

# **Product Data Sheet**

### Anti-KCNN2 Antibody

Catalog #	Source	Reactivity	Applications		
CPA9405	Rabbit	Н, М, R	WB, IH		
Description	R	abbit polyclonal antibody to	o KCNN2		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence of human KCNN2. The			
	е	xact sequence is proprietar	<i>I</i> .		
Purification	Т	he antibody was purified by	immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous leve	ls of KCNN2 protein.		
Clonality Polyclonal					
Conjugation					
Form	Li	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	nd 0.01% sodium azide.			
Dilution	V	VB (1/1000 - 1/2000), IH (1/1	00 - 1/200)		
Gene Symbol	K	CNN2			
Alternative Na	<mark>ames</mark> S	mall conductance calcium-a	ctivated potassium channel protein 2; SK2; SKCa 2;		
	S	KCa2; KCa2.2			
Entrez Gene	3	781 (Human)			
SwissProt	С	Q9H2S1 (Human)			
Storage/Stabi	lity S	hipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	fr	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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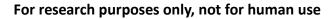
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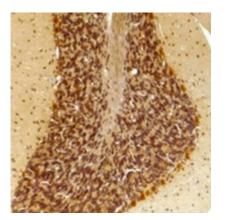
200

140



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Western blot analysis of KCNN2 expression in HepG2 (A), mouse brain (B), rat brain (C) whole cell lysates. (Predicted band size: 63 kD; Observed band size: 70; 26 kD)



Immunohistochemical analysis of KCNN2 staining in human brain, mouse 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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