

Product Data Sheet

Anti-Kv11.1 Antibody

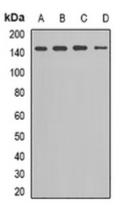
Catalog #	Source	e Reactivity	Applications	
CQA1155	Rabbit	t H, M	WB, IH	
Description		Rabbit polyclonal antibody to	o Kv11.1	
Immunogen		KLH-conjugated synthetic pe	otide of human Kv11.1	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	ls of Kv11.1 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/2000), IH (1/50	- 1/200)	
Gene Symbol		KCNH2		
Alternative Names		ERG; ERG1; HERG; Potassium voltage-gated channel subfamily H member 2; Eag		
		homolog; Ether-a-go-go-rela	ed gene potassium channel 1; ERG-1; Eag-related	
		protein 1; Ether-a-go-go-rela	ted protein 1; H-ERG; hERG-1; hERG1; Voltage-gated	
		potassium channel subunit K	v11.1	
Entrez Gene		3757 (Human); 16511 (Mous	e)	
SwissProt		Q12809 (Human); O35219 (N	/louse)	
Storage/Stabi	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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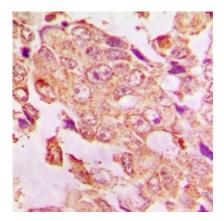




For research purposes only, not for human use

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Western blot analysis of Kv11.1 expression in SW620 (A), MCF7 (B), K562 (C), mouse brain (D) whole cell lysates. (Predicted band size: 60; 90; 97; 115; 121; 126 kD; Observed band size: 148 kD)



Immunohistochemical analysis of Kv11.1 staining in human breast 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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