Human IL-17A Neutralizing (D13C2) Rabbit mAb

🗹 100 μg

Sterile

:12739

rev. 12/27/13



Species Cross-Reactivity: H

Description: Neutralizing antibodies can be used to inhibit normal biological function through their binding to biological molecules. These reagents can be used to determine the effects that a particular molecule has in biological systems. Human IL-17A Neutralizing (D13C2) Rabbit mAb has been shown to neutralize the production of IL-6 from human dermal fibroblast cells *in vitro* with an ND₅₀ in the range of 8-50 ng/ml.

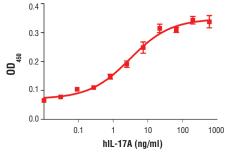
Background: IL-17A is a cystine-linked homodimeric pro-inflammatory cytokine produced by Th₁₇ cells, a distinct CD4+ T cell lineage (1,2). IL-17A stimulates the production of the pro-inflammatory cytokines IL-1β, TNF-α, and IL-6. IL-17A also induces production of the neutrophil chemoattractants IL-8, CXCL1, and CXCL6 thereby bridging adaptive and innate immunity (1,2). IL-17A is intimately involved in mucosal immunity against bacterial infections (1,3) and has a putative role in some autoimmune disorders (1,4). IL-17A effects appear to be exerted primarily through binding to the IL-17RA (5). IL-17A binding induces production of cytokines, chemokines and other proteins through activation of the Erk1/2 MAP kinase, PI3K/Akt, p38, and NF-κB pathways (3,4,6).

Endotoxin: Less than 0.1 EU/µg of antibody.

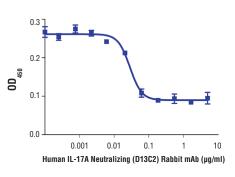
Specificity/Sensitivity: Human IL-17A Neutralizing (D13C2) Rabbit mAb binds to human IL-17A (hIL-17A) and neutralizes its ability to induce IL-6 production in human dermal fibroblast cells. This antibody shows 50% cross- reactivity with human IL-17A/F. This antibody does not cross-react with mouse IL-17A, human IL-25 or mouse IL-25.

Source/Purification: Monoclonal antibody is produced by immunizing animals with a recombinant human IL-17A protein.

Notes on Use: CST recommends incubation of the neutralizing antibody with the intended target for 1 hr at 37°C before addition to the experiment at an optimal concentration determined by the user



The production of IL-6 by human dermal fibroblasts cultured with increasing concentrations of Human IL-17A (hIL-17A) #8928 was assessed. Media from cells incubated with IL-17A for 48 hours was collected and assayed for IL-6 by ELISA and the OD₄₅₀ was determined.



The ability of Human IL-17A Neutralizing (D13C2) Rabbit mAb to inhibit IL-17A-induced IL-6 production was assessed. Human dermal fibroblasts were incubated with increasing concentrations of antibody in the presence of Human Interleukin-17A (hIL-17A) #8928 (15 ng/mI). After 48 hr, culture supernatants were harvested and assayed for IL-6 by ELISA and the OD_{450} was determined.

Entrez-Gene ID #3605 Swiss-Prot Acc. #Q16552

Formulation: Lyophilized from a 0.2 μm filtered solution in 10 mM HEPES with trehalose.

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Reconstitution: Add sterile 10 mM HEPES pH 7.0 to a final concentration of greater than 50 µg/ml. Solubilize for 20 min at room temperature with occasional gentle vortexing.

Storage: Store lyophilized material at -20°C. After reconstitution, recommended storage at 4°C for 1 month or -20°C for 6 months. *Avoid repeated freeze/thawing.*

Background References:

- (1) Kolls, J.K. and Lindén, A. (2004) Immunity 21, 467-76.
- (2) Liang, S.C. et al. (2006) J Exp Med 203, 2271-9.
- (3) Dubin, P.J. and Kolls, J.K. (2008) *Immunol Rev* 226, 160-71.
- (4) Zrioual, S. et al. (2009) J Immunol 182, 3112-20.
- (5) Wright, J.F. et al. (2008) J Immunol 181, 2799-805.
- (6) Rahman, M.S. et al. (2006) J Immunol 177, 4064-71.

 Applications Key:
 W—Western
 IP—Immunoprecipitation
 IHC—Immunohistochemistry
 ChIP—Chromatin Immunoprecipitation
 IF—Immunofluorescence
 F—Flow cytometry
 E-P—ELISA-Peptide

 Species Cross-Reactivity Key:
 H—human
 M—mouse
 R—rat
 Hm—hamster
 Mk—monkey
 Mi—mink
 C—chicken
 Dm—D. melanogaster
 X—zebrafish
 B—bovine

 Dg—dog
 Pg—pig
 Sc—S. cerevisiae
 Cerevisiae
 Cerevisiae
 Cerevisiae
 AII—all species expected
 Species enclosed in parentheses are predicted to react based on 100% homology.