

## **Product Data Sheet**

### **Recombinant Anti-SCP2 Rabbit mAb**

	Catalog #	Source	e Reactivity	Applications		
	CMA4293	Rabbit	: H	WB, IH, IF/IC		
	Description		Recombinant rabbit monoclonal antibody to SCP2			
	Immunogen		KLH-conjugated synthetic p	peptide encompassing a sequence within human SCP2.		
			The exact sequence is proprietary.			
Purification			The antibody was purified by immunogen affinity chromatography.			
Specificity			Recognizes endogenous levels of SCP2 protein			
	Clonality	Clonality Monoclonal				
	Conjugation					
Form			Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium azide.			
	Dilution		WB (1/500 - 1/1000), IH (1/5	50 - 1/200), IF/IC (1/50 - 1/200)		
	Gene Symbol SCP2					
	Alternative Na	mes	Non-specific lipid-transfer	protein; NSL-TP; Propanoyl-CoA C-acyltransferase;		
			SCP-chi; SCPX; Sterol carrie	er protein 2; SCP-2; Sterol carrier protein X; SCP-X		
Entrez Gene			6342 (Human)			
	SwissProt		P22307 (Human)			
	Storage/Stabil	ity	Shipped at 4°C. Upon deliv	ery 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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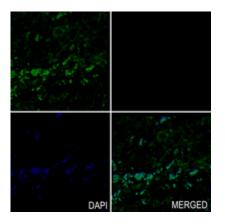
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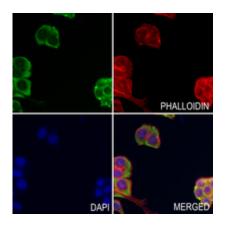
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Western blot analysis of SCP2 expression in HEK293T (A), THP1 (B) whole cell lysates. (Predicted band size: 58 kD; Observed band size: 58 kD)



Immunohistochemical analysis of SCP2 staining in human kidney 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. Tyramide-AF488 (green) was used as the chromogen. The section was then counterstained with DAPI (blue).



Immunofluorescent analysis of SCP2 staining in MCF7 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 an 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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