

## **Product Data Sheet**

### **Anti-Factor VIII Antibody**

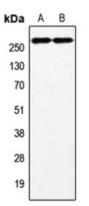
Catalog #	Source	Reactivity	Applications		
CPA1407	Rabbit	H, D	WB, IH		
Description	Ra	Rabbit polyclonal antibody to Factor VIII			
Immunogen	KLI	H-conjugated synthetic p	eptide encompassing a sequence within the C-term		
	reg	gion of human Factor VIII.	The exact sequence is proprietary.		
Purification	Th	e antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous lev	els of Factor VIII protein.		
Clonality	Po	lyclonal			
Conjugation					
Form	Liq	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	d 0.01% sodium azide.			
Dilution	WE	B (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	F8				
Alternative Na	ames F80	C; Coagulation factor VIII;	Antihemophilic factor; AHF; Procoagulant component		
Entrez Gene		2157 (Human)			
SwissProt	PO	0451 (Human)			
Storage/Stabi	<b>lity</b> Shi	ipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fre	eze/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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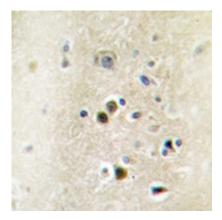
# Cohesion



For research purposes only, not for human use

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Western blot analysis of Factor VIII expression in HeLa (A), HepG2 (B) whole cell lysates. (Predicted band size: 267 kD; Observed band size: 280 kD)



Immunohistochemical analysis of Factor VIII 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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