

# **Product Data Sheet**

# Anti-Kv1.5 Antibody

Catalog #	Source	Reactivity	Applications		
CPA1643	Rabbit	H, M, R, B, D, P	WB, IH		
Description	R	Rabbit polyclonal antibody to	<v1.5< td=""></v1.5<>		
Immunogen	К	LH-conjugated synthetic pept	ide encompassing a sequence within the center		
	re	egion of human Kv1.5. The ex	act sequence is proprietary.		
Purification	Т	he antibody was purified by i	mmunogen affinity chromatography.		
Specificity	R	Recognizes endogenous levels	of Kv1.5 protein.		
Clonality	P	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	nd 0.01% sodium azide.			
Dilution	V	NB (1/500 - 1/1000), IH (1/50 -	1/100)		
Gene Symbol	К	CNA5			
Alternative Na	ames P	otassium voltage-gated chan	nel subfamily A member 5; HPCN1; Voltage-gated		
	р	otassium channel HK2; Volta	ge-gated potassium channel subunit Kv1.5		
Entrez Gene	3	8741 (Human); 16493 (Mouse	; 25470 (Rat)		
SwissProt	Р	22460 (Human); Q61762 (Mc	use); P19024 (Rat)		
Storage/Stabi	lity S	hipped at 4°C. Upon delivery	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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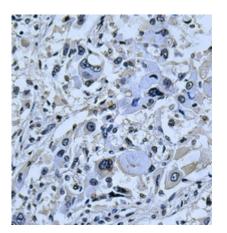
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kDa A B C

For research purposes only, not for human use

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Western blot analysis of Kv1.5 expression in MCF7 (A), Raw264.7 (B), H9C2 (C) whole cell lysates. (Predicted band size: 67 kD; Observed band size: 56 kD)



Immunohistochemical analysis of Kv1.5 staining in human liver 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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