

Human IL-27 Rα/WSX-1/TCCR Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: AF1479

DESCRIPTION			
Species Reactivity	Human		
Specificity	Detects human IL-27 Rα/WSX-1/TCCR in direct ELISAs and Western blots. In direct ELISAs, approximately 20% cross-reactivity with recombinant mouse TCCR is observed.		
Source	Polyclonal Goat IgG		
Purification	Antigen Affinity-purified		
lmmunogen	Mouse myeloma cell line NS0-derived recombinant human IL-27 Rα/WSX-1/TCCR Gly34-Lys516 Accession # Q6UWB1		
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details.		

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Western Blot	0.1 μg/mL	Recombinant Human IL-27 Rα/WSX-1/TCCR Fc Chimera (Catalog # 1479-TC)
Immunohistochemistry	5-15 μg/mL	Immersion fixed paraffin-embedded sections of human lymph node

PREPARATION AND STORAGE

Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles.	
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.	
Reconstitution	Reconstitute at 0.2 mg/mL in sterile PBS.	

- 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

IL-27 Rα (also known as WSX-1 and TCCR) is a 96-100 kDa member of the type I, group 2 cytokine receptor family (1, 2, 3, 4, 5, 6). Mature IL-27 Rα is a type I transmembrane glycoprotein that contains a 484 amino acid (aa) extracellular region, a 21 aa transmembrane segment and a 99 aa cytoplasmic domain. Consistent with type I cytokine receptors, the extracellular region contains four positionally conserved cysteine residues, a WSxWS motif (for receptor folding and ligand binding), and three fibronectin type III repeats. The intracellular domain contains a "box-1" motif that may be involved with Janus kinases (3). One potential alternate splice form has been hypothesized that involves a 58 aa addition to the cytoplasmic domain and, based on mouse, a soluble 33 kDa splice form that shows a 20 aa substitution for aa 257-636 may also occur in human (3, 7). The human IL-27 Rα extracellular region shares 63% amino acid identity with the mouse IL-27 Rα extracellular domain (2, 3). IL-27 Rα is expressed in mast cells, endothelial cells, NK cells, macrophages, monocytes, B cells, dendritic cells, and naïve T cells (1, 2, 4, 8). Typical of other class I cytokine receptor chains, the ligand binding IL-27 Rα molecule is known to heterodimerize with a signal-transducing subunit (gp130) to form a functional IL-27 receptor (9, 10). In addition, IL-27 Rα is reported to complex with CNTFRα and gp130 form a humanin receptor on neurons (7, 11), and to complex with gp130 and IL-6 R to form a receptor for a p28:CLF heterodimeric cytokine on lymphocytes (12). Studies using IL-27 Rα/WSX-1-/- mice reveal that IL-27 known to act on naïve T cells, blocking their differentiation into a Th17 phenotype. Notably, cells committed to a Th17 phenotype, although they express a functional IL-27 receptor, are unresponsive to the effects of IL-27 (15). Activated T cells that are CD4+ and CD8+, and which express the IL-27 receptor, can be induced by IL-27 to form a double-positive CD25+ FoxP3- IFN-γ plus IL-10 secreting phenotype that both promot

References:

- 1. Villarino, A.V. et al. (2004) J. Immunol. 173:715.
- 2. Chen, Q. et al. (2000) Nature 407:916.
- 3. Sprecher, C.A. et al. (1998) Biochem. Biophys. Res. Commun. 246:82.
- 4. Artis, D. et al. (2004) J. Immunol. 173:5626.
- 5. Yoshida, H. & Y. Miyazaki (2008) Int. J. Biochem. Cell Biol. 40:2379.
- Yoshida, H. & M. Yoshiyuki (2008) Immunol. Rev. 226:234.
- 7. Hashimoto, Y. et al. (2009) Biochem. Biophys. Res. Commun. 389:95.
- 8. Holscher, C. et al. (2005) J. Immunol. 174:3534.
- 9. Pflanz, S. et al. (2004) J. Immunol. 172:2225.
- 10. Scheller, J. et al. (2005) Biochem. Biophys. Res. Commun. 326:724.
- 11. Hashimoto, Y. et al. (2009) Mol. Biol. Cell 20:2864.
- 12. Crabe, S. et al. (2009) J. Immunol. 183:7692.
- 13. Villarino, A. et al. (2003) J. Immunol. 170:645.
- 14. Hamano., S. et al. (2003) Immunity 19:657.
- 15. El-behi, M. *et al.* (2009) J. Immunol. **183**:4957.
- 16. Fitzgerald, D.C. et al. (2007) Nat. Immunol. 8:1372.

