

# **Product Data Sheet**

## **Anti-TOLLIP Antibody**

Catalog #	Source	Reactivity	Applications			
CPA2693	Rabbit	H, M, R	WB, IH, IP			
Description	Rat	Rabbit polyclonal antibody to TOLLIP				
Immunogen	KLH	H-conjugated synthetic p	eptide encompassing a sequence within the center			
	reg	region of human TOLLIP. The exact sequence is proprietary.				
Purification	The	The antibody was purified by immunogen affinity chromatography.				
Specificity	Red	cognizes endogenous lev	els of TOLLIP protein.			
Clonality	Pol	yclonal				
Conjugation						
Form	Liq	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	and	d 0.01% sodium azide.				
Dilution	WE	3 (1/500 - 1/1000), IH (1/10	00 - 1/200), IP (1/10 - 1/100)			
Gene Symbol	TO	LLIP				
Alternative Na	ames Tol	l-interacting protein				
Entrez Gene	544	54472 (Human)				
SwissProt	Q9	H0E2 (Human)				
Storage/Stabi	lity Shi	pped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid			
	fre	eze/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

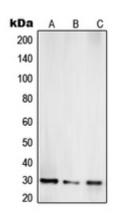
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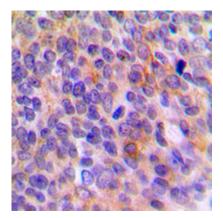


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Western blot analysis of TOLLIP expression in Jurkat (A), LNCaP (B), HepG2 (C) whole cell lysates. (Predicted band size: 22; 30 kD; Observed band size: 30 kD)



Immunohistochemical analysis of TOLLIP 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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