

# **Product Data Sheet**

### **Anti-Calponin 2 Antibody**

Catalog #	Source	Reactivity	Applications		
CPA3862	Rabbit	H, M, R, B	WB, IF/IC		
Description		Rabbit polyclonal antibody to Calponin 2			
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the center		
		region of human Calponin 2. The exact sequence is proprietary.			
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of Calponin 2 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 - 1/1000), IF/IC (1/	100 - 1/500)		
Gene Symbol		CNN2			
Alternative Na	cernative Names Calponin-2; Calponin H2, smooth muscle; Neutral calponin		ooth muscle; Neutral calponin		
Entrez Gene		1265 (Human); 12798 (Mouse)			
SwissProt		Q99439 (Human); Q08093 (I	/louse)		
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

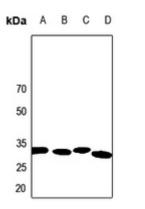
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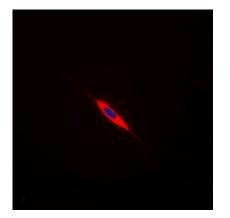


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Western blot analysis of Calponin 2 expression in Hela (A), mouse muscle (B), rat heart (C), rat muscle (D) whole cell lysates. (Predicted band size: 33 kD; Observed band size: 34 kD)



Immunofluorescent analysis of Calponin 2 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 hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-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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