

Product Data Sheet

Anti-SPARC Antibody

Catalog #	Source	Reactivity	Applications		
CPA3016	Rabbit	H, M, R, B, S, Z	WB, IH		
Description	Ra	abbit polyclonal antibody to	SPARC		
Immunogen	KL	H-conjugated synthetic pep.	tide encompassing a sequence within the center		
	re	gion of human SPARC. The e	xact sequence is proprietary.		
Purification	Th	ne antibody was purified by i	mmunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous levels	of SPARC protein.		
Clonality	Pc	blyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	nd 0.01% sodium azide.			
Dilution	W	'B (1/500 - 1/1000), IH (1/100	- 1/200)		
Gene Symbol	SP	PARC			
Alternative N	ames Of	N; SPARC; Basement-membr	ane protein 40; BM-40; Osteonectin; ON; Secreted		
	pr	otein acidic and rich in cyste	ine		
Entrez Gene	66	578 (Human); 20692 (Mouse); 24791 (Rat)		
SwissProt	PC)9486 (Human); P07214 (Mc	use); P16975 (Rat)		
Storage/Stabi	lity Sh	nipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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

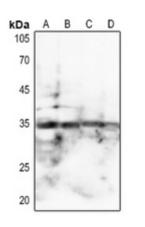
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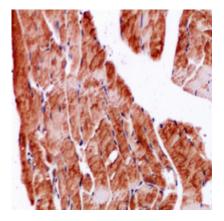




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Western blot analysis of SPARC expression in A549 (A), mouse kidney (B), mouse brain (C), rat brain (D) whole cell lysates. (Predicted band size: 34 kD; Observed band size: 34 kD)



Immunohistochemical analysis of SPARC staining in mouse muscle 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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