

## **Product Data Sheet**

### Anti-ADAM15 Antibody

Catalog #	Source	e Reactivity	Applications	
CQA1798	Rabbit	t H, M, R	WB, IF/IC	
Description		Rabbit polyclonal antibody t	D ADAM15	
Immunogen		Recombinant full length prot	ein of human ADAM15	
Purification		The antibody was purified by	<i>immunogen affinity chromatography.</i>	
Specificity		Recognizes endogenous leve	ls of ADAM15 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 - 1/2000), IF/IC (1/	50 - 1/200)	
Gene Symbol		ADAM15		
Alternative Names		MDC15; Disintegrin and metalloproteinase domain-containing protein 15; ADAM 15;		
		Metalloprotease RGD disinte	grin protein; Metalloproteinase-like, disintegrin-like,	
		and cysteine-rich protein 15,	MDC-15; Metargidin	
Entrez Gene		8751 (Human); 11490 (Mous	se); 57025 (Rat)	
SwissProt		Q13444 (Human); O88839 (I	Mouse); Q9QYV0 (Rat)	
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y 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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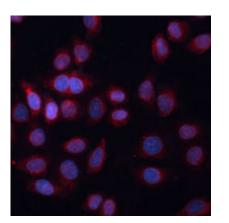
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For research purposes only, not for human use

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Western blot analysis of ADAM15 expression in mouse kidney (A), mouse brain (B) whole cell lysates. (Predicted band size: 55; 68; 83-92 kD; Observed band size: 93 kD)



Immunofluorescent analysis of ADAM15 staining in HeLa 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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