

## **Product Data Sheet**

### **Anti-PABPC4** Antibody

Catalog #	Source	Reactivity	Applications
CQA3138	Rabbit	H, M, R	WB, IH, IF/IC
Description	Rabb	oit polyclonal antibody t	o PABPC4
Immunogen	KLH-	conjugated synthetic pe	eptide of human PABPC4
Purification	The	antibody was purified b	y immunogen affinity chromatography.
Specificity	Reco	ognizes endogenous leve	els of PABPC4 protein.
Clonality	Poly	clonal	
Conjugation			
Form	Liqu	id in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1/500 - 1/2000) <i>,</i> IH (1/50	- 1/200), IF/IC (1/50 - 1/200)
Gene Symbol	PABI	PC4	
Alternative Na	ames APP:	1; PABP4; Polyadenylate	-binding protein 4; PABP-4; Poly(A)-binding protein 4;
	Activ	vated-platelet protein 1	APP-1; Inducible poly(A)-binding protein; iPABP
Entrez Gene	8761	(Human)	
SwissProt	Q13	310 (Human)	
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid
	freez	ze/thaw cycles.	

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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## **Product Data Sheet**

kDa
A
B

130
95

95
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70
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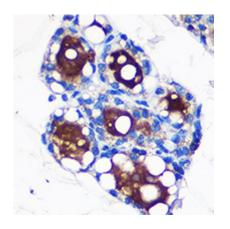
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38
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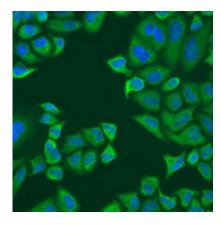
28
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19
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Western blot analysis of PABPC4 expression in Jurkat (A), mouse heart (B) whole cell lysates. (Predicted band size: 69; 70; 72 kD; Observed band size: 79 kD)



Immunohistochemical analysis of PABPC4 staining in human brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of PABPC4 staining in U2OS cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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