) by the addition of a
Reactivity:	poly(A) tail of 200-250 nt to the upstream cleavage product (By similarity). Stimulates poly(A) polymerase (PAPOLA) conferring processivity on the poly(A) tail elongation
Human	reaction and controls also the poly(A) tail length (By similarity). Increases the affinity of poly(A) polymerase for RNA (By similarity). Is also present at various stages of mRNA metabolism including nucleocytoplasmic trafficking and nonsense-mediated decay (NMD) of mRNA. Cooperates with SKIP to synergistically activate E-box-mediated transcription through MYOD1 and may regulate the expression of muscle-specific genes . Binds to poly(A) and to poly(G) with high affinity (By similarity). May protect the poly(A) tail from degradation (By similarity). <b>Gene ID:</b> PABPN1
Source:	
Homo sapiens (Human)	
lsotype:	
Rabbit lgG	
Applications:	
ELISA, WB, IHC, IF, FC	Uniprot
Recommended dilutions:	Q86U42
WB:1:500-1:5000, IHC:1:50-1:200, IF:1:20- 1:200, FC:1:20-1:200	Synonyms:
	Polyadenylate-binding protein 2 (PABP-2) (Poly(A)-binding protein 2) (Nuclear poly(A)- binding protein 1) (Poly(A)-binding protein II) (PABII) (Polyadenylate-binding nuclear protein 1), PABPN1, PAB2 PABP2
	Immunogen:
	A synthesized peptide derived from human PABPN1.

## Storage:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.



IHC image of RACO0333 diluted at 1:100 and staining in paraffinembedded human testis tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a Goat antirabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.

IHC image of RACO0333 diluted at 1:100 and staining in paraffinembedded human bladder cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a Goat antirabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.

## Western Blot

Positive WB detected in(293 whole cell lysate) MCF-7 whole cell lysate) Raji whole cell lysate) HepG2 whole cell lysate) All lanes: PABPN1 antibody at 1:2000 Secondary Goat polyclonal to rabbit IgG at 1:50000 dilution

Predicted band size: 33, 32, 38 kDa Observed band size: 50 kDa