

# **Product Data Sheet**

### **Anti-Cerebellin 4 Antibody**

Catalog #	Source	Reactivity	Applications
CPA2868	Rabbit	H, M, R, B, Mk, P	WB, IH
Description	Rabb	it polyclonal antibody to Ce	erebellin 4
Immunogen	KLH-c	conjugated synthetic peptid	le encompassing a sequence within the C-term
	regio	n of human Cerebellin 4. Th	ne exact sequence is proprietary.
Purification	The a	ntibody was purified by im	munogen affinity chromatography.
Specificity	Reco	gnizes endogenous levels of	f Cerebellin 4 protein.
Clonality	Polyc	lonal	
Conjugation			
Form	Liquio	d in 0.42% Potassium phosp	ohate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and C	.01% sodium azide.	
Dilution	WB (	I/500 - 1/1000), IH (1/100 - 1	/200)
Gene Symbol	CBLN	4	
Alternative Na	ames CBLN	L1; Cerebellin-4; Cerebellin	-like glycoprotein 1
Entrez Gene	1406	89 (Human); 228942 (Mous	se)
SwissProt	Q9N1	U7 (Human); Q8BME9 (Mc	ouse)
Storage/Stabi	lity Shipp	ed at 4°C. Upon delivery al	iquot and store at -20°C for one year. Avoid
	freez	e/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

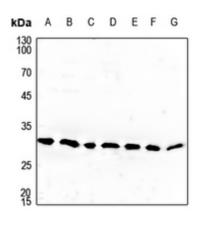
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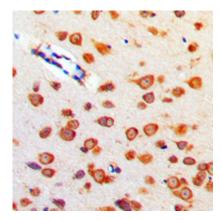


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Western blot analysis of Cerebellin 4 expression in HEK293T (A), Hela (B), A549 (C), mouse lung (D), mouse liver (E), rat lung (F), rat liver (G) whole cell lysates. (Predicted band size: 21 kD; Observed band size: 30 kD)



Immunohistochemical analysis of Cerebellin 4 staining in human 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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