

Mouse TLR2 Biotinylated Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: BAF1530

DESCRIPTION		
Species Reactivity	y Mouse	
Specificity	Detects mouse TLR2 in Western blots. In Western blots, approximately 5% cross-reactivity with recombinant human (rh) TLR2 observed and less than 1% cross-reactivity with recombinant mouse (rm) TLR1, rhTLR3, rmTLR4 and rmTLR6 is observed.	is
Source	Polyclonal Goat IgG	
Purification	Antigen Affinity-purified	
Immunogen	Mouse myeloma cell line NS0-derived recombinant mouse TLR2 Gln25-Leu590 Accession # Q9QUN7	
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with BSA as a carrier protein. See Certificate of Analysis for details.	
APPLICATIONS Please Note: Optimal diluti	filutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.	
	Recommended Sample Concentration	
Western Blot	0.1 μg/mL Recombinant Mouse TLR2 Fc Chimera (Catalog # 1530-TR)	
PREPARATION AND	D STORAGE	
Reconstitution	Reconstitute at 0.2 mg/mL in sterile PBS.	
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.	
Stability & Storage	 Use a manual defrost freezer and avoid repeated freeze-thaw cycles. 12 months from date of receipt, -20 to -70 °C as supplied. 1 month, 2 to 8 °C under sterile conditions after reconstitution. 6 months, -20 to -70 °C under sterile conditions after reconstitution. 	

BACKGROUND

The Toll-like family of molecules are a group of integral membrane proteins that serve as pattern recognition receptors for microbial pathogens (1-4). To date, there are at least eleven mouse and ten human members that activate the innate immune system following exposure to a variety of microbial species (1, 3). All Toll-like receptors (TLRs) are type I transmembrane (TM) proteins that exist either in the plasma membrane or in the membranes of endosomal structures (where they bind intracellular nucleic acids) (3). All TLRs also contain a large number of extracellular leucine-rich repeats (LRRs) and a cytoplasmic tail with a Toll/IL-1 receptor (TIR) domain. Mouse Toll-like receptor-2 (TLR2) is a 97 kDa, 760 amino acid (aa) glycoprotein that contains a 563 aa extracellular region, a 21 aa TM segment, and a 176 aa cytoplasmic domain (5, 6). The extracellular region contains 16 leucine-rich repeats, while the cytoplasmic tail shows one 146 aa TIR domain. The receptor is expressed on a number of cell types including T cells ($\alpha\beta$ and $\gamma\delta$), monocytes, dendritic cells, neutrophils, B cells, endothelial cells, mast cells, NK cells, macrophages and hepatocytes (1, 4, 5, 7, 8). TLR2 functions as part of a heterodimeric complex with either TLR1 or TLR6 (1, 3, 4). These complexes recognize lipoproteins and glycolipids from gram-positive and gram-negative bacteria as well as mycoplasma and yeast. TLR2/TLR1 heterodimers recognize triacylated lipopeptides from a variety of microorganisms. The TLR2/TLR6 heterodimer preferentially recognizes diacylated lipopeptides (9). Biglycan is also known to activate TLR2, but the context is unclear (8). Notably, in human, TLR2 also dimerizes with TLR10. But the TLR10 gene in mouse (but not rat) is mutationally inactive, and thus this complex is unclear (10). Upon ligand recognition, TLR2 delivers an activating signal via the associated adapter molecules, MyD88 and TIRAP (1, 11). Activation via TLR2 also demorphic in rat, human, hamster and canine, respectively.

References:

- 1. Wetzler, L. (2003) Vaccine **21**:S2/55.
- 2. Netea, M. et al. (2004) J. Leukoc. Biol. **75**:749.
- 3. Dunne, A. and L. O'Neill (2005) FEBS. Lett. 579:3330.
- 4. Hopkins, P.A. and S. Sriskandan (2005) Clin. Exp. Immunol. 140:395.
- 5. Matsuguchi, T. et al. (2000) Blood 95:1378.
- 6. Meng, G. et al. (2005) Immunol. Lett. 98:200.
- 7. Flo T. et al. (2001) J. Leukoc. Biol. 69:474.
- 8. Schaefer, L. et al. (2005) J. Clin. Invest. 115:2223.
- 9. Akira S. (2003) Curr. Opin. Immunol. **15**:5.
- 10. Hasan, U. et al. (2005) J. Immunol. 174:2942.
- 11. Yamamoto M. et al. (2002) Nature 420:324.

