

# **Product Data Sheet**

### Anti-GRK2 Antibody

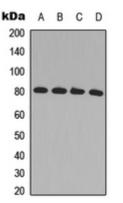
Catalog #	Source	Reactivity	Applications		
CPA4555	Rabbit	H, M, R, B	WB, IH		
Description	R	Rabbit polyclonal antibody t	o GRK2		
Immunogen	K	(LH-conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	r	egion of human GRK2. The	exact sequence is proprietary.		
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	Recognizes endogenous leve	ls of GRK2 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	ADRBK1			
Alternative Na	a <mark>mes</mark> B	3ARK; BARK1; GRK2; Beta-ad	Irenergic receptor kinase 1; Beta-ARK-1; G-protein		
	С	coupled receptor kinase 2			
Entrez Gene	1	.56 (Human); 25238 (Rat)			
SwissProt	Р	25098 (Human); Q99MK8 (	Mouse); P26817 (Rat)		
Storage/Stabi	lity S	hipped at 4°C. Upon delive	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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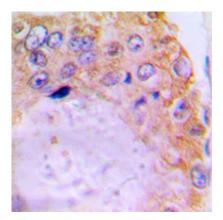




For research purposes only, not for human use

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Western blot analysis of GRK2 expression in Jurkat (A), Raji (B), mouse kidney (C), rat kidney (D) whole cell lysates. (Predicted band size: 79 kD; Observed band size: 80 kD)



Immunohistochemical analysis of GRK2 staining in human prostate 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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