

Human IL-17D Biotinylated Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: BAF1504

DESCRIPTION			
Species Reactivity	Human		
Specificity	Detects human IL-17D in ELISAs and Western blots. In sandwich immunoassays, approximately 6% cross-reactivity with recombinant mous (rm) IL-17D is observed and less than 0.02% cross-reactivity with recombinant human (rh) IL-17, rhIL-17B, rhIL-17C, rhIL-17E, rhIL-17F, rmIL 17, and rmIL-17C is observed.		
Source	Polyclonal Goat IgG		
Purification	Antigen Affinity-purified		
Immunogen	E. coli-derived recombinant human IL-17D Ala18-Pro202 Accession # Q8TAD2		
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 dilut	ions should be determined by each laboratory for each applicat	tion. General Protocols are available in the Technical Information section on our website.	
	Recommended Concentration	Sample	
Western Blot	0.1 µa/ml	Decembinant Human II. 47D (Catalan # 4504 II.)	

	Concentration	
Western Blot	0.1 µg/mL	Recombinant Human IL-17D (Catalog # 1504-IL)
Human IL-17D Sandwich Immunoassay		Reagent
ELISA Capture	2-8 µg/mL	Human IL-17D Antibody (Catalog # MAB1504)
ELISA Detection	0.1-0.4 μg/mL	Human IL-17D Biotinylated Antibody (Catalog # BAF1504)
Standard		Recombinant Human IL-17D (Catalog # 1504-IL)
PREPARATION AND STORAGE		

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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	e 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 Interleukin-17 (IL-17) family proteins, comprising six members (IL-17, IL-17B through IL-17F), are secreted, structurally related proteins that share a conserved cysteine-knot fold near the C-terminus, but have considerable sequence divergence at the N-terminus (1, 2). With the exception of IL-17B, which exists as a non-covalently linked dimer, all IL-17 family members are disulfide-linked dimers (3). IL-17 family proteins are pro-inflammatory cytokines that induce local cytokine production and are involved in the regulation of immune functions (1, 2). Two receptors (IL-17 R, and IL-17B R), which are activated by IL-17 family members, have been identified. In addition, at least three additional orphan type I transmembrane receptors with homology to IL-17 R, including IL-17 RL (IL-17 RC), IL-17 RD, and IL-17 RE, have also been reported (1-4). The functions of IL-17 RC, D, and E are not known.

Human IL-17D cDNA encodes a 202 amino acid (aa) residues protein with a putative 17 aa signal peptide (5). Human and mouse IL-17D share 78% sequence identity. Among IL-17 family members, IL-17D is most closely related to IL-17B, sharing 27% aa sequence homology (5, 6). IL-17D is expressed preferentially in skeletal muscle, heart, adipose tissue, lung, pancreas, and nervous system (1, 5). Like other IL-17 family members, IL-17D modulates immune responses indirectly by stimulating the production of myeloid growth factors and chemokines including IL-6, IL-8, and GM-CSF (5). IL-17D has also been shown to suppress the proliferation of myeloid progenitors in colony formation assays. The receptor of IL-17D has not yet been identified. However, stimulation of IL-8 production by IL-17D is mediated through the activation of nuclear factor kappa-B (5). The IL-17D preparations from R&D Systems have been found to bind immobilized recombinant IL17B R/Fc in a functional ELISA.

References:

- 1. Aggarwal, S. and A.L. Gurney (2002) J. Leukoc. Biol. 71:1.
- 2. Moseley, T.A. et al. (2003) Cytokine & Growth Factor Rev. 14:155.
- 3. Hymowitz, S.G. et al. (2001) EMBO J. 20:5332.
- 4. Haudenschild, D. et al. (2002) J. Biol. Chem. 277:4309.
- 5. Starnes, T. et al. (2002) J. Immunol. 169:642.
- 6. Li, H. et al. (2000) Proc. Natl. Acad. Sci. USA **97**:773.

