

Product Data Sheet

Anti-GABRG1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA5211	Rabbit	H, M, R	WB, IH		
			· ·		
Description		abbit polyclonal antibody t			
Immunogen	KI	LH-conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	re	egion of human GABRG1. T	he exact sequence is proprietary.		
Purification	Tł	ne antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	els of GABRG1 protein.		
Clonality	Po	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol	G	ABRG1			
Alternative Na	ames Ga	amma-aminobutyric acid ı	eceptor subunit gamma-1; GABA(A) receptor subunit		
	ga	amma-1			
Entrez Gene		2565 (Human); 14405 (Mouse); 140674 (Rat)			
SwissProt	Q	8N1C3 (Human); Q9R0Y8	Mouse); P23574 (Rat)		
Storage/Stabi	lity Sł	nipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fr	eeze/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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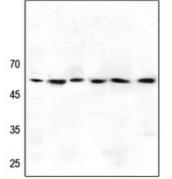
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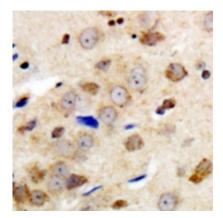
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KDA A B C D E F



Western blot analysis of GABRG1 expression in HEK293T (A), LOVO (B), Hela (C), mouse brain (D), mouse kidney (E), rat kidney (F) whole cell lysates. (Predicted band size: 53 kD; Observed band size: 54 kD)



Immunohistochemical analysis of GABRG1 staining in human brain 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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