

Product Data Sheet

Anti-GRK3 Antibody

Catalog #	Source	Reactivity	Applications		
CPA7193	Rabbit	H, M, R, B	WB, IH		
Description	R	abbit polyclonal antibody t	o GRK3		
Immunogen	K	(LH-conjugated synthetic pe	ptide encompassing a sequence within the center		
	r	egion of human GRK3. The	exact sequence is proprietary.		
Purification	Т	he antibody was purified by	y immunogen affinity chromatography.		
Specificity	R	Recognizes endogenous leve	els of GRK3 protein.		
Clonality	Р	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	and 0.01% sodium azide.			
Dilution	V	NB (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol	А	ADRBK2			
Alternative Na	ames B	3ARK2; GRK3; Beta-adrener	gic receptor kinase 2; Beta-ARK-2; G-protein-coupled		
	r	eceptor kinase 3			
Entrez Gene	1	.57 (Human); 320129 (Mous	se); 25372 (Rat)		
SwissProt	Р	235626 (Human); Q3UYH7 (I	Mouse); P26819 (Rat)		
Storage/Stabi	lity S	hipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	fi	reeze/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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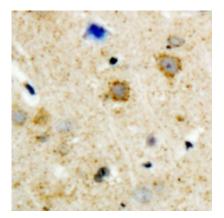
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Western blot analysis of GRK3 expression in Hela (A), HL60 (B) whole cell lysates. (Predicted band size: 79 kD; Observed band size: 83 kD)



Immunohistochemical analysis of GRK3 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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