

PRODUCT DATA SHEET

Bioworld Technology,Inc.

SARM1 polyclonal antibody

Catalog: BS78063 Host: Rabbit Reactivity: Human

BackGround:

NAD(+ hydrolase, which plays a key role in axonal degeneration following injury by regulating NAD(+ metabolism. Acts as a negative regulator of MYD88- and TRIF-dependent toll-like receptor signaling pathway by promoting Wallerian degeneration, an injury-induced form of programmed subcellular death which involves degeneration of an axon distal to the injury site. Wallerian degeneration is triggered by NAD(+ depletion: in response to injury, SARM1 is activated and catalyzes cleavage of NAD(+ into ADP-D-ribose (ADPR, cyclic ADPR (cADPR and nicotinamide; NAD(+ cleavage promoting cytoskeletal degradation and axon destruction. Also able to hydrolyze NADP(+, but not other NAD(+-related molecules. Can activate neuronal cell death in response to stress. Regulates dendritic arborization through the MAPK4-JNK pathway (By similarity. Involved in innate immune response: inhibits both TI-CAM1/TRIF- and MYD88-dependent activation of JUN/AP-1, TRIF-dependent activation of NF-kappa-B and IRF3, and the phosphorylation of MAPK14/p38.

Product:

1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2 $\,$

Molecular Weight:

73KDa

Swiss-Prot:

Q6SZW1

Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

Applications:

WB,1:500 - 1:2000|IP,1:50 - 1:200

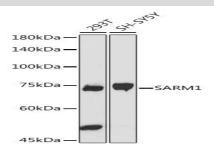
Storage&Stability:

Store at $4 \, \mathbb{C}$ short term. Aliquot and store at $-20 \, \mathbb{C}$ long term. Avoid freeze-thaw cycles.

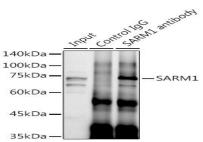
Modification:

Unmodification

DATA:



Western blot analysis of extracts of various cell lines, using SARM1 antibody at 1:1000 dilution.
Secondary antibody: HRP Goat Anti-Rabbit IgG at 1:10000 dilution.
br/>Lysates/proteins: 25ug per lane.
br/>Blocking buffer: 3% nonfat dry milk in TBST.
Detection: ECL Basic Kit .
br/>Exposure time: 90s.



Immunoprecipitation analysis of 300ug extracts of SH-SY5Y cells using 3ug SARM1 antibody. Western blot was performed from the immunoprecipitate using SARM1 antibody at a dilution of 1:1000.

Note:

For research use only, not for use in diagnostic procedure.

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