

## **Product Data Sheet**

## **Anti-NRG4 Antibody**

Catalog #	Source	Reactivity	Applications			
CPA4133	Rabbit	H, M, R, Mk	WB, IH			
Description	Rab	Rabbit polyclonal antibody to NRG4				
Immunogen	KLH	-conjugated synthetic pe	ptide encompassing a sequence within the center			
	regi	region of human NRG4. The exact sequence is proprietary.				
Purification	The	The antibody was purified by immunogen affinity chromatography.				
Specificity	Reco	Recognizes endogenous levels of NRG4 protein.				
Clonality	ity Polyclonal					
Conjugation						
Form	Liqu	id in 0.42% Potassium pł	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycero	ol,		
	and	0.01% sodium azide.				
Dilution	WB	(1/500 - 1/1000), IH (1/50	- 1/100)			
Gene Symbol	NRG	64				
Alternative Names Pro-ne		neuregulin-4, membrane	uregulin-4, membrane-bound isoform; Pro-NRG4			
Entrez Gene	Entrez Gene 145957 (		(Human); 83961 (Mouse)			
SwissProt	Q8V	Q8WWG1 (Human); Q9WTX4 (Mouse)				
Storage/Stabi	lity Ship	Shipped at 4°C. Upon delivery 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

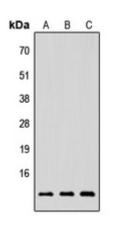
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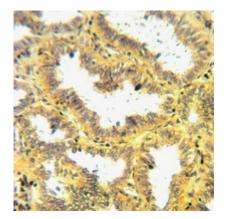
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Western blot analysis of NRG4 expression in HEK293T (A), Raw264.7 (B), H9C2 (C) whole cell lysates. (Predicted band size: 12 kD; Observed band size: 12 kD)



Immunohistochemical analysis of NRG4 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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