

Product Data Sheet

Anti-ALP Antibody

Catalog #	Source	e Reactivity	Applications	
CPA3764	Rabbit	Н, М	WB, IH	
Description		Rabbit polyclonal antibody	o ALP	
Immunogen		KLH-conjugated synthetic pe	eptide encompassing a sequence within the N-term	
		region of human ALP. The ex	act sequence is proprietary.	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of ALP protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/1000), IH (1/50	- 1/200)	
Gene Symbol		ALPP; ALPPL2		
Alternative N	ames	ALPP; PLAP; Alkaline phospl	natase, placental type; Alkaline phosphatase Regan	
		isozyme; Placental alkaline j	phosphatase 1; PLAP-1; ALPPL2; ALPPL; Alkaline	
		phosphatase, placental-like;	ALP-1; Alkaline phosphatase Nagao isozyme; Germ cell	
		alkaline phosphatase; GCAP	; Placental alkaline phosphatase-like; PLAP-like	
Entrez Gene		250, 251 (Human); 11650 (N	1ouse)	
SwissProt		P05187, P10696 (Human); P	24823 (Mouse)	
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry 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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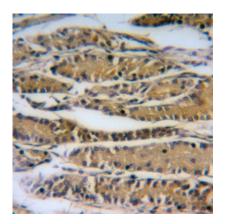
kDa A B

140 100

For research purposes only, not for human use

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Western blot analysis of ALP expression in HEK293T (A), Raw264.7 (B) whole cell lysates. (Predicted band size: 57 kD; Observed band size: 58 kD)



Immunohistochemical analysis of ALP staining in human muscle 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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