

Product Data Sheet

Anti-CstF-64 Antibody

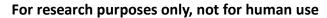
Catalog #	Source	Reactivity	Applications		
CPA1290	Rabbit	H, M, R, B, Z	WB, IH, IF/IC		
Description	Ral	bbit polyclonal antibody t	o CstF-64		
Immunogen	KLł	-conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	reg	ion of human CstF-64. Th	e exact sequence is proprietary.		
Purification	The	e antibody was purified by	v immunogen affinity chromatography.		
Specificity	Red	cognizes endogenous leve	ls of CstF-64 protein.		
Clonality	Pol	lyclonal			
Conjugation					
Form	Liq	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	d 0.01% sodium azide.			
Dilution	WE	3 (1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	CST	TF2			
Alternative Na	ames Cle	avage stimulation factor	ubunit 2; CF-1 64 kDa subunit; Cleavage stimulation		
	fac	tor 64 kDa subunit; CSTF	64 kDa subunit; CstF-64		
Entrez Gene	147	78 (Human); 108062 (Mou	ise)		
SwissProt	P33	3240 (Human); Q8BIQ5 (N	1ouse)		
Storage/Stabi	lity Shi	pped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	fre	eze/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

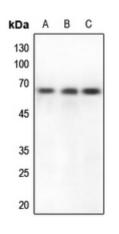
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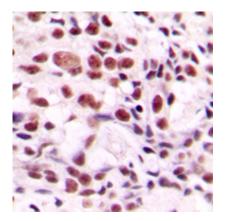
Coherion



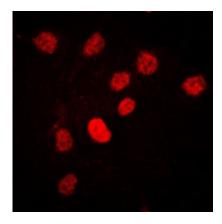
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Western blot analysis of CstF-64 expression in mouse spleen (A), rat spleen (B), rat brain (C) whole cell lysates. (Predicted band size: 60 kD; Observed band size: 64 kD)



Immunohistochemical analysis of CstF-64 staining in human breast 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 CstF-64 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 humidified 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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