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# anti-Sema4a antibody (Center)

3 Images



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Alternative Name:

Quantity:	100 μL	
Target:	Sema4a	
Binding Specificity:	Center	
Reactivity:	Human, Monkey	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This Sema4a antibody is un-conjugated	
Application:	Western Blotting (WB), Immunofluorescence (IF), Immunoprecipitation (IP),	
	Immunochromatography (IC)	
Product Details		
Immunogen:	KLH-conjugated synthetic peptide encompassing a sequence within the center region of human	
	Semaphorin 4A.	
Specificity:	Recognizes endogenous levels of Semaphorin 4A protein.	
Characteristics:	Rabbit polyclonal antibody to Semaphorin 4A	
Purification:	The antibody was purified by immunogen affinity chromatography.	
Target Details		
Target:	Sema4a	

Semaphorin 4A (Sema4a Products)

#### **Target Details**

Background:	SEMAB, SEMB, Semaphorin-4A, Semaphorin-B, Sema B	
Gene ID:	64218	
UniProt:	Q9H3S1	

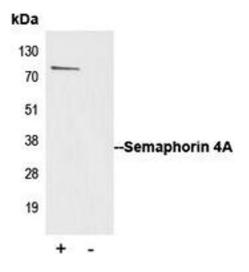
# **Application Details**

Application Notes:	WB (1:500 - 1:1000), IF/IC (1:100 - 1:500), IP (1:10 - 1:100)
Restrictions:	For Research Use only

# Handling

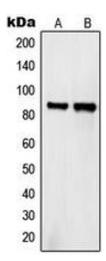
Format:	Liquid
Buffer:	Liquid in 0.42 % Potassium phosphate, 0.87 % Sodium chloride, pH 7.3, 30 % glycerol, and 0.01 % sodium azide.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.
Expiry Date:	12 months

### **Images**



# Immunoprecipitation

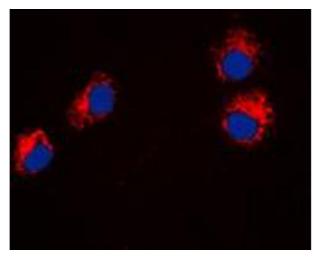
**Image 1.** Immunoprecipitation of Semaphorin 4A from 0.5mg Jurkat whole cell extract lysate, using 5ug of Anti-Semaphorin 4A Antibody and 50ul of protein G magnetic beads (+). No antibody was added to the control (-). The antibody was incubated under agitation with Protein G beads for 10min, Jurkat whole cell extract lysate diluted in RIPA buffer was added to each sample and incubated for a further 10min under agitation. Proteins were eluted by addition of 40ul SDS loading buffer and incubated for 10min



at 70°C; 10ul of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with Anti-Semaphorin 4A Antibody.

#### **Western Blotting**

**Image 2.** Western blot analysis of Semaphorin 4A expression in HeLa (A), COS7 (B) whole cell lysates.



#### **Immunofluorescence**

Image 3. Immunofluorescent analysis of Semaphorin 4A 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).