

RAT anti-MOUSE CD16/32

Publication No. MAN0006739

Store at 2° to 8°C

Rev. 2.00

Catalog No. Form		Amount	Excitation	Peak Emission	
A14717	FITC	0.05 mL (25 μg)	488 nm	519 nm	
A14718	APC	0.25 mL (25 μg)	650 nm	660 nm	
A14789	PerCP-Cy [®] 5.5	0.125 mL (25 μg)	482 nm	695 nm	
A14719	PE-Cy®7	0.25 mL (25 μg)	488 nm	767 nm	

Product Description

The Rat anti-Mouse CD16/32 Monoclonal Antibody (mAb) recognizes a conformational epitope formed by lymphocyte FcyII and FcyIII receptors. The Fcy receptors (FcyR) recognize the Fc portion of IgG, presented as immune complexes or free antibody. The Fc type II receptor is expressed on a wide variety of cells including B cells, hematopoietic cells, monocyte/macrophages, neutrophils, platelets, Langerhans cells, eosinophils, basophils, trophoblasts, and endothelial cells of the placenta. The Fcy type III receptors are expressed on macrophages, NK cells, and neutrophils. Both types of receptors can be constitutively expressed on the same cell in varying ratios, and can be modulated cytokines and lymphokines. The Rat anti-Mouse CD16/32 mAb can identify FcyR+ cells, and block Fc receptor binding.

Product Specifications

Clonality: Monoclonal Host/Class: Rat IgG

Reactivity: Mouse CD16/32

Gene ID: 14131
Sequence Identity: Mouse
Clone/PAD: 93
Isotype: IgG_{2a}

Lot: See product label

Product Applications

Applications reported for the Rat anti-Mouse CD16/32 mAb include flow cytometry, and blocking Fcy receptors.

Because conditions may vary, it is recommended that each investigator determine the optimal amount of antibody to be used for each application.

Storage and Handling

Store reagents at 2° to 8°C. If the reagent is being diluted, it is recommended that only the quantity to be used within one week be diluted. Cells should be analyzed within 18 hours of staining for best results.

Avoid light exposure with fluorochrome-conjugated antibodies. Use dim light during handling, incubation with cells, and prior to analysis.

Stability

When stored as instructed, expires one year from date of receipt unless otherwise indicated on product label.

Storage Buffer

Phosphate buffered saline (PBS) with 0.1% sodium azide, and a non-BSA stabilizing agent. FITC conjugates contain PBS with 0.1% sodium azide. PerCP-Cy $^{\odot}$ 5.5 conjugates contain an aqueous buffer with 0.09% sodium azide, and may contain carrier protein/stabilizer.

Caution: Sodium azide is an extremely toxic and dangerous compound particularly when combined with acids or metals. Properly dispose of solutions containing sodium azide.

Product Documentation

To obtain a Certificate of Analysis or Safety Data Sheets (SDSs), visit www.lifetechnologies.com/support.

Related Products

Product Name	Quantity	Catalog no.
AbC [™] Anti-Mouse Bead Kit	1 kit	A10344
AbC [™] anti-Rat/Hamster Bead Kit	1 kit	A10389
Protein A Agarose	5 mL	15918-014
Recombinant Protein G (rProtein G) Agarose	5 mL	15920-010

Explanation of symbols

Symbol	Description	Symbol	Description
REF	Catalogue Number	LOT	Batch code
RUO	Research Use Only	IVD	In vitro diagnostic medical device
X	Use by	1	Temperature limitation
***	Manufacturer	EC REP	European Community authorised representative
[-]	Without, does not contain	[+]	With, contains
from Light	Protect from light	À	Consult accompanying documents
$\prod i$	Directs the user to consult instructions for use (IFU), accompanying the product.		

Limited Product Warranty

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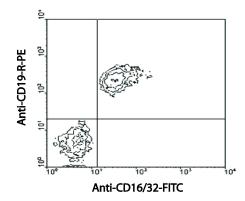


Figure 1 Two-color analysis of CD16/32 expression on mouse splenocytes.

BALB/c spleen cells were double-stained with Rat anti-Mouse CD16/32-FITC Monoclonal Antibody (Cat. no. A14717) and a rat anti-mouse CD19-R-PE antibody. Lymphocytes were gated and analyzed on a FACScan[™] flow cytometer (BDIS, San Jose, CA).

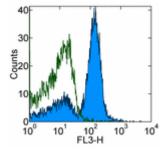


Figure 2 CD16/32 expression on mouse splenocytes.

Staining of BALB/c splenocytes with 0.06 μg of a rat IgG2a K-PerCP-Cy®5.5 isotype control (open histogram) or 0.06 μg of Rat anti-Mouse CD16/32-PerCP-Cy®5.5 Monoclonal Antibody (Cat. no. A14789) (filled histogram). Total viable cells were used for analysis.

Note: All flow cytometric data shown may not necessarily have been generated using the enclosed lot of reagent. For this reason, and due to differences in flow cytometers and cytometer settings, results may vary from those illustrated above. It is suggested that investigators titrate reagents to determine optimal conditions for use in their systems.

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