

Product Data Sheet

Anti-G15 Antibody

Catalog #	Source	e Reactivity	Applications		
CPA4194	Rabbit	H, M, R, B	WB, IH		
Description		Rabbit polyclonal antibody t	o G15		
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the center		
		region of human G15. The ex	act sequence is proprietary.		
Purification		The antibody was purified by	<i>immunogen affinity chromatography.</i>		
Specificity		Recognizes endogenous leve	ls of G15 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 (1/10	0 - 1/200)		
Gene Symbol		RBM5			
Alternative Na	ames	RNA-binding protein 5; Prote	in G15; Putative tumor suppressor LUCA15;		
		RNA-binding motif protein 5	; Renal carcinoma antigen NY-REN-9		
Entrez Gene		10181 (Human); 83486 (Mouse); 300996 (Rat)			
SwissProt		P52756 (Human); Q91YE7 (N	1ouse); B2GV05 (Rat)		
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

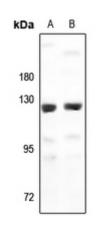
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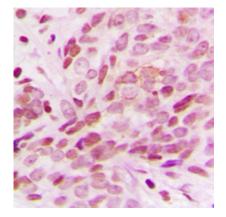
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Western blot analysis of G15 expression in K562 (A), Jurkat (B) whole cell lysates. (Predicted band size: 92 kD; Observed band size: 125 kD)



Immunohistochemical analysis of G15 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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