

# **Product Data Sheet**

### **Anti-CAD Antibody**

Catalog #	Source	Reactivity	Applications		
CQA1862	Rabbit	Н	WB, IH, IF/IC		
Description		Rabbit polyclonal antibody	r to CAD		
Immunogen		Recombinant full length pr	otein of human CAD		
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous le	vels of CAD 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), IH (1/5	60 - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol		CAD			
Alternative Na	ames	CAD protein			
Entrez Gene		790 (Human)			
SwissProt		P27708 (Human)			
Storage/Stabi	lity	Shipped at 4 $^\circ$ C. Upon del	ivery aliquot and store at -20 $^\circ$ 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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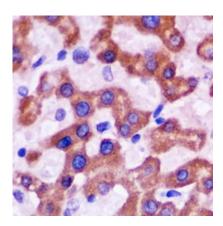
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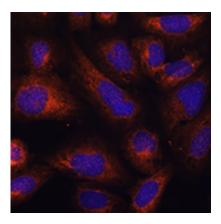
KDa A B C 200 140 100 80 60 50 For research purposes only, not for human use

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Western blot analysis of CAD expression in HepG2 (A), Hela (B), HL60 (C) whole cell lysates. (Predicted band size: 242 kD; Observed band size: 243 kD)



Immunohistochemical analysis of CAD staining in human liver 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.



Immunofluorescent analysis of CAD staining in U2OS 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  $^{\circ}$  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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