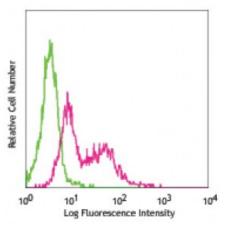
Product Data Sheet

PerCP/Cy5.5 anti-mouse IL-17A

Catalog # / Size:	3134595 / 25 μg 3134600 / 100 μg
Clone:	TC11-18H10.1
Isotype:	Rat IgG1, κ
Immunogen:	<i>E. coli</i> expressed, recombinant mouse IL-17A
Reactivity:	Mouse
Preparation:	The antibody was purified by affinity chromatography, and conjugated with PerCP/Cy5.5 under optimal conditions. The solution is free of unconjugated PerCP/Cy5.5 and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).
Concentration:	0.2



PMA (20 ng/ml) + ionomycin (1 microg/ml) -stimulated (6 hours + monensin, 2 μ M) mouse thymoma cell line EL-4 intracellularly stained with TC11-18H10.1 PerCP/Cy5.5

Applications:

- Applications: Flow Cytometry
- **Recommended** Usage: Each lot of this antibody is quality control tested by intracellular immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is 5 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

* PerCP/Cy5.5 has a maximum absorption of 482 nm and a maximum emission of 690 nm.

Application Notes: ELISA Capture^{3,4} and ELISPOT Capture⁵: The purified TC11-18H10.1 antibody is useful as the capture antibody in a sandwich ELISA, when used in conjunction with the biotinylated TC11-8H4 antibody (Cat. No. 507002) as the detecting antibody and recombinant mouse IL-17 (Cat. No. 576009) as the standard.

Flow Cytometry^{2-4,7,8,11,12}: The TC11-18H10.1 antibody is useful for intracellular immunofluorescent staining and flow cytometric analysis to identify IL-17-producing cells within mixed cell populations.

Neutralization^{6,9}: The LEAF^m purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for neutralization of mouse IL-17 bioactivity *in vivo* and *in vitro* (Cat. No. 506906).

Additional reported applications (for the relevant formats) include: Western blotting.

Application References:	 Kennedy J, <i>et al.</i> 1996. <i>J. Interferon Cytokine Res.</i> 16:611. Schubert D, <i>et al.</i> 2004. <i>J. Immunol.</i> 172:4503. (ICFC) Infante-Duarte C, <i>et al.</i> 2000. <i>J. Immunol.</i> 165:6107. (ICFC, ELISA Capture) Harrington LE, <i>et al.</i> 2005. <i>Nature Immunol.</i> doi:10.1038/ni1254. (ICFC, ELISA Capture) Nekrasova T, <i>et al.</i> 2005. <i>J. Immunol.</i> 175:2734. (ELISPOT Capture) Yen D, <i>et al.</i> 2006. <i>J. Clin. Invest.</i> 116:1310. (Neut)
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Description: IL-17, also known as CTLA-8, is a T cell-expressed pleiotropic cytokine that exhibits a high degree of homology to a protein encoded by the ORF13 gene of herpes virus Saimiri. IL-17 is produced by Th cells (Th17) that are distinct from the traditional Th1- and Th2-cell subsets. IL-23 plays an important role in triggering IL-17 production. Both recombinant and natural IL-17 have been shown to exist as disulfide linked homodimers. IL-17 exhibits multiple biological activities on a variety of cells including: the induction of IL-6 and IL-8 production in fibroblasts, activation of NF-κB, and costimulation of T cell proliferation. IL-17 is an essential inflammatory mediator in the development of autoimmune diseases. Neutralization of IL-17 with monoclonal antibody is able to ameliorate the disease course.

Antigen1. Fitzgerald K, et al. Eds. 2001. The Cytokine FactsBook. Academic Press SanReferences:Diego.

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