

# **Product Data Sheet**

### Anti-pro-NGF beta Antibody

Catalog #	Source	Reactivity	Applications		
CPA4124	Rabbit	H, M, R	WB, IH		
Description	Rab	Rabbit polyclonal antibody to pro-NGF beta			
Immunogen	KLH	I-conjugated synthetic p	eptide encompassing a sequence within the center		
	regi	ion of human pro-NGF b	eta. The exact sequence is proprietary.		
Purification	The	e antibody was purified l	by immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous lev	els of pro-NGF beta protein.		
Clonality	Poly	yclonal			
Conjugation					
Form	Liqu	uid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	l 0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/5	0 - 1/100)		
Gene Symbol	NGI	F			
Alternative Na	ames NGI	FB; Beta-nerve growth f	actor; Beta-NGF		
Entrez Gene	480	)3 (Human); 18049 (Mo	ıse); 310738 (Rat)		
SwissProt	P01	138 (Human); P01139 (	Mouse); P25427 (Rat)		
Storage/Stabi	lity Ship	pped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
	free	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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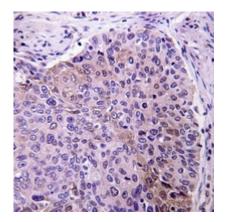
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Western blot analysis of pro-NGF beta expression in H1792 (A), mouse lung (B), rat lung (C) whole cell lysates. (Predicted band size: 26 kD; Observed band size: 27 kD)



Immunohistochemical analysis of pro-NGF 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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