

Product Data Sheet

Anti-TAF5L Antibody

Catalog #	Source	Reactivity	Applications		
CPA5123	Rabbit	H, D, Mk, Rb	WB, IH		
Description	I	Rabbit polyclonal antibody to	TAF5L		
Immunogen	I	KLH-conjugated synthetic pep	tide encompassing a sequence within the center		
	I	region of human TAF5L. The e	kact sequence is proprietary.		
Purification	-	The antibody was purified by	mmunogen affinity chromatography.		
Specificity	I	Recognizes endogenous levels	of TAF5L protein.		
Clonality	I	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution	v	WB (1/500 - 1/1000), IH (1/100	- 1/200)		
Gene Symbol	-	TAF5L			
Alternative Na	ames	PAF65B; TAF5-like RNA polym	erase II p300/CBP-associated factor-associated factor		
	(65 kDa subunit 5L; PCAF-asso	iated factor 65 beta; PAF65-beta		
Entrez Gene		27097 (Human)			
SwissProt	(075529 (Human)			
Storage/Stabi	lity S	Shipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	f	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

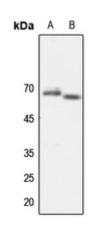
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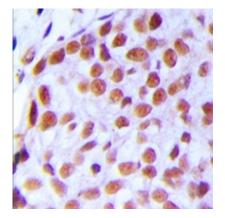
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For research purposes only, not for human use

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Western blot analysis of TAF5L expression in HEK293T (A), Hela (B) whole cell lysates. (Predicted band size: 66 kD; Observed band size: 66 kD)



Immunohistochemical analysis of TAF5L 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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