

## **Product Data Sheet**

### Anti-IL-1 beta Antibody

Catalog #	Source	Reactivity	Applications		
CPA1586	Rabbit	H, M, R, Mk	WB, IH		
Description	Rab	Rabbit polyclonal antibody to IL-1 beta			
Immunogen	KLH	-conjugated synthetic pe	otide encompassing a sequence within the center		
	regi	on of human IL-1 beta. Tl	ne exact sequence is proprietary.		
Purification	The	antibody was purified by	immunogen affinity chromatography.		
Specificity	Reco	ognizes endogenous leve	ls of IL-1 beta protein.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqu	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 (1/100	) - 1/200)		
Gene Symbol	IL1B				
Alternative Na	ames IL1F	2; Interleukin-1 beta; IL-2	beta; Catabolin		
Entrez Gene	3553	3 (Human)			
SwissProt	P01	584 (Human)			
Storage/Stabi	lity Ship	ped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	free	ze/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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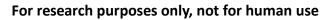
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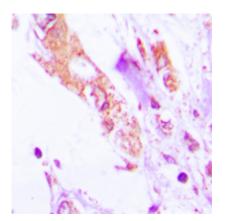
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kDa A B C D E F



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Western blot analysis of IL-1 beta expression in Raw264.7 (A), mouse spleen (B), rat spleen (C), rat thymus (D), mouse thymus (E), Jurkat (F), THP1 (G) whole cell lysates. (Predicted band size: 30 kD; Observed band size: 43; 35; 17 kD)



Immunohistochemical analysis of IL-1 beta 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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