

## **Product Data Sheet**

## Anti-Synaptophysin Antibody

Catalog #	Source	Reactivity	Applications		
CPA2132	Rabbit	H, M, R, B, Mk, P	WB, IH, IF/IC		
Description	R	Rabbit polyclonal antibody to Synaptophysin			
Immunogen	K	LH-conjugated synthetic peptide	e encompassing a sequence within the center		
	re	region of human Synaptophysin. The exact sequence is proprietary.			
Purification	TI	he antibody was purified by imr	nunogen affinity chromatography.		
Specificity	R	ecognizes endogenous levels of	Synaptophysin protein.		
Clonality	Po	olyclonal			
Conjugation					
Form	Li	iquid in 0.42% Potassium phosp	hate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	aı	nd 0.01% sodium azide.			
Dilution	W	VB (1/500 - 1/1000), IH (1/50 - 1/1	00), IF/IC (1/50 - 1/200)		
Gene Symbol	S	YP			
Alternative N	ames Sy	ynaptophysin; Major synaptic ve	esicle protein p38		
Entrez Gene	68	855 (Human); 20977 (Mouse); 2	24804 (Rat)		
SwissProt	P	08247 (Human); Q62277 (Mous	e); P07825 (Rat)		
Storage/Stabi	lity Sł	hipped at 4°C. Upon delivery ali	quot and store at -20°C for one year. Avoid		
	fr	reeze/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

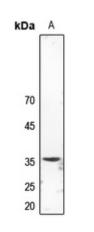
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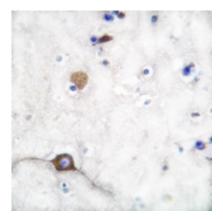
# **Cohesion**

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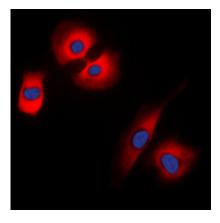
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Western blot analysis of Synaptophysin expression in rat muscle (A) whole cell lysates. (Predicted band size: 33 kD; Observed band size: 38 kD)



Immunohistochemical analysis of Synaptophysin staining in human brain 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.



Immunofluorescent analysis of Synaptophysin 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).

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