

Product Data Sheet

Anti-APBA2 Antibody

Catalog #	Source	Reactivity	Applications		
CPA3129	Rabbit	H, M, R	WB, IH, IF/IC		
Description		Rabbit polyclonal antibody t	o APBA2		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
		region of human APBA2. The exact sequence is proprietary.			
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	els of APBA2 protein.		
Clonality Polyclonal					
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol		APBA2			
Alternative Na	ames	MINT2; X11L; Amyloid beta	A4 precursor protein-binding family A member 2;		
		Adapter protein X11beta; N	euron-specific X11L protein; Neuronal		
		Munc18-1-interacting prote	in 2; Mint-2		
Entrez Gene		321 (Human); 11784 (Mouse	e); 83610 (Rat)		
SwissProt		Q99767 (Human); P98084 (M	Aouse); O35431 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
		freeze/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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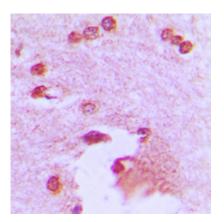
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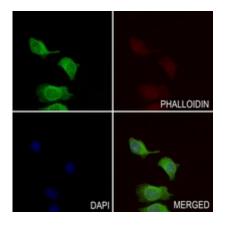
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Western blot analysis of APBA2 expression in H4 (A), HeLa (B) whole cell lysates. (Predicted band size: 82 kD; Observed band size: 82 kD)



Immunohistochemical analysis of APBA2 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 APBA2 staining in A549 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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