

## **Product Data Sheet**

## **Anti-UBF Antibody**

Catalog #	Source	Reactivity	Applications	
CPA7067	Rabbit	H, M, R	WB, IH	
Description		Rabbit polyclonal antibody t	o UBF	
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the center	
		region of human UBF. The ex	act sequence is proprietary.	
Purification		The antibody was purified by	/ immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	ls of UBF protein.	
Clonality Polyclonal				
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 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)	
Gene Symbol		UBTF		
Alternative Na	imes	UBF; UBF1; Nucleolar transc	ription factor 1; Autoantigen NOR-90; Upstream-binding	
		factor 1; UBF-1		
Entrez Gene		7343 (Human); 25574 (Rat)		
SwissProt		P17480 (Human); P25976 (N	louse); P25977 (Rat)	
Storage/Stabil	lity	Shipped at 4°C. Upon deliver	y 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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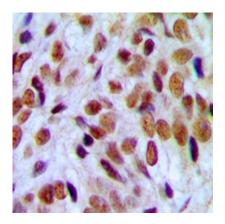
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# Coherion

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Western blot analysis of UBF expression in Jurkat (A), rat muscle (B) whole cell lysates. (Predicted band size: 89 kD; Observed band size: 97 kD)



Immunohistochemical analysis of UBF 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.

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