Product Data Sheet

PE anti-mouse CD71

Catalog # / Size: $1169035 / 50 \mu g$

1169040 / 200 µg

Clone: RI7217

Isotype: Rat IgG2a, κ

Reactivity: Mouse

Preparation: The antibody was purified by affinity

chromatography, and conjugated with PE under optimal conditions. The solution is free of unconjugated PE and

unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

Concentration: 0.2



C57BL/6 mouse bone marrow cells stained with RI7217 FITC and

Ter119 PE

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 10^6 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 cellular proliferation. The LEAF $^{\text{TM}}$ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 113810).

This clone may also be known as R17217 or R17 217.

Application References:

- 1. Trowbridge I, et al. 1982. J. Cell. Physiol. 112:403.
- 2. Grisendi S, et al. 2005. Nature 437:147.
- 3. van Rooy I, et al. 2010. J. Control Release 150:30. PubMed
- 4. Willhelm BT, et al. 2011. Blood 117:27. PubMed
- 5. Okasi Y, et al. 2012. Exp Hematol. 40:143PubMed
- 6. Stojanov K, et al. 2012. Mol Pharm. 9:1620. PubMed

Description: CD71 is a 95 kD type II heterodimeric transmembrane glycoprotein that is also

known as T9 and transferrin receptor. CD71 is expressed on proliferating cells, reticulocytes, and erythroid precursors. Its expression is very low on resting leukocytes. CD71 plays a role in the control of cellular proliferation by facilitating the uptake of iron via ferrotransferrin binding and the recycling of apotransferrin

to the cell surface.

Antigen References:

- 1. Hentze MW, et al. 1996. P. Natl. Acad. Sci. USA 93:8175.
- 2. Trowbridge IS, et al. 1993. Annu. Rev. Cell Biol. 9:129.
- 3. Trowbridge I, *et al.* 1982. *J. Cell Physiol.* 112:403.

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