

Product Data Sheet

Anti-Factor XIII B Antibody

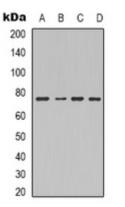
Catalog #	Source	Reactivity	Applications		
CPA7291	Rabbit	Н, М	WB, IH		
Description	Rabbit polyclonal antibody to Factor XIII B				
Immunogen		KLH-conjugated synthetic pe	otide encompassing a sequence within the center		
		region of human Factor XIII E	. The exact sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of Factor XIII B 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) - 1/200)		
Gene Symbol		F13B			
Alternative Na	ames	Coagulation factor XIII B chai	n; Fibrin-stabilizing factor B subunit; Protein-glutamine		
		gamma-glutamyltransferase	B chain; Transglutaminase B chain		
Entrez Gene		2165 (Human); 14060 (Mous	e)		
SwissProt		P05160 (Human); Q07968 (N	louse)		
Storage/Stabi	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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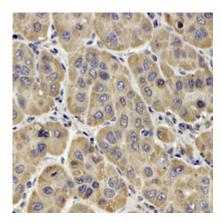




For research purposes only, not for human use

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Western blot analysis of Factor XIII B expression in Hela (A), mouse kidney (B), mouse heart (C), mouse liver (D) whole cell lysates. (Predicted band size: 75 kD; Observed band size: 75 kD)



Immunohistochemical analysis of Factor XIII B staining in human liver 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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