

Product Data Sheet

Anti-Tachykinin Receptor 1 Antibody

Catalog #	Source	Reactivity	Applications			
CPA4268	Rabbit	H, M, R, C, D	WB, IF/IC			
Description	Rab	bit polyclonal antibody t	o Tachykinin Receptor 1			
Immunogen	KLH	-conjugated synthetic pe	ptide encompassing a sequence within the center			
	regi	on of human Tachykinin	Receptor 1. The exact sequence is proprietary.			
Purification	The	antibody was purified b	y immunogen affinity chromatography.			
Specificity	Reco	ognizes endogenous leve	els of Tachykinin Receptor 1 protein.			
Clonality	Poly	vclonal				
Form	Liqu	iid in 0.42% Potassium p	hosphate, 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	TAC	R1				
Alternative Na	ames NK1	R; TAC1R; Substance-P r	eceptor; SPR; NK-1 receptor; NK-1R; Tachykinin receptor			
	1					
Entrez Gene	686	6869 (Human); 21336 (Mouse); 24807 (Rat)				
SwissProt	P25	103 (Human); P30548 (N	1ouse); P14600 (Rat)			
Storage/Stabi	lity Ship	pped at 4°C. Upon delive	ry 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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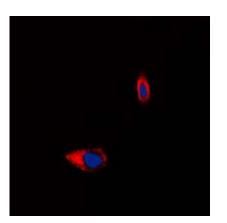
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A B

For research purposes only, not for human use

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Western blot analysis of Tachykinin Receptor 1 expression in COLO205 (A), mouse brain (B) whole cell lysates.



Immunofluorescent analysis of Tachykinin Receptor 1 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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