

# **Product Data Sheet**

# Anti-elF4Al Antibody

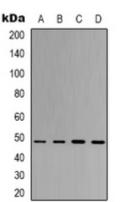
Catalog #	Source	Reactivity	Applications	
CPA9155	Mouse	H, M, R	WB, IH, IF/IC	
Description	М	louse monoclonal antibod	y to eIF4AI	
Immunogen	KL	H-conjugated synthetic pe	eptide encompassing a sequence of human eIF4AI. The	
	ex	act sequence is proprieta	ту.	
Purification				
Specificity	Re	ecognizes endogenous leve	els of eIF4AI protein.	
Clonality	М	Ionoclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ar	nd 0.01% sodium azide.		
Dilution	W	/B (1/1000 - 1/3000), IH (1/1	00 - 1/200), IF/IC (1/100 - 1/200)	
Gene Symbol	EI	F4A1		
Alternative Na	ames DI	DX2A; EIF4A; Eukaryotic in	itiation factor 4A-I; eIF-4A-I; eIF4A-I; ATP-dependent	
	RI	NA helicase eIF4A-1		
Entrez Gene	19	973 (Human); 13681 (Mou	se)	
SwissProt	PE	50842 (Human); P60843 (N	/louse)	
Storage/Stabi	<b>lity</b> Sh	nipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
	fre	eeze/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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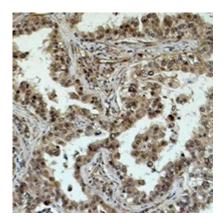




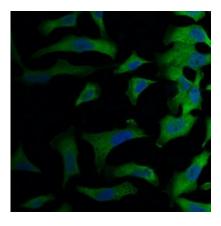
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Western blot analysis of eIF4AI expression in 293T (A), Hela (B), HepG2 (C), mouse brain (D) whole cell lysates. (Predicted band size: 46 kD; Observed band size: 46 kD)



Immunohistochemical analysis of eIF4AI staining in human lung cancer 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 eIF4AI staining in Hela 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 hidified chamber. Cells were washed with PBST and incubated with a FITC-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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