

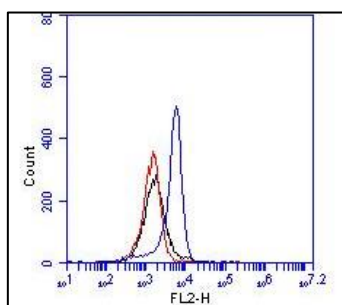
CERTIFICATE OF ANALYSIS – TECHNICAL DATA SHEET

Product name	TLR4/MD-2, Mouse, clone MTS510, biotinylated		
Catalog number	HM1029BT-50UG		
Lot number	-	Expiry date	-
Volume	500 µl	Amount	50 µg
Formulation	0.2 µm filtered in PBS+0.1%BSA+0.02%NaN3	Concentration	100 µg/ml
Host Species	Rat IgG2a	Conjugate	Biotin
Endotoxin	N.A.	Purification	Protein G
Storage	4°C		

Application notes

	IHC-F	IHC-P	IF	FC	FS	IA	IP	W
Reference #	4			1,3,5,8	2,3,6,7		2	
Yes	•			•	•		•	
No								
N.D.		•	•			•		•

N.D. = Not Determined; IHC = Immuno histochemistry; F = Frozen sections; P = Paraffin sections; IF = Immuno Fluorescence; FC = Flow Cytometry; FS = Functional Studies; IA = Immuno Assays; IP = Immuno Precipitation; W = Western blot



FC: RAW264.7 cells (105) were stained with 4µg/ml mAb for 1h at 4°C (black- isotype control, red- irrelevant mAb, Blue- HM1029)

Dilutions to be used depend on detection system applied. It is recommended that users test the reagent and determine their own optimal dilutions. The typical starting working dilution is 1:50.

- IHC-F: 6µm acetone fixed sections blocked with 0.3% hydrogen peroxidase in methanol and subsequently with normal serum. Sections were incubated for 2h at RT at 1/100 dilution of clone MTS510.
- FC: Cells were incubated with 0.1µg mAb for 30 minutes at 4°C. Postive control: RAW264.7 cells.
- FS: Cells were pre-incubated with 10µg/ml of antagonistic mAb MTS510.

General Information
Description

The monoclonal antibody MTS510 reacts with the Toll-like receptor 4 (TLR4, CD284) that is associated with MD2. TLRs are expressed by various cells of the immune system, such as macrophages and dendritic cells. TLRs are class I receptors, with a single α -helix that spans the cell membrane. They recognize and respond to molecules derived from bacterial, viral and fungal pathogens, such as lipopolysaccharide (LPS) from the outer membrane of Gram negative bacteria, peptidoglycan fragments from bacterial cell walls and single-stranded and double-stranded RNA from viruses. Toll-like receptor 4 (TLR4; CD284) has been identified, next to MD-2 and CD14, as a receptor that is central to the innate immune response to LPS of Gram-negative bacteria. TLR4 is unique among TLRs in its ability to activate two distinct signaling pathways; one pathway is activated by the adaptors TIRAP (Toll/interleukin-1- receptor (TIR)-domain-containing adaptor protein) and MyD88, which leads to the induction of pro-inflammatory cytokines. The second pathway is activated by the adaptors TRIF (TIR-domaincontaining adaptor protein inducing interferon- β) and TRAM (TRIF-related adaptor molecule), which leads to the induction of type I interferons.

MD-2 exists as a cell surface protein in association with TLR4. It also exists as secreted forms consisting of MD-2 monomer and multimers. Circulating sMD-2 is mainly present as a doublet of ~20 and 25 kD, representing differentially glycosylated forms. Unlike TLR4, sMD-2 binds directly LPS without the need of soluble CD14 (sCD14). However, LPS-

MD-2 interactions are increased when LPS is pretreated with CD14. Only monomeric sMD-2 is biologically active and able to associate with TLR4 and LPS. sMD-2 circulates in plasma of healthy individuals as a non-active, polymeric protein. In septic plasma, the total amount of sMD-2 was strongly elevated and contained both sMD-2 polymers and monomers. Soluble MD-2 is proposed to be an important mediator of organ inflammation during sepsis. During experimental human endotoxemia, the monomeric and total sMD-2 content in plasma increased with the kinetics of an acute phase protein. This parallels enhanced TLR4 costimulatory activity. In vitro studies revealed that sMD-2 release appears to be restricted to endothelial and dendritic cells.

The monoclonal antibody MTS510 reacts preferentially, especially in flow cytometry, with mouse TLR4 that is associated with MD-2. MTS510 is a TLR4 function-blocking antibody that is useful for studies on the role of TLR4 as a receptor for LPS induced cytokine production by TLR4 bearing cells. The antibody was shown to coprecipitate MD-2 (30 kDa) with TLR4 (100 kDa).

Immunogen	Ba/F3 cells expressing mouse TLR4 and MD-2 (Ref.2)
Aliases	Toll-like receptor 4: TLR4, CD284, ARMD10 MD2: ESOP-1; myeloid differentiation protein-2; lymphocyte antigen 96 (Ly-96)
References	<ol style="list-style-type: none">1. Nomura, F et al ; Cutting edge: Endotoxin tolerance in mouse peritoneal macrophages correlate with down-regulation of surface toll-like receptor 4 expression. <i>J Immunol</i> 2000, <i>164</i>: 34762. Akashi, S et al; Cutting edge: Cell surface expression and lipopolysaccharide signalling via the toll-like receptor 4-MD-2 complex on mouse peritoneal macrophages. <i>J Immunol</i> 2000, <i>164</i>: 34713. Sato, S et al; Synergy and cross-tolerance between toll-like receptor (TLR)2- and TLR4-mediated signaling pathways. <i>J Immunol</i> 2000, <i>165</i>: 70964. Ortega-Cava, C et al; Strategic compartmentalization of toll-like receptor 4 in the mouse gut. <i>J Immunol</i> 2003, <i>170</i>: 39775. Tsujimoto, H. A critical role of CpG motifs in a murine peritonitis model by their binding to highly expressed toll-like receptor-9 on liver NTK cells. <i>J hepatology</i> 2006, <i>45</i>: 8366. Cook,C et al. Lipopolysaccharide, Tumor Necrosis Factor alpha, or Interleukin-1β triggers reactivation of latent CMV in immunocompetent mice. <i>J virology</i> 2006, <i>80</i>: 91517. Tsukumo_D et al. Loss-of-function mutation in Toll-like receptor 4 prevents diet-induced obesity and insulin resistance. <i>Diabetes</i> 2007, <i>56</i>: 19868. Burton, O et al. Importance of TLR2 in the direct response of T lymphocytes to Schistosoma mansoni antigens. <i>Eur J Immunol</i> 2010, <i>40</i>: 2221
Storage&stability	Product should be stored at 4°C. Under recommended storage conditions, product is stable for at least one year.
Precautions	For research use only. Not for use in or on humans or animals or for diagnostics. It is the responsibility of the user to comply with all local/state and federal rules in the use of this product. Hycult Biotech is not responsible for any patent infringements that might result from the use or derivation of this product.

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