

## **Product Data Sheet**

## **Anti-IDOL Antibody**

Out also in	6.	Describerts	Andreathers
Catalog #	Source	Reactivity	Applications
CPA4106	Rabbit	H, M, R, B, Mk	WB, IH
Description		Rabbit polyclonal antibody to	DOL
Immunogen		KLH-conjugated synthetic pept	ide encompassing a sequence within the center
		region of human IDOL. The exa	ct sequence is proprietary.
Purification		The antibody was purified by i	mmunogen affinity chromatography.
Specificity		Recognizes endogenous levels	of IDOL protein.
Clonality		Polyclonal	
Conjugation			
Form		Liquid in 0.42% Potassium pho	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
		and 0.01% sodium azide.	
Dilution		WB (1/500 - 1/1000), IH (1/100	1/200)
Gene Symbol		MYLIP	
Alternative Na	ames	BZF1; IDOL; E3 ubiquitin-prote	in ligase MYLIP; Inducible degrader of the
		LDL-receptor; Idol; Myosin reg	ulatory light chain interacting protein; MIR
Entrez Gene		29116 (Human); 218203 (Mou	se)
SwissProt		Q8WY64 (Human); Q8BM54 (I	/louse)
Storage/Stabi	lity	Shipped at 4°C. Upon delivery	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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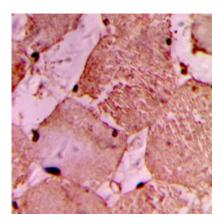
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Western blot analysis of IDOL expression in Hela (A), COS7 (B), MCF7 (C), PC12 (D), CT26 (E) whole cell lysates. (Predicted band size: 49 kD; Observed band size: 50 kD)



Immunohistochemical analysis of IDOL staining in human heart 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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