

Product Data Sheet

Anti-Factor XIIIa Antibody

Catalog #	Source	Reactivity	Applications	
CPA5720	Rabbit	H, M, R	WB, IH	
Description	Rabbit polyclonal antibody to Factor XIIIa			
Immunogen	I	KLH-conjugated synthetic per	tide encompassing a sequence within the N-term	
	I	region of human Factor XIIIa.	The exact sequence is proprietary.	
Purification	-	The antibody was purified by	immunogen affinity chromatography.	
Specificity	I	Recognizes endogenous level	s of Factor XIIIa protein.	
Clonality	I	Polyclonal		
Conjugation				
Form	I	Liquid in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
	ä	and 0.01% sodium azide.		
Dilution	v	WB (1/500 - 1/1000), IH (1/50 ·	1/100)	
Gene Symbol	I	F13A1		
Alternative Na	ames	F13A; Coagulation factor XIII	A chain; Coagulation factor XIIIa; Protein-glutamine	
	ł	gamma-glutamyltransferase	A chain; Transglutaminase A chain	
Entrez Gene		2162 (Human); 74145 (Mous	e); 60327 (Rat)	
SwissProt	I	P00488 (Human); Q8BH61 (N	louse); O08619 (Rat)	
Storage/Stabi	lity S	Shipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid	
	t	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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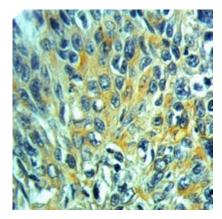
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kDa A B C D E F 180 130 95 72 Western blot analysis of Factor XIIIa expression in mouse testis (A), rat testis (B), K562 (C), EC9706 (D), MCF7 (E), HCT116 (F) whole cell lysates. (Predicted band size: 83 kD; Observed band size: 100 kD)



Immunohistochemical analysis of Factor XIIIa staining in human lung 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 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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