

# **Product Data Sheet**

# **Anti-NEFH Antibody**

Catalog #	Source	Reactivity	Applications		
CPA6012	Rabbit	H, M, R	WB, IH, IC		
Description	Ra	abbit polyclonal antibody t	o NEFH		
Immunogen	KL	.H-conjugated synthetic pe	ptide encompassing a sequence within the C-term		
	re	gion of human NEFH. The	exact sequence is proprietary.		
Purification	Th	ie antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	els of NEFH protein.		
Clonality	Ро	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IH (1/50	- 1/200), IC (1/100 - 1/500)		
Gene Symbol	NE	EFH			
Alternative Na	ames Kl	AA0845; NFH; Neurofilam	ent heavy polypeptide; NF-H; 200 kDa neurofilament		
	pr	otein; Neurofilament tripl	et H protein		
Entrez Gene	47	744 (Human); 380684 (Mo	use)		
SwissProt	P1	12036 (Human); P19246 (N	1ouse); P16884 (Rat)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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

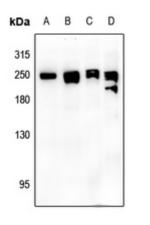
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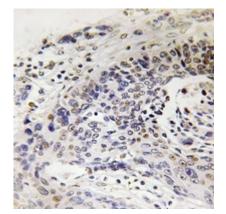


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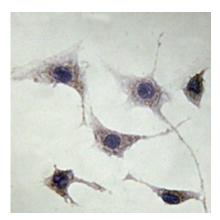
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Western blot analysis of NEFH expression in C6 (A), CT26 (B), HepG2 (C), A2780 (D) whole cell lysates. (Predicted band size: 112 kD; Observed band size: 250 kD)



Immunohistochemical analysis of NEFH 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.



Immunocytochemistry analysis of NEFH staining in NIH3T3 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a HRP-conjugated secondary antibody in PBS at room temperature. DAB was used as the chromogen.

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