Product datasheet Anti-MYD88 Antibody (Clone#OTI1B4) Catalog Number: M00025-5

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-MYD88 Antibody (Clone#OTI1B4)
Gene Name	MYD88
Source	Mouse
Clonality	Monoclonal
lsotype	lgG2a
Species Reactivity	human
Tested Application	WB, ICC/IF, FCM
Contents	PBS (PH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Immunogen	Full length human recombinant protein of human MYD88 (NP_002459) produced in HEK293T cell.
Concentration	500 ug/ml
Purification	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)
Observed MW	33.1 kDa
Dilution Ratios	Western blot (WB):1:2000Immunocytochemistry/Immunofluorescence (ICC/IF):1:100Flow cytometry (FCM):1:100

Storage

Stable for 12 months from date of receipt. Store at -20°C as received.

Background Information

MYD88(MYELOID DIFFERENTIATION PRIMARY RESPONSE GENE 88), is a protein that, in humans, is encoded by the MYD88 gene. MyD88 is a key downstream adapter for most Toll-like receptors (TLRs) and interleukin-1 receptors (IL1Rs). And it is mapped on 3p22.2. MYD88 encodes a cytosolic adapter protein that plays a central role in the innate and adaptive immune response. This protein functions as an essential signal transducer in the interleukin-1 and Toll-like receptor signaling pathways. Overexpression of MYD88 caused an increase in the level of transcription from the interleukin-8 promoter. The C-terminal domain of MYD88 has significant sequence similarity to the cytoplasmic domain of IL1RAP. Inhibiting the IL1R-MYD88 pathway in vivo could block the damage from acute inflammation that occurs in response to sterile cell death, and do so in a way that might not compromise tissue repair or host defense against pathogens.

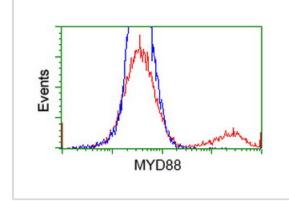
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BOSTER® antibody and ELISA experts

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Selected Validation Data



HEK293T cells transfected with either overexpress plasmid (Red) or empty vector control plasmid (Blue) were immunostained by anti-MYD88 antibody, and then analyzed by flow cytometry.