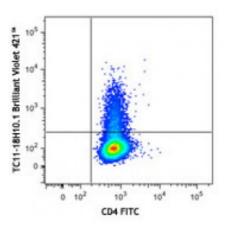
Product Data Sheet

Brilliant Violet 421[™] anti-mouse IL-17A

Catalog # / Size:	3134625 / 125 μl 3134630 / 50 μg
Clone:	TC11-18H10.1
Isotype:	Rat IgG1, к
Immunogen:	<i>E. coli</i> expressed, recombinant mouse IL-17A
Reactivity:	Mouse
Preparation:	The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 421 [™] under optimal conditions. The solution is free of unconjugated Brilliant Violet 421 [™] and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).
Concentration:	microL size: Lot-specific nd.com/concentrationlookup" t



Th17-polarized C57BL/6 mouse CD4+ lymphocytes were stimulated with PMA + lonomycin for 6 hours in the presence of monensin, stained with CD4 FITC, fixed, permeabilized, and then stained with IL-17A (clone TC11-18H10.1) Brilliant Violet 421[™] (top) or

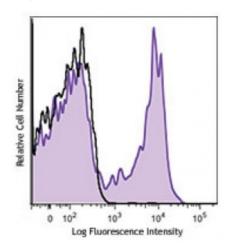
Applications:

Applications: F	low Cytometry
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Each lot of this antibody is quality Recommended Usage: control tested by intracellular immunofluorescent staining with flow cytometric analysis. For flow cytometric staining using the microL size, the suggested use of this reagent is ≤ 5 microL per million cells or 5 microL per 100 microL of whole blood. For flow cytometric staining using the microg size, the suggested use of this reagent is ≤ 0.25 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.

> Brilliant Violet 421[™] excites at 405 nm and emits at 421 nm. The standard bandpass filter 450/50 nm is recommended for detection. Brilliant Violet 421[™] is a trademark of Sirigen Group Ltd.

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Application ELISA Capture^{3,4} and **ELISPOT** Notes: 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. Cytometry^{2-4,7,8,11,12} Flow 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 LEAFTM 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.

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9. Smith E, <i>et al.</i> 2008. <i>J. Immunol.</i> 181:1357. (Neut) <u>PubMed</u> 10. Neufert C, <i>et al.</i> 2007. <i>Eur. J. Immunol.</i> 37:1809. <u>PubMed</u>	
11. Wang C, <i>et al.</i> 2009. <i>Mucosal Immunol</i> 2:173. (ICFC) <u>PubMed</u> 12. Cui Y, <i>et al.</i> 2009. <i>Invest. Ophth. Vis. Sci.</i> 50:5811. (ICFC) <u>PubMed</u>	
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15. Ma Y, <i>et al.</i> 2012. <i>PLoS One.</i> 7:e40763. <u>PubMed</u> 16. Murakami R, <i>et al.</i> 2013. <i>PLoS One.</i> 8:73270. <u>PubMed</u>	

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.
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Antigen 1. Fitzgerald K, *et al.* Eds. 2001. The Cytokine FactsBook. Academic Press San
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 2. Numasaki M, *et al.* 2002. *Blood* 101:2620.

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