

Product Data Sheet

Anti-Protein C Antibody

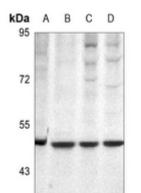
Catalog #	Source	Reactivity	Applications		
CPA4174	Rabbit	H, M, R	WB, IH		
Description	F	Rabbit polyclonal antibody to Protein C			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
	r	region of human Protein C. 1	he exact sequence is proprietary.		
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity	F	Recognizes endogenous leve	els of Protein C protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/50 - 1/100)			
Gene Symbol	F	PROC			
Alternative Names		Vitamin K-dependent protein C; Anticoagulant protein C; Autoprothrombin IIA;			
	E	Blood coagulation factor XIV			
Entrez Gene		5624 (Human); 19123 (Mouse); 25268 (Rat)			
SwissProt	F	P04070 (Human); P33587 (Mouse); P31394 (Rat)			
Storage/Stabi	ability Shipped at 4°C. Upon delivery 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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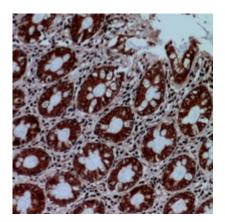




For research purposes only, not for human use

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Western blot analysis of Protein C expression in Raw264.7 (A), HepG2 (B), Jurkat (C), HEK293T (D) whole cell lysates. (Predicted band size: 52 kD; Observed band size: 52 kD)



Immunohistochemical analysis of Protein C staining in human colon 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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