

# **Product Data Sheet**

### **Anti-AMD1** Antibody

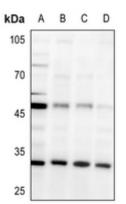
Catalog #	Source	Reactivity	Applications		
CPA1046	Rabbit	H, M, R, B	WB, IF/IC		
Description	Rat	bit polyclonal antibody	to AMD1		
Immunogen	KLF	I-conjugated synthetic po	eptide encompassing a sequence within the center		
	reg	ion of human AMD1. The	e exact sequence is proprietary.		
Purification	The	e antibody was purified b	y immunogen affinity chromatography.		
Specificity	Rec	cognizes endogenous lev	els of AMD1 protein.		
Clonality	Pol	yclonal			
Conjugation					
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	d 0.01% sodium azide.			
Dilution	WB	8 (1/500 - 1/1000), IF/IC (1	/50 - 1/200)		
Gene Symbol	AM	ID1			
Alternative Na	ames AM	ID; S-adenosylmethionin	e decarboxylase proenzyme; AdoMetDC; SAMDC		
Entrez Gene	262	2 (Human); 81640 (Rat)			
SwissProt	P17	7707 (Human); P17708 (F	at)		
Storage/Stabi	lity Shi	pped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	free	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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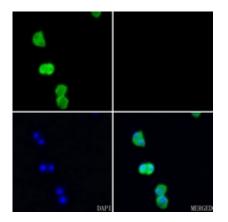




For research purposes only, not for human use

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Western blot analysis of AMD1 expression in AML12 (A), HCT116 (B), LO2 (C), A549 (D) whole cell lysates. (Predicted band size: 38 kD; Observed band size: 34 kD)



Immunofluorescent analysis of AMD1 staining in C6 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 hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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