

Product Data Sheet

Anti-Semaphorin 3C Antibody

Catalog #	Source	Reactivity	Applications		
CPA6213	Rabbit	H, M, R, B, D, P	WB, IH		
Description	Rab	Rabbit polyclonal antibody to Semaphorin 3C			
Immunogen	KLH	-conjugated synthetic pep	tide encompassing a sequence within the C-term		
	regi	on of human Semaphorin	3C. The exact sequence is proprietary.		
Purification	The	antibody was purified by	immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous level	s of Semaphorin 3C protein.		
Clonality	Poly	rclonal			
Conjugation					
Form	Liqu	id in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/50 -	1/200)		
Gene Symbol	SEM	IA3C			
Alternative N	ames SEM	IAE; Semaphorin-3C; Sem	aphorin-E; Sema E		
Entrez Gene	105	12 (Human); 20348 (Mou	se)		
SwissProt	Q99	985 (Human); Q62181 (N	ouse)		
Storage/Stabi	lity Ship	pped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	free	ze/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

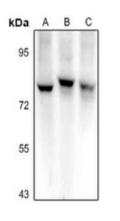
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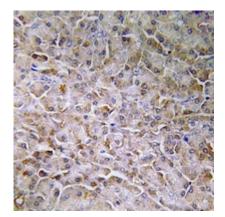
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Western blot analysis of Semaphorin 3C expression in C6 (A), MCF7 (B), A375 (C) whole cell lysates. (Predicted band size: 85 kD; Observed band size: 75 kD)



Immunohistochemical analysis of Semaphorin 3C staining in human pancreas 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.

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