

**CD282 (Toll-Like Receptor 2) Antibody - With BSA and Azide**  
**Mouse Monoclonal Antibody [Clone TLR2/221 ]**  
**Catalog # AH12424**

**Specification**

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**CD282 (Toll-Like Receptor 2) Antibody - With BSA and Azide - Product Information**

Application	,2,3,4,
Primary Accession	<a href="#">O60603</a>
Other Accession	<a href="#">7097</a> , <a href="#">519033</a>
Reactivity	Human, Monkey, Drosophila
Host	Mouse
Clonality	Monoclonal
Isotype	Mouse / IgG2a, kappa
Calculated MW	90kDa KDa

**CD282 (Toll-Like Receptor 2) Antibody - With BSA and Azide - Additional Information**

**Gene ID** 7097

**Other Names**

Toll-like receptor 2, Toll/interleukin-1 receptor-like protein 4, CD282, TLR2, TIL4

**Storage**

Store at 2 to 8°C. Antibody is stable for 24 months.

**Precautions**

CD282 (Toll-Like Receptor 2) Antibody - With BSA and Azide is for research use only and not for use in diagnostic or therapeutic procedures.

**CD282 (Toll-Like Receptor 2) Antibody - With BSA and Azide - Protein Information**

**Name** TLR2 ([HGNC:11848](#))

**Synonyms** TIL4

**Function**

Cooperates with LY96 to mediate the innate immune response to bacterial lipoproteins and other microbial cell wall components. Cooperates with TLR1 or TLR6 to mediate the innate immune response to bacterial lipoproteins or lipopeptides (PubMed: [17889651](http://www.uniprot.org/citations/17889651), PubMed: [21078852](http://www.uniprot.org/citations/21078852)). Acts via MYD88 and TRAF6, leading to NF-kappa-B activation, cytokine secretion and the inflammatory response. May also activate immune cells and promote apoptosis in response to the lipid moiety of lipoproteins (PubMed: [10426995](http://www.uniprot.org/citations/10426995), PubMed: [10426996](http://www.uniprot.org/citations/10426996)). Recognizes mycoplasmal macrophage-activating lipopeptide-2kD (MALP-2), soluble tuberculosis factor (STF), phenol-soluble modulins (PSM) and B.burgdorferi outer surface protein A lipoprotein (OspA-L) cooperatively with TLR6 (PubMed: [10426996](#)).

<http://www.uniprot.org/citations/11441107> target="\_blank">11441107</a>). Stimulation of monocytes in vitro with M.tuberculosis PstS1 induces p38 MAPK and ERK1/2 activation primarily via this receptor, but also partially via TLR4 (PubMed:<a href="http://www.uniprot.org/citations/16622205" target="\_blank">16622205</a>). MAPK activation in response to bacterial peptidoglycan also occurs via this receptor (PubMed:<a href="http://www.uniprot.org/citations/16622205" target="\_blank">16622205</a>). Acts as a receptor for M.tuberculosis lipoproteins LprA, LprG, LpqH and PstS1, some lipoproteins are dependent on other coreceptors (TLR1, CD14 and/or CD36); the lipoproteins act as agonists to modulate antigen presenting cell functions in response to the pathogen (PubMed:<a href="http://www.uniprot.org/citations/19362712" target="\_blank">19362712</a>). M.tuberculosis HSP70 (dnaK) but not HSP65 (groEL-2) acts via this protein to stimulate NF-kappa-B expression (PubMed:<a href="http://www.uniprot.org/citations/15809303" target="\_blank">15809303</a>). Recognizes M.tuberculosis major T-antigen EsxA (ESAT-6) which inhibits downstream MYD88-dependent signaling (shown in mouse) (By similarity). Forms activation clusters composed of several receptors depending on the ligand, these clusters trigger signaling from the cell surface and subsequently are targeted to the Golgi in a lipid-raft dependent pathway. Forms the cluster TLR2:TLR6:CD14:CD36 in response to diacylated lipopeptides and TLR2:TLR1:CD14 in response to triacylated lipopeptides (PubMed:<a href="http://www.uniprot.org/citations/16880211" target="\_blank">16880211</a>). Required for normal uptake of M.tuberculosis, a process that is inhibited by M.tuberculosis LppM (By similarity).

#### Cellular Location

Membrane {ECO:0000250|UniProtKB:Q9QUN7}; Single-pass type I membrane protein. Cytoplasmic vesicle, phagosome membrane {ECO:0000250|UniProtKB:Q9QUN7}; Single-pass type I membrane protein. Membrane raft. Note=Does not reside in lipid rafts before stimulation but accumulates increasingly in the raft upon the presence of the microbial ligand. In response to diacylated lipoproteins, TLR2:TLR6 heterodimers are recruited in lipid rafts, this recruitment determines the intracellular targeting to the Golgi apparatus. Triacylated lipoproteins induce the same mechanism for TLR2:TLR1 heterodimers.

#### Tissue Location

Highly expressed in peripheral blood leukocytes, in particular in monocytes, in bone marrow, lymph node and in spleen. Also detected in lung and in fetal liver. Levels are low in other tissues

### CD282 (Toll-Like Receptor 2) Antibody - With BSA and Azide - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

### CD282 (Toll-Like Receptor 2) Antibody - With BSA and Azide - Images

### CD282 (Toll-Like Receptor 2) Antibody - With BSA and Azide - Background

This MAbs reacts with human Toll-like receptor 2 (TLR2). It is a member of the Toll-like receptor (TLR) family, which plays a fundamental role in pathogen recognition and activation of innate immunity. TLRs are highly conserved from Drosophila to humans and share structural and functional similarities. They recognize pathogen-associated molecular patterns (PAMPs) that are expressed on infectious agents, and mediate the production of cytokines necessary for the

development of effective immunity. The various TLRs exhibit different patterns of expression. This gene is expressed most abundantly in peripheral blood leukocytes, and mediates host response to Gram-positive bacteria and yeast via stimulation of NF-kappaB.

#### **CD282 (Toll-Like Receptor 2) Antibody - With BSA and Azide - References**

Lien E, Sellati TJ, Yoshimura A, Flo TH, Rawadi G, Finberg RW, Carroll JD, Espevik T, Ingalls RR, Radolf JD, Golenbock DT. Toll-like receptor 2 functions as a pattern recognition receptor for diverse bacterial products. *J Biol Chem.* 1999 Nov 19;274(47):33419-25