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anti-APLP1 antibody (AA 505-539)

2 Images



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Overview		
Quantity:	200 μL	
Target:	APLP1	
Binding Specificity:	AA 505-539	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This APLP1 antibody is un-conjugated	
Application:	Western Blotting (WB), Flow Cytometry (FACS)	
Product Details		
Immunogen:	This APLP1 antibody is generated from a rabbit immunized with a KLH conjugated synthetic	
	peptide between 505-539 amino acids from the human region of human APLP1.	
Clone:	RB58020	
Isotype:	lg Fraction	
Purification:	This antibody is purified through a protein A column, followed by peptide affinity purification.	
Target Details		
Target:	APLP1	
Alternative Name:	APLP1 (APLP1 Products)	
Background:	May play a role in postsynaptic function. The C-terminal gamma-secretase processed	

Target Details

fragment, ALID1, activates transcription activation through APBB1 (Fe65) binding (By similarity). Couples to JIP signal transduction through C-terminal binding. May interact with cellular G-protein signaling pathways. Can regulate neurite outgrowth through binding to components of the extracellular matrix such as heparin and collagen I.

Molecular Weight: 72176

UniProt: P51693

Pathways: cAMP Metabolic Process

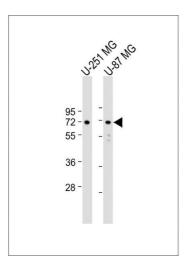
Application Details

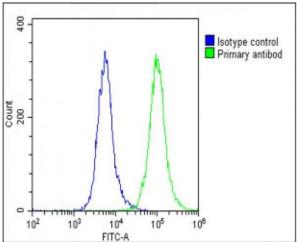
Application Notes: WB: 1:2000. FC: 1:25

Restrictions: For Research Use only

Handling

Format:	Liquid	
Buffer:	Purified polyclonal antibody supplied in PBS with 0.09 % (W/V) sodium azide.	
Preservative:	Sodium azide	
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.	
Storage:	4 °C,-20 °C	
Expiry Date:	6 months	





Western Blotting

Image 1. All lanes: Anti-LP1 Antibody (C-Term) at 1:2000 dilution Lane 1: U-251 MG whole cell lysate Lane 2: U-87 MG whole cell lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size: 72 kDa Blocking/Dilution buffer: 5 % NFDM/TBST.

Flow Cytometry

Image 2. Overlay histogram showing U-87 MG cells stained with (ABIN6243047 and ABIN6578879)(green line). The cells were fixed with 2% paraformaldehyde (10 min) and then permeabilized with 90% methanol for 10 min. The cells were then icubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody ((ABIN6243047 and ABIN6578879), 1:25 dilution) for 60 min at 37 °C. The secondary antibody used was Goat-Anti-Rabbit IgG, DyLight® 488 Conjugated Highly Cross-Adsorbed(1583138) at 1/200 dilution for 40 min at 37 °C. Isotype control antibody (blue line) was rabbit IgG1 (1 μ g/1x10^6 cells) used under the same conditions. Acquisition of >10,000 events was performed.