Brilliant Violet 510™ anti-mouse CD25

Catalog # / Size: 1110210 / 500 μl

1110205 / 125 µl

Clone: PC61

Isotype: Rat IgG1, λ

Immunogen: IL-2-dependent cytolytic mouse T-cell

clone B6.1

Reactivity: Mouse

Preparation: The antibody was purified by affinity

chromatography and conjugated with Brilliant Violet $510^{\,\text{\tiny IM}}$ under optimal conditions. The solution is free of unconjugated Brilliant Violet $510^{\,\text{\tiny IM}}$ and

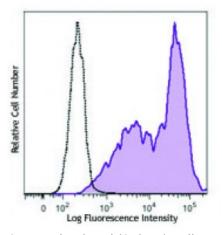
unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and BSA

(origin USA).

Concentration: Lot-specific



Con A-stimulated (3 days) Balb/c splenocytes were stained with CD25 (clone PC61) Brilliant Violet 510™ (filled histogram). Unstained control cells are represented by the open histogram.

Applications:

Applications: Flow Cytometry

Recommended

Usage:

Each lot of this antibody is quality control tested by 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.

Brilliant Violet 510^{TM} excites at 405 nm and emits at 510 nm. The bandpass filter 510/50 nm is recommended for detection, although filter optimization may be required depending on other fluorophores used. **Be sure to verify that your cytometer configuration and software setup are appropriate for detecting this channel.** Refer to your instrument manual or manufacturer for support. Brilliant Violet 510^{TM} is a trademark of Sirigen Group Ltd.

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Application Notes:

Additional reported applications (for the relevant formats) include: immunoprecipitation 1,2 , *in vitro* blocking of IL-2 binding to low- and high-affinity receptors $^{1-4}$, growth inhibition of IL-2-dependent T-cell lines $^{1-4}$, *in vivo* depletion of CD25+CD4+ Treg cells $^{5-8,10}$, and immunohistochemical staining of acetone-fixed frozen sections 2. PC61 antibody recognizes a different epitope than 3C7 antibody (Cat. No. 101902). The LEAF $^{\text{TM}}$ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 102014). For *in vivo* studies or highly sensitive assays, we recommend Ultra-LEAF $^{\text{TM}}$ purified antibody (Cat. No. 102040) with a lower endotoxin limit than standard LEAF $^{\text{TM}}$ purified antibodies (Endotoxin <0.01 EU/microg).

Application References:

- 1. Lowenthal JW, et al. 1985. Nature 315:669. (IP, Block)
- 2. Ceredig R, et al. 1985. Nature 314:98. (IP, IHC, Block)
- 3. Lowenthal JW, et al. 1985. J. Immunol. 135:3988. (Block)
- 4. Moreau JL, et al. 1987. Eur. J. Immunol. 17:929. (Block)
- 5. Takahashi T, *et al.* 2000. *J. Exp. Med.* 192:303. (Deplete)
- 6. Onizuka S, et al. 1999. Cancer Res. 59:3128. (Deplete)
- 7. Lei TC, *et al.* 2005. *Blood* 105:4865. (Deplete)
- 8. Pasare C, et al. 2004. Immunity 21:733. (Deplete)
- 9. León-Ponte M, et al. 2007. Blood 109:3139.
- 10. Cao OW, et al. 2007. Blood doi:10.1182/blood-2007-02-073304. (Deplete)
- 11. Benson MJ, et al. 2007. J. Exp. Med. doi:10.1084/jem.20070719.
- 12. Anguela XM, et al. 2013. Diabetes. 62:551. PubMed.

Description:

CD25 is a 55 kD glycoprotein also known as the low affinity IL-2R α , Ly-43, p55, or Tac. It is expressed on activated T and B cells, thymocyte subsets, pre-B cells, and T regulatory cells. In association with CD122 (IL-2R β) and CD132 (common γ chain), CD25 forms the high affinity signaling IL-2 receptor.

Antigen References:

- 1. Taniguchi T, et al. 1993. Cell 73:5.
- 2. Waldmann TA. 1991. J. Biol. Chem. 266:2681.
- 3. Read S, et al. 2000. J. Exp. Med. 192:295.
- 4. Lowenthal JW, et al. 1985. J. Immunol.