

Product Data Sheet

Anti-Syntaxin 1A Antibody

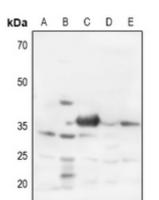
Catalog #	Source	Reactivity	Applications		
CPA3331	Rabbit	H, M, R, B, Mk	WB, IH, IF/IC		
Description	Rabb	Rabbit polyclonal antibody to Syntaxin 1A			
Immunogen	KLH-	conjugated synthetic pep	tide encompassing a sequence within the N-term		
	regio	n of human Syntaxin 1A.	The exact sequence is proprietary.		
Purification	The a	antibody was purified by	immunogen affinity chromatography.		
Specificity	Reco	gnizes endogenous levels	s of Syntaxin 1A protein.		
Clonality	Polyc	lonal			
Conjugation					
Form	Liqui	d in 0.42% Potassium pho	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and ().01% sodium azide.			
Dilution	WB (1/500 - 1/1000), IH (1/50 -	1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	STX1	A			
Alternative Na	ames STX1	; Syntaxin-1A; Neuron-sp	ecific antigen HPC-1		
Entrez Gene	6804	(Human); 20907 (Mouse	e); 116470 (Rat)		
SwissProt	Q166	523 (Human); O35526 (M	ouse); P32851 (Rat)		
Storage/Stabi	lity Shipp	oed at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	freez	e/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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Western blot analysis of Syntaxin 1A expression in HEK293T (A), Hela (B), mouse brain (C), mouse heart (D), mouse liver (E) whole cell lysates. (Predicted band size: 33 kD; Observed band size: 33 kD)



Immunohistochemical analysis of Syntaxin 1A 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 Syntaxin 1A staining in NIH3T3 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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