antibody and ELISA experts BOSTER BIOLOGICAL TECHNOLOGY Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

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Basic Information	
Product Name	Anti-MMD Antibody
Gene Name	MMD
Source	Rabbit
Clonality	Polyclonal
lsotype	lgG
Species Reactivity	human, mouse, rat
Tested Application	WB, FCM, ELISA
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.
Immunogen	E.coli-derived human MMD recombinant protein (Position: R20-A182). Human MMD shares 100% amino acid (aa) sequence identity with both mouse and rat MMD.
Concentration	500 ug/ml
Purification	Immunogen affinity purified.
Observed MW	28 kDa
Dilution Ratios	Western blot (WB):1:500-2000Flow Cytometry (Fixed):1:50-200Enzyme linked immunosorbent assay (ELISA):1:100-1000

Storage

12 months from date of receipt, -20°C as supplied.

Background Information

This protein is expressed by in vitro differentiated macrophages but not freshly isolated monocytes. Although sequence analysis identifies seven potential transmembrane domains, this protein has little homology to G-protein receptors and it has not been positively identified as a receptor. A suggested alternative function is that of an ion channel protein in maturing macrophages.

Reference

Anti-MMD Antibody被引用在1文献中。

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Selected Validation Data



Western blot analysis of MMD using anti-MMD antibody (A06022). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human HepG2 whole cell lysates,

Lane 2: human SH-SY5Y whole cell lysates,

Lane 3: human Caco-2 whole cell lysates,

Lane 4: human U2OS whole cell lysates,

Lane 5: rat C6 whole cell lysates,

Lane 6: rat RH35 whole cell lysates,

Lane 7: mouse Neuro-2a whole cell lysates,

Lane 8: mouse Hepa1-6 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-MMD antigen affinity purified polyclonal antibody (A06022) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for MMD at approximately 28 kDa. The expected band size for MMD is at 28 kDa.



Flow Cytometry analysis of SH-SY5Y cells using anti-MMD antibody (A06022).

Overlay histogram showing SH-SY5Y cells stained with A06022 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-MMD Antibody (A06022) at 1:100 dilution for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample (Red line) was also used as a control.