

## **Product Data Sheet**

## **Anti-CLIC4 Antibody**

Catalog #	Source	Reactivity	Applications		
CPA9269	Rabbit	H, M, R	WB, IH		
Description	R	abbit polyclonal antibody t	D CLIC4		
Immunogen	K	LH-conjugated synthetic pe	otide encompassing a sequence of human CLIC4. The		
	e	xact sequence is proprietar	<i>I</i> .		
Purification	T	he antibody was purified by	immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous leve	ls of CLIC4 protein.		
Clonality	P	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	a	nd 0.01% sodium azide.			
Dilution	W	VB (1/1000 - 1/2000), IH (1/1	00 - 1/200)		
Gene Symbol	C	LIC4			
Alternative Na	ames C	hloride intracellular channe	l protein 4; Intracellular chloride ion channel protein		
	p	64H1			
Entrez Gene	2	5932 (Human)			
SwissProt	Q	29Y696 (Human)			
Storage/Stabi	lity Sl	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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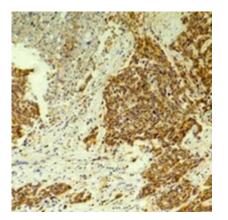
**kDa** A B 200

140

For research purposes only, not for human use

# **Product Data Sheet**

Western blot analysis of CLIC4 expression in mouse brain (A), rat brain (B) whole cell lysates. (Predicted band size: 28 kD; Observed band size: 30 kD)



Immunohistochemical analysis of CLIC4 staining in human breast cancer, 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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