

Product Data Sheet

Anti-KCNK10 Antibody

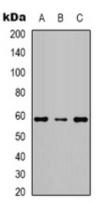
Catalog #	Source	Reactivity	Applications		
CPA9403	Rabbit	H, M, R	WB, IH		
Description		Rabbit polyclonal antibody to KCNK10			
Immunogen		KLH-conjugated synthetic pe	otide encompassing a sequence of human KCNK10. The		
		exact sequence is proprietary	Ι.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	s of KCNK10 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 (1/1000 - 1/2000), IH (1/10	00 - 1/200)		
Gene Symbol		KCNK10			
Alternative Na	ames	TREK2; Potassium channel su	bfamily K member 10; Outward rectifying potassium		
	1	channel protein TREK-2; TREI	<-2 K(+) channel subunit		
Entrez Gene		54207 (Human)			
SwissProt		P57789 (Human)			
Storage/Stabil	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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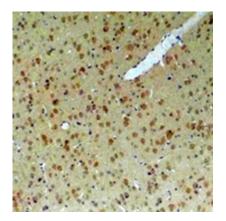




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Western blot analysis of KCNK10 expression in Jurkat (A), mouse brain (B), rat brain (C) whole cell lysates. (Predicted band size: 59 kD; Observed band size: 59 kD)



Immunohistochemical analysis of KCNK10 staining in rat 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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