

# **Product Data Sheet**

# Anti-elF3C Antibody

Catalog #	Source	Reactivity	Applications		
CQA1516	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Ra	abbit polyclonal antibody	to eIF3C		
Immunogen	Re	ecombinant full length pro	tein of human eIF3C		
Purification	Tł	ne antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous lev	els of eIF3C protein.		
Clonality	Pc	olyclonal			
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 (1/500 - 1/2000), IH (1/50	) - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	EI	F3C			
Alternative Na	ames El	F3S8; Eukaryotic translation	on initiation factor 3 subunit C; eIF3c; Eukaryotic		
	tra	anslation initiation factor	3 subunit 8; eIF3 p110		
Entrez Gene	86	563 (Human); 56347 (Mou	se); 293484 (Rat)		
SwissProt	Q	99613 (Human); Q8R1B4	Mouse); B5DFC8 (Rat)		
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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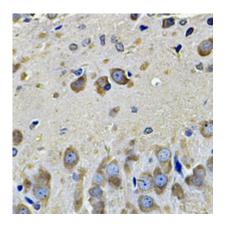
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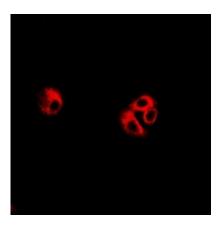
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Western blot analysis of eIF3C expression in HepG2 (A), SW620 (B), mouse testis (C) whole cell lysates. (Predicted band size: 104; 105 kD; Observed band size: 105 kD)



Immunohistochemical analysis of eIF3C staining in rat 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 eIF3C 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 hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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