For Research Use Only

Phospho-MARCKS (Ser167/170) Polyclonal antibody



Catalog Number: 29145-1-AP

Basic Information

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Size: 900 µg/ml Source: Rabbit

Isotype: IgG GenBank Accession Number:

BC089040
GeneID (NCBI):
4082
UNIPROT ID:
P29966
Full Name:

myristoylated alanine-rich protein

kinase C substrate Calculated MW: 32 kDa Observed MW: 80 kDa Purification Method: Antigen affinity purification Recommended Dilutions:

WB 1:2000-1:10000 IF/ICC 1:200-1:800

Applications

Tested Applications: WB, IF/ICC, ELISA Species Specificity: ...

Positive Controls:

WB: λ phosphatase treated HeLa cells, IF/ICC: λ phosphatase treated HeLa cells,

Background Information

The Myristoylated Alanine Rich C-Kinase Substrate (MARCKS) is a ubiquitous, highly conserved protein among vertebrates, which is essential for postnatal survival, and has been widely studied for its functions in the brain and nervous system. Being highly expressed in nervous tissue, particularly during early development but persisting in the adult, it plays numerous roles related to brain growth, neuronal migration, neurite outgrowth, neurotransmitter release, and synaptic plasticity. Protein kinase C (PKC) phosphorylates MARCKS, which converts MARCKS from a membrane-bound protein to a cytoplasmic protein. The phosphorylation site of MARCKS protein is called the effector domain (ED). Its structure is highly conserved. It can be combined with cell membrane, PKC, calcium/calmodulin-dependent kineses (CaMK) and F-actin. Studies have shown that increased membrane-bound, non-phosphorylated MARCKS might be conducive to the stabilization of synaptic morphology. Phosphorylated MARCKS protein (P-MARCKS) can regulate the stability of actin network and alter the synaptic structure. (PMID: 30655546, PMID: 30155805)

Storage

Storage:

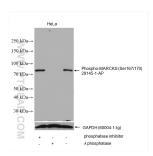
Store at -20°C. Stable for one year after shipment.

Storage Buffer

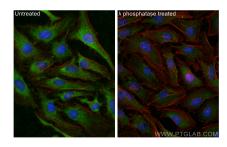
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

Aliquoting is unnecessary for -20°C storage

Selected Validation Data



Non-treated HeLa cells, phosphatase inhibitor treated and λ phosphatase treated HeLa cells were subjected to SDS PAGE followed by western blot with 29145-1-AP (Phospho-MARCKS (Ser167/170) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.



Immunofluorescent analysis of (-20°C Ethanol) fixed $^{\lambda}$ phosphatase treated HeLa cells using Phospho-MARCKS (Ser167/170) antibody (29145-1-AP) at dilution of 1:400 and CoraLite®488-Conjugated AffiniPure Goat Anti-Rabbit IgG(H+L) (SA00013-2), CL594-Phalloidin (red).