

Product Data Sheet

Anti-ALPL Antibody

Catalog #	Source	Reactivity	Applications
CQA3800	Rabbit	H, M, R	WB, IH
Description	Ral	bbit polyclonal antibody	to ALPL
Immunogen	Red	combinant fusion proteir	of human ALPL. The exact sequence is proprietary.
Purification	The	e antibody was purified b	y immunogen affinity chromatography.
Specificity	Red	cognizes endogenous lev	els of ALPL protein
Clonality	Pol	lyclonal	
Conjugation			
Form	Liq	uid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	d 0.01% sodium azide.	
Dilution	WE	3 (1/500 - 1/2000), IH (1/5) - 1/200)
Gene Symbol	ALI	PL	
Alternative N	ames Alk	aline phosphatase tissue	-nonspecific isozyme; AP-TNAP; TNSALP; Alkaline
	pho	osphatase liver/bone/kic	ney isozyme
Entrez Gene	249	9 (Human); 11647 (Mous	e); 25586 (Rat)
SwissProt	PO!	5186 (Human); P09242 (Mouse); P08289 (Rat)
Storage/Stabi	i lity Shi	pped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid
	fre	eze/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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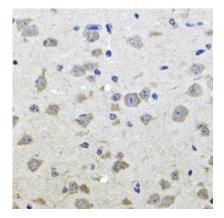


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kDa <u>A</u> <u>B</u> <u>C</u> 130 95 72 55 43 34

Western blot analysis of ALPL expression in HeLa (A), mouse kidney (B), rat kidney (C) whole cell lysates. (Predicted band size: 48; 51; 57 kD; Observed band size: 80 kD)



Immunohistochemical analysis of ALPL staining in 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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