

Product Data Sheet

Anti-AARSD1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA2796	Rabbit	H, M, R, B	WB, IH		
Description	F	Rabbit polyclonal antibody to	DAARSD1		
Immunogen	k	KLH-conjugated synthetic pe	otide encompassing a sequence within the center		
	r	region of human AARSD1. Th	e exact sequence is proprietary.		
Purification	Т	The antibody was purified by	immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	ls of AARSD1 protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	a	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	Ļ	AARSD1			
Alternative Na	ames A	Alanyl-tRNA editing protein	arsd1; Alanyl-tRNA synthetase domain-containing		
	ŗ	protein 1			
Entrez Gene	1	100885850, 80755 (Human);	69684 (Mouse); 619440 (Rat)		
SwissProt	C	Q9BTE6 (Human); Q3THG9 (Mouse); Q5XI97 (Rat)		
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	f	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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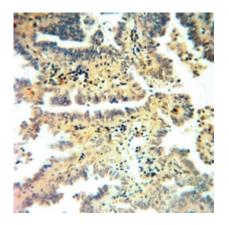
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kDa A B C 105 70 45 35 25 Western blot analysis of AARSD1 expression in Jurkat (A), 3T3L1 (B), PMVEC (C) whole cell lysates. (Predicted band size: 45 kD; Observed band size: 45; 65 kD)



Immunohistochemical analysis of AARSD1 staining in human lung cancer 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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