

Product Data Sheet

Anti-NEIL3 Antibody

Catalog #	Source	e Reactivity	Applications		
CPA2705	Rabbit	H, M, R	WB, IH		
Description		Rabbit polyclonal antibody t	o NEIL3		
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the C-term		
		region of human NEIL3. The	exact sequence is proprietary.		
Purification		The antibody was purified by	<i>immunogen affinity chromatography.</i>		
Specificity		Recognizes endogenous leve	ls of NEIL3 protein.		
Clonality		Polyclonal			
Conjugation					
Form		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 (1/50	- 1/100)		
Gene Symbol		NEIL3			
Alternative Na	ames	Endonuclease 8-like 3; DNA	glycosylase FPG2; DNA glycosylase/AP lyase Neil3;		
		Endonuclease VIII-like 3; Nei	-like protein 3		
Entrez Gene		55247 (Human); 234258 (Mouse)			
SwissProt		Q8TAT5 (Human); Q8K203 (N	/louse)		
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
		freeze/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

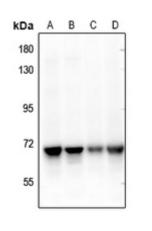
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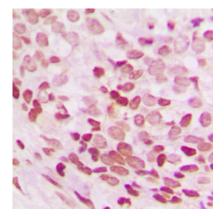


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Western blot analysis of NEIL3 expression in MCF7 (A), H1792 (B), rat thymus (C), mouse testis (D) whole cell lysates. (Predicted band size: 67 kD; Observed band size: 68 kD)



Immunohistochemical analysis of NEIL3 staining in human breast 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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