

Product Data Sheet

Anti-HSP27 (Phospho-S78) Antibody

Catalog #	Source	e Reactivity	Applications		
CPA6040	Rabbit	t H, R	WB, IH		
Description		Rabbit polyclonal antibody to HSP27 (Phospho-S78)			
Immunogen		KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding			
	S78 of human HSP27 protein. The exact sequence is proprietary.				
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	els of HSP27 protein only when phosphorylated at S78.		
Clonality		Polyclonal			
Conjugation					
Form	m Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerc				
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/50	- 1/200)		
Gene Symbol		HSPB1			
Alternative N	ames	HSP27; HSP28; Heat shock p	rotein beta-1; HspB1; 28 kDa heat shock protein;		
		Estrogen-regulated 24 kDa p	rotein; Heat shock 27 kDa protein; HSP 27;		
		Stress-responsive protein 27	; SRP27		
Entrez Gene		3315 (Human); 24471 (Rat)			
SwissProt		P04792 (Human); P42930 (F	at)		
Storage/Stabi	ility	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

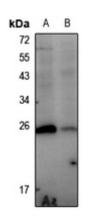
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Western blot analysis of HSP27 (Phospho-S78) expression in Hela (A), rat spleen (B) whole cell lysates. (Predicted band size: 22 kD; Observed band size: 27 kD)



Immunohistochemical analysis of HSP27 (Phospho-S78) 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.

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