

Product Data Sheet

Anti-ANKRD1 Antibody

Catalog #	Source	Reactivity	Applications			
CPA3404	Rabbit	H, M, R, B, C, P, Rb	WB, IH			
Description	Ra	bbit polyclonal antibody to	ANKRD1			
Immunogen	KL	H-conjugated synthetic pept	ide encompassing a sequence within the C-term			
	re	gion of human ANKRD1. Thε	exact sequence is proprietary.			
Purification	Th	e antibody was purified by i	mmunogen affinity chromatography.			
Specificity	Re	cognizes endogenous levels	of ANKRD1 protein.			
Clonality	Ро	lyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	an	d 0.01% sodium azide.				
Dilution	W	B (1/500 - 1/1000), IH (1/100	- 1/200)			
Gene Symbol	AN	IKRD1				
Alternative Na	ames C1	93; CARP; HA1A2; Ankyrin r	epeat domain-containing protein 1; Cardiac ankyrin			
	rep	peat protein; Cytokine-induc	tible gene C-193 protein; Cytokine-inducible nuclear			
	pro	otein				
Entrez Gene	27	063 (Human); 107765 (Mou	se); 27064 (Rat)			
SwissProt	Q1	.5327 (Human); Q9CR42 (M	ouse); Q8R560 (Rat)			
Storage/Stabi	lity Sh	ipped at 4°C. Upon delivery	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

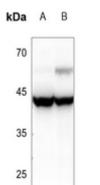
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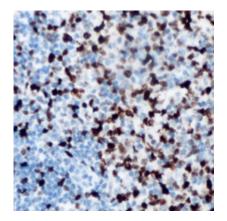


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Western blot analysis of ANKRD1 expression in mouse muscle (A), rat muscle (B) whole cell lysates. (Predicted band size: 36 kD; Observed band size: 43 kD)



Immunohistochemical analysis of ANKRD1 staining in human lymph node 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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