

Product Data Sheet

Anti-SMARCAD1 Antibody

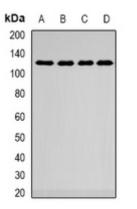
Catalog #	Source	Reactivity	Applications
CQA2241	Rabbit	H, M, R	WB, IF/IC
Description	Rabl	oit polyclonal antibody	to SMARCAD1
Immunogen	Reco	ombinant full length pro	tein of human SMARCAD1
Purification	The	antibody was purified b	y immunogen affinity chromatography.
Specificity	Reco	ognizes endogenous lev	els of SMARCAD1 protein.
Clonality	Poly	clonal	
Conjugation			
Form	Liqu	id in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1/500 - 1/1000), IF/IC (1	/50 - 1/200)
Gene Symbol	SMA	RCAD1	
Alternative Na	ames KIAA	1122; SWI/SNF-related	matrix-associated actin-dependent regulator of
	chro	matin subfamily A cont	aining DEAD/H box 1; ATP-dependent helicase 1; hHEL1
Entrez Gene	5692	L6 (Human); 13990 (Mc	use); 312398 (Rat)
SwissProt	Q9H	4L7 (Human); Q04692 (Mouse); D3Z9Z9 (Rat)
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid
	free	ze/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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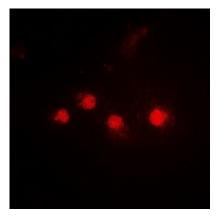




For research purposes only, not for human use

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Western blot analysis of SMARCAD1 expression in A549 (A), HepG2 (B), mouse kidney (C), mouse lung (D) whole cell lysates. (Predicted band size: 68; 117 kD; Observed band size: 120 kD)



Immunofluorescent analysis of SMARCAD1 staining in HepG2 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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