

Product Data Sheet

Anti-BRF1 Antibody

Catalog #	Source	Reactivity	Applications	
CPA1518	Rabbit	H, M, R	WB, IH, IF/IC	
Description		Rabbit polyclonal antibody t	o BRF1	
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center		
		region of human BRF1. The exact sequence is proprietary.		
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of BRF1 protein.	
Clonality		Polyclonal		
Conjugation				
Form Liquid in 0.42% Potassium		Liquid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/1000), IH (1/10	0 - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol		BRF1		
Alternative Na	ames	BRF; GTF3B; TAF3B2; TAF3C;	Transcription factor IIIB 90 kDa subunit; TFIIIB90;	
		hTFIIIB90; B-related factor 1	; BRF-1; hBRF; TAF3B2; TATA box-binding	
		protein-associated factor, RI	NA polymerase III, subunit 2	
Entrez Gene		2972 (Human)		
SwissProt		Q92994 (Human); Q8CFK2 (Mouse)	
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
		freeze/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

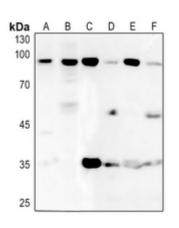
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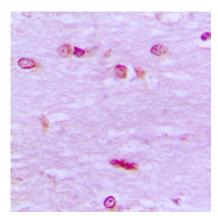
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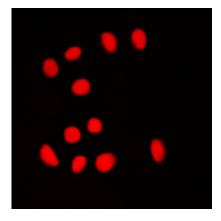
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Western blot analysis of BRF1 expression in A549 (A), H1688 (B), mouse muscle (C), mouse testis (D), rat muscle (E), rat testis (F) whole cell lysates. (Predicted band size: 73 kD; Observed band size: 90 kD)



Immunohistochemical analysis of BRF1 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 BRF1 staining in NIH3T3 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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