

Product Data Sheet

Anti-ZNF148 Antibody

Catalog #	Source	Reactivity	Applications	
CPA4320	Rabbit	Н, В	WB, IH, IF/IC	
		·		
Description	F	Rabbit polyclonal antibody	to ZNF148	
Immunogen	ł	KLH-conjugated synthetic p	eptide encompassing a sequence within the N-term	
	r	region of human ZNF148. T	he exact sequence is proprietary.	
Purification	٦	The antibody was purified b	by immunogen affinity chromatography.	
Specificity	F	Recognizes endogenous lev	els of ZNF148 protein.	
Clonality	ł	Polyclonal		
Form	l	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ä	and 0.01% sodium azide.		
Dilution	١	WB (1/500 - 1/1000), IH (1/1	00 - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	Z	ZNF148		
Alternative N	ames Z	ZBP89; Zinc finger protein 2	.48; Transcription factor ZBP-89; Zinc finger DNA-binding	
	ł	protein 89		
Entrez Gene	7	7707 (Human)		
SwissProt	(Q9UQR1 (Human)		
Storage/Stabi	ility S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
	f	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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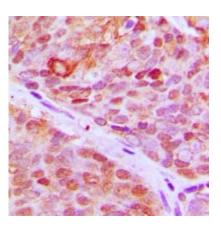
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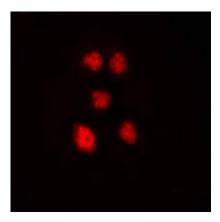
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Western blot analysis of ZNF148 expression in HeLa (A), HepG2 (B) whole cell lysates.



Immunohistochemical analysis of ZNF148 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of ZNF148 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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