

Store at
4°C

CD127/IL-7R α (A7R34) Rat mAb (APC Conjugate)

#28470

Support: +1-978-867-2388 (U.S.)
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UniProt ID #P16872

New 07/19

For Research Use Only. Not For Use In Diagnostic Procedures.**Applications**
F
Endogenous**Species Cross-Reactivity**
M**Isotype**
Rat IgG2a

Description: This Cell Signaling Technology antibody is conjugated to APC and tested in-house for direct flow cytometric analysis in mouse cells.

Background: The IL-7 receptor (IL-7R) is comprised of two protein subunits, CD127/IL-7R α (IL-7R α) and the common gamma chain (CD132), which is the major signaling component for several cytokines including IL-2, IL-4, IL-9, IL-15, and IL-21 (1). CD127/IL-7R α is a transmembrane protein belonging to the cytokine receptor homology class 1 (CRH1) and is expressed by a wide variety of cells including immature B cells, thymic natural killer cells, bone marrow stromal cells, and T cells (5-6). On its own, CD127/IL-7R α functions as a receptor for two cytokine receptor complex signaling cascades: IL-7 and thymic stromal lymphopoietin (TSLP) (2). IL-7 signaling contributes to T cell development and homeostasis whereas TSLP receptor signaling contributes to dendritic cell activation and B cell development. IL-7 signaling is an essential component in regulating the homeostasis of naive and memory T cells as differential expression of CD127/IL-7R α is observed on naive and activated T cells, which occurs following TCR activation. Specifically, CD127/IL-7R α expression is downregulated on activated T cells and the subsequent re-expression of CD127/IL-7R α on these cells is indicative of cells that will differentiate into memory T cells (3-4).

The A7R34 antibody is widely used to identify CD127/IL-7R α expression on both B and T cells (6).

Specificity/Sensitivity: CD127/IL-7R α (A7R34) Rat mAb (APC Conjugate) recognizes endogenous levels of total CD127/IL-7R α protein. This antibody detects an epitope within the extracellular domain.

Source/Purification: This monoclonal antibody was purified from tissue culture supernatant via affinity chromatography. The purified antibody was conjugated under optimal conditions, with unreacted dye removed from the preparation.

Storage: Supplied in 10 mM NaH₂PO₄, 150 mM NaCl, 0.09% Na₂S₂O₃, 0.1% gelatin, pH 7.2. This product is stable for 6 months when stored at 4°C. *Do not aliquot the antibody. Protect from light. Do not freeze.*

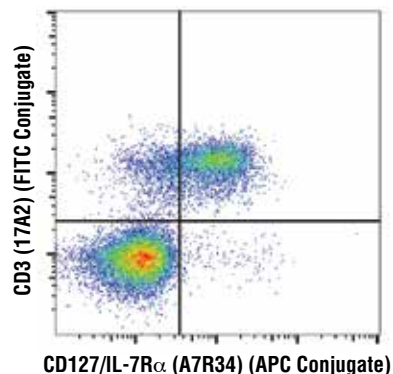
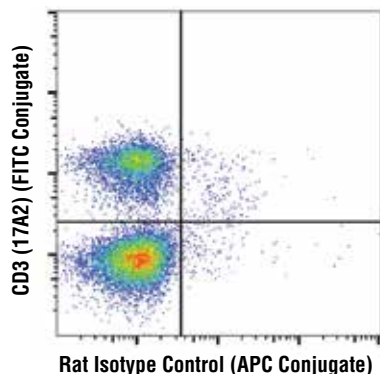
Recommended Antibody Dilutions:

Flow Cytometry 1:80

For product specific protocols and a complete listing of recommended companion products please see the product web page at www.cellsignal.com.

Background References:

- (1) Rochman, Y. et al. (2009) *Nat Rev Immunol* 9, 480-90.
- (2) Levin, S.D. et al. (1999) *J Immunol* 162, 677-83.
- (3) Schluns, K.S. et al. (2000) *Nat Immunol* 1, 426-32.
- (4) Rochman, Y. and Leonard, W.J. (2008) *J Immunol* 181, 7699-705.
- (5) McElroy, C.A. et al. (2012) *Proc Natl Acad Sci U S A* 109, 2503-8.
- (6) Sudo, T. et al. (1993) *Proc Natl Acad Sci U S A* 90, 9125-9.



Flow cytometric analysis of live mouse splenocytes using CD127/IL-7R α (A7R34) Rat mAb (APC Conjugate) and co-stained with CD3 (17A2) Rat mAb (FITC Conjugate) #86603 (right), compared to concentration-matched Rat Isotype Control (APC Conjugate) (left).

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Applications: W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation IF—Immunofluorescence F—Flow cytometry E-P—ELISA-Peptide **Species Cross-Reactivity:** H—human M—mouse R—rat Hm—hamster Mk—monkey Mi—mink C—chicken Dm—D. melanogaster X—Xenopus Z—zebrafish B—bovine Dg—dog Pg—pig Sc—S. cerevisiae Ce—C. elegans Hr—Horse All—all species expected **Species** enclosed in parentheses are predicted to react based on 100% homology.