

Product Data Sheet

Anti-IFN beta Antibody

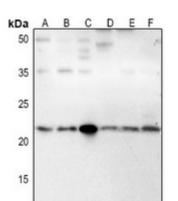
Catalog #	Source	Reactivity	Applications			
CPA6218	Rabbit	H, M, R	WB, IH			
Description	R	Rabbit polyclonal antibody to IFN beta				
Immunogen	K	KLH-conjugated synthetic peptide encompassing a sequence within the center				
	r	region of human IFN beta. The exact sequence is proprietary.				
Purification	Т	he antibody was purified by	/ immunogen affinity chromatography.			
Specificity	R	Recognizes endogenous leve	ls of IFN beta protein.			
Clonality	Р	Polyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	а	nd 0.01% sodium azide.				
Dilution	V	VB (1/500 - 1/1000), IH (1/50	- 1/200)			
Gene Symbol	I	FNB1				
Alternative Na	ames II	FB; IFNB; Interferon beta; IF	N-beta; Fibroblast interferon			
Entrez Gene		3456 (Human); 15977 (Mouse); 24481 (Rat)				
SwissProt	ot P01574 (Human); P01575 (Mouse); P70499 (Rat)		louse); P70499 (Rat)			
Storage/Stabi	lity S	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid				
	fı	reeze/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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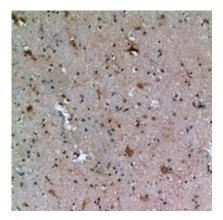
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For research purposes only, not for human use

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Western blot analysis of IFN beta expression in Hela (A), H1688 (B), H446 (C), mouse lung (D), mouse kidney (E), rat spleen (F) whole cell lysates. (Predicted band size: 22 kD; Observed band size: 22 kD)



Immunohistochemical analysis of IFN beta 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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