CD115 antibodies, mouse

For research use only

30 µg equal 100 tests, 150 µg equal 500 tests. One test corresponds to labeling of 10 cells.

Product	Content	Order no.
CD115-Biotin	30 µg in 200 µL	130-112-826
CD115-FITC	30 µg in 200 µL	130-112-827
CD115-FITC	150 µg in 1 mL	130-112-638
CD115-PE	30 µg in 200 µL	130-112-828
CD115-PE	150 µg in 1 mL	130-112-639
CD115-APC	30 µg in 200 µL	130-112-829
CD115-APC	150 µg in 1 mL	130-112-640
CD115-PE-Vio615	30 µg in 200 µL	130-112-833
CD115-PE-Vio615	150 µg in 1 mL	130-112-644
CD115-PE-Vio770	30 µg in 200 µL	130-112-830
CD115-PE-Vio770	150 µg in 1 mL	130-112-641
CD115-Biotin	150 µg in 1 mL	130-112-637

Warnings

Reagents contain sodium azide. Under acidic conditions sodium azide yields hydrazoic acid, which is extremely toxic. Azide compounds should be diluted with running water before discarding. These precautions are recommended to avoid deposits in plumbing where explosive conditions may develop.

Technical data and background information

Antigen	CD115
Clone	REA827
Isotype	recombinant human IgG1
Isotype control	REA Control antibodies
Alternative names of antigen	Csfmr, Fim-2, FMS, M-CSFR, CSF-1R, c-fms
Entrez Gene ID	12978
Molecular mass of antigen [kDa]	107
Distribution of antigen	breast, dendritic cells, macrophages, myeloid leukemia cells, osteoclasts, smooth muscle
Product format	Reagents are supplied in buffer containing stabilizer and 0.05% sodium azide.
Fixation	Cells should be stained prior to fixation, if formaldehyde is used as a fixative.
Storage	Store protected from light at 2–8 °C. Do not freeze.

Clone REA827 recognizes the protein tyrosine kinase transmembrane receptor for macrophage colony-stimulating factor (M-CSF), also known as CSF-1 and interleukin 34 (IL-34). CD115 is a single-pass type I membrane protein. Receptor activation induces homo-dimerization, which leads to phosphorylation and ubiquitinylation of intracellular residues. CD115 is expressed on

mouse monocytes, macrophages, and osteoclasts, as well as on common dendritic cell precursors and macrophage/dendritic cell precursors. Additional information: Clone REA827 displays negligible binding to Fc receptors.

Reagent requirements

- Buffer: Prepare a solution containing phosphate-buffered saline (PBS), pH 7.2, 0.5% bovine serum albumin (BSA), and 2 mM EDTA by diluting MACS® BSA Stock Solution (# 130-091-376) 1:20 with autoMACS® Rinsing Solution (# 130-091-222). Keep buffer cold (2−8 °C).
  Note: EDTA can be replaced by other supplements such as anticoagulant citrate dextrose formula-A (ACD-A) or citrate phosphate dextrose (CPD). Buffers or media containing Ca^{2+} or Mg^{2+} are not recommended for use.
- (Optional) Fluorochrome-conjugated anti-biotin antibodies, e.g., Anti-Biotin-PE (# 130-090-756) as secondary antibody reagent in combination with biotinylated antibodies.
- (Optional) Propidium Iodide Solution (# 130-093-233) for flow cytometric exclusion of dead cells without fixation.
- (Optional) Fixation and Dead Cell Discrimination Kit (# 130-091-163) for cell fixation and flow cytometric exclusion of dead cells.

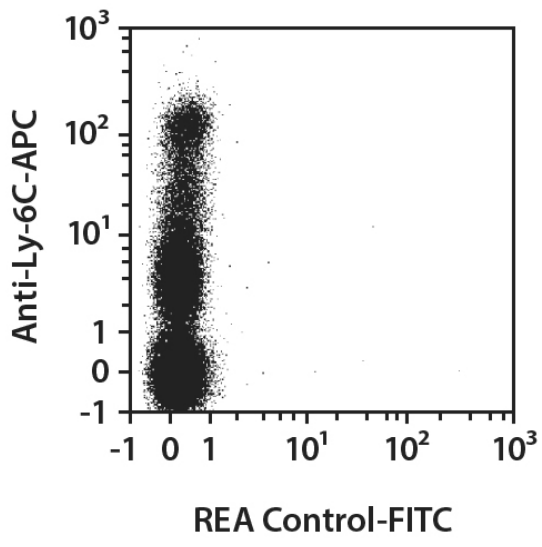
Protocol for cell surface staining

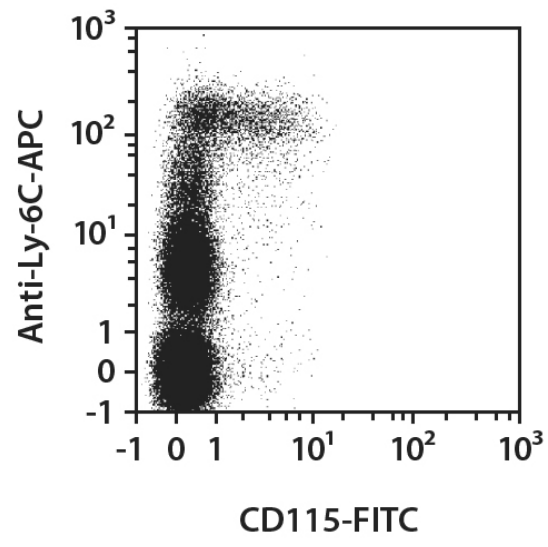
- The recommended antibody dilution for labeling of cells and subsequent analysis by flow cytometry is 1:50 for up to 10^6 cells/100 µL.
- Volumes given below are for up to 10^6 nucleated cells. When working with fewer than 10^6 cells, use the same volumes as indicated. When working with higher cell numbers, scale up all reagent volumes and total volumes accordingly.

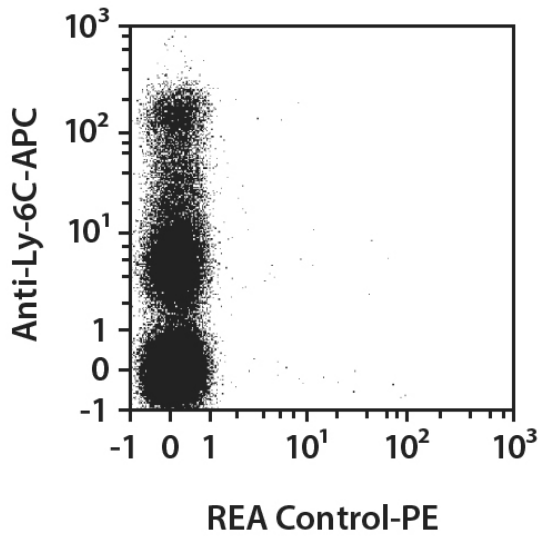
1. Determine cell number.
2. Centrifuge cell suspension at 300×g for 10 minutes. Aspirate supernatant completely.
3. Resuspend up to 10^6 nucleated cells per 98 µL of buffer.
4. Add 2 µL of the antibody.
5. Mix well and incubate for 10 minutes in the dark in the refrigerator (2−8 °C).
   Note: Higher temperatures and/or longer incubation times may lead to non-specific cell labeling. Working on ice requires increased incubation times.
6. Wash cells by adding 1−2 mL of buffer and centrifuge at 300×g for 10 minutes. Aspirate supernatant completely.
7. (Optional) If biotinylated antibody was used, resuspend the cell pellet in buffer and stain with fluorochrome-conjugated anti-biotin antibody according to the manufacturer's recommendations.
8. Resuspend cell pellet in a suitable amount of buffer for analysis by flow cytometry or fluorescence microscopy.

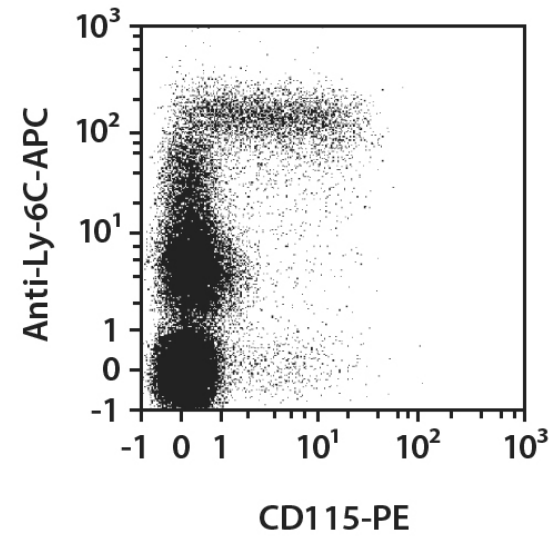
Examples of immunofluorescent staining

Splenocytes of C57BL/6 mice were stained with CD115 antibodies or the corresponding REA Control antibodies (left image) as well as with Anti-Ly-6C antibodies. Flow cytometry was performed using the MACSQuant®Analyzer. Cell debris and dead cells were excluded from the analysis based on scatter signals and propidium iodide fluorescence or 4',6-diamidino-2-phenylindole (DAPI) fluorescence, as in the case of tandem conjugates.






Anti-Ly-6C-APC
REA Control-PE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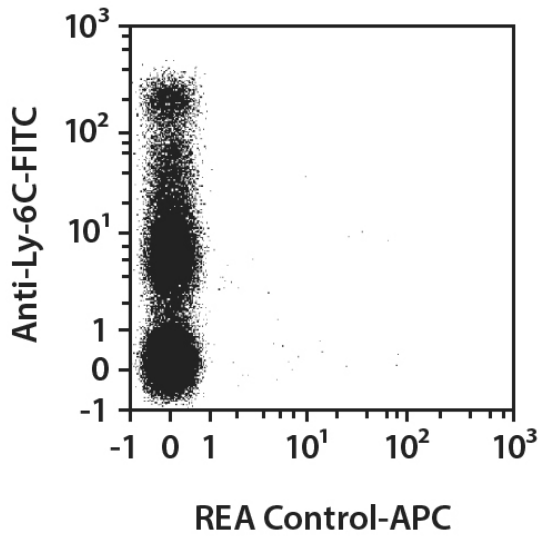
CD115-PE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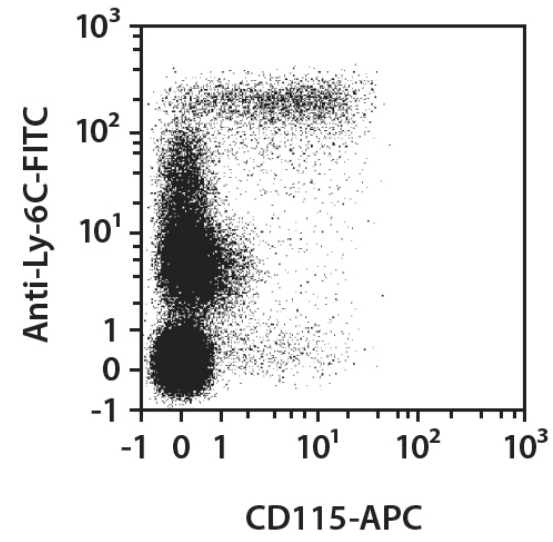
REA Control-APC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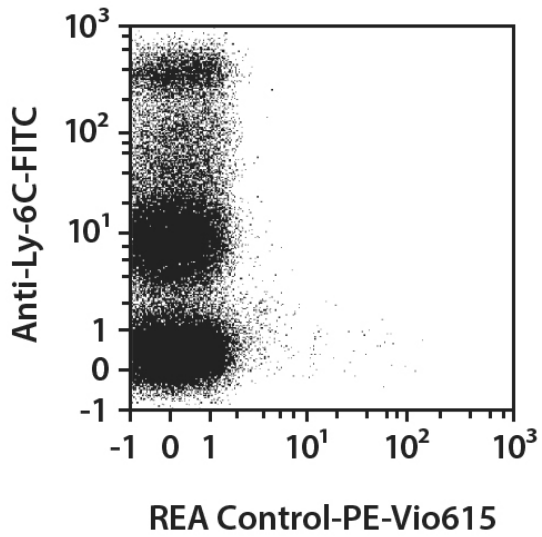
CD115-APC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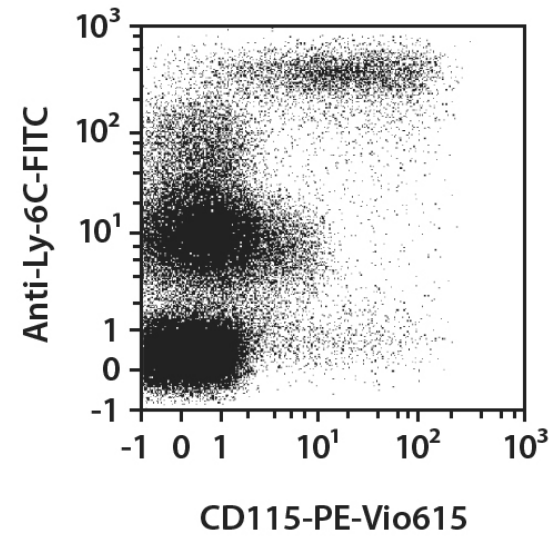
REA Control-PE-Vio615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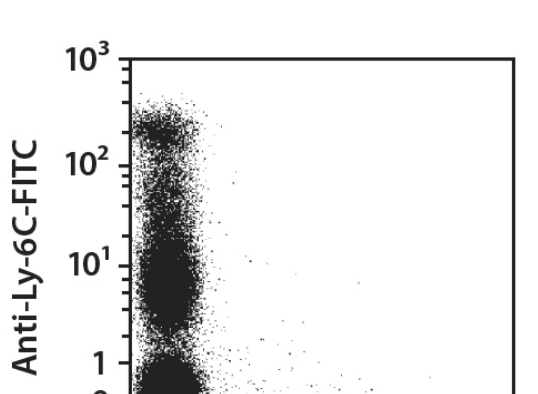
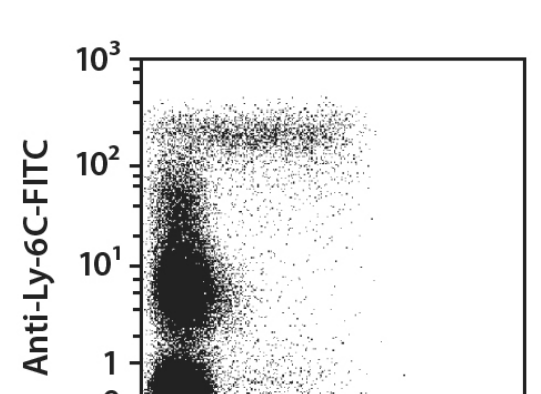
CD115-PE-Vio615


Anti-Ly-6C-FITC
REA Control-PE-Vio770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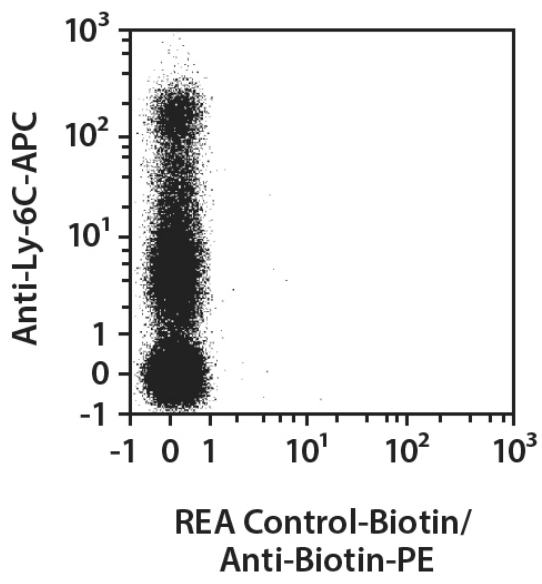
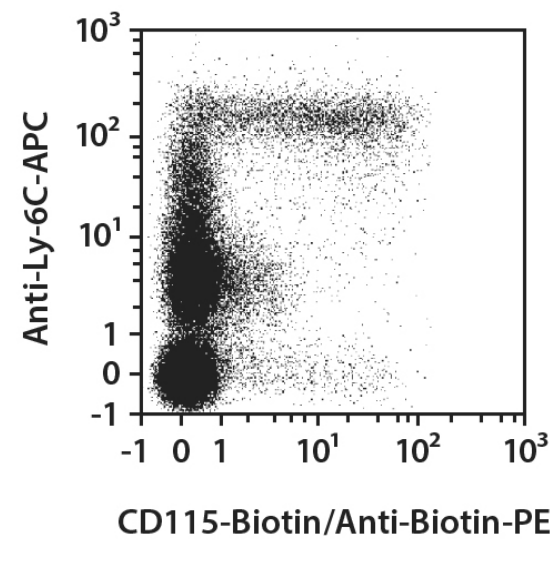
CD115-PE-Vio770


Anti-Ly-6C-APC
REA Control-Biotin/
Anti-Biotin-PE


Anti-Ly-6C-APC
CD115-Biotin/Anti-Biotin-PE

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