Purified anti-mouse CD115 (CSF-1R)

Catalog # / Size: 1277510 / 500 µg

1277505 / 50 μg

Clone: AFS98

Isotype: Rat IgG2a, ĸ

Reactivity: Mouse

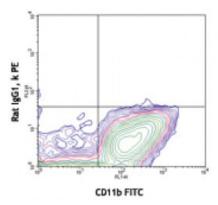
The antibody was purified by affinity **Preparation:**

chromatography.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

Concentration:



Thioglycolate-elicited BALB/c mouse peritoneal macrophages stained with rat IgG1, k PE and CD11b FITC.

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 ≤ 0.25 microg per 10⁶ cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.

Application Notes: Additional reported applications (for the relevant formats) include: blocking of ligand binding1. The LEAF™ purified

antibody (Endotoxin < 0.1 EU/microg, Azide-Free, 0.2 µm filtered) is recommended for functional assays.

CD11b FITC

Thioglycolate-elicited BALB/c mouse peritoneal macrophages stained with AFS98 PE and CD11b FITC.

Application References:

1. Sudo T, et al. 1995. Oncogene 11:2469.

2. Murayama T, et al. 1999. Circulation 99:1740.

osteoclasts.

3. Jaeger BN, et al. 2012. J. Exp. Med. 209:565. PubMed

Description:

CSF-1R, also known as CD115 and M-CSFR, is a single-pass type I membrane protein and member of the platelet-derived growth factor receptor family. This cfms (Fms proto-oncogene) gene product's natural ligands include M-CSF and IL-34. Structural studies of CD115 have described an Ig-like extracellular domain, a transmembrane domain, an intracellular juxtamembrane domain, a split tyrosine kinase domain, and a C-terminal tail receptor. Receptor activation induces homodimerization in addition to phosphorylation and ubiquitination of intracellular residues. CD115 directly influences tissue macrophage and osteoclast differentiation and proliferation. It is expressed on monocytes/macrophages, peritoneal exudate cells, plasmacytoid and conventional dendritic cells, and

Antigen References:

- Sudo T, et al. 1995 Oncogene 11:2469.
 Murayama T, et al. 1999 Circulation 99:1740.
- 3. Goswami S, et al. 2005 Cancer Res. 65:5278.
- 4. Yu W, et al. 2008 J. Leuko. Bio