

# Mouse Fc gamma RIII (CD16) APC-conjugated Antibody Monoclonal Rat IgG<sub>2A</sub> Clone # 275003

Catalog Number: FAB19601A

100 TESTS, 25 TESTS

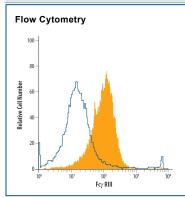
DESCRIPTION			
Species Reactivity	Mouse		
Specificity	Detects mouse Fcγ RIIIA/B (CD16) in direct ELISAs. In direct ELISAs, no cross-reactivity with recombinant mouse Fcrl3/CD16-2, rmFcγ RI or rhFcγ RIIA is observed.		
Source	Monoclonal Rat IgG <sub>2A</sub> Clone # 275003		
Purification	Protein A or G purified from hybridoma culture supernatant		
Immunogen	Mouse myeloma cell line NS0-derived recombinant mouse Fcγ RIII Ala31-Thr215 Accession # Q5D5J5		
Conjugate	Allophycocyanin Excitation Wavelength: 620-650 nm Emission Wavelength: 660-670 nm		
Formulation	Supplied in a saline solution containing BSA and Sodium Azide. See Certificate of Analysis for details.		
	*Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Shee (SDS) for additional information and handling instructions.		

#### **APPLICATIONS**

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Flow Cytometry	10 μL/10 <sup>6</sup> cells	See Below

# DATA



Detection of Fc gamma RIII (CD16) in RAW 264.7 Mouse Cell Line by Flow Cytometry. RAW 264.7 mouse monocyte/ macrophage cell line was stained with Rat Anti-Mouse Fc gamma RIII (CD16) APCconjugated Monoclonal Antibody (Catalog # FAB19601A, filled histogram) or isotype control antibody (Catalog # IC006A, open histogram). View our protocol for Staining Membrane-associated Proteins

## PREPARATION AND STORAGE

Shipping The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.

Stability & Storage

Protect from light. Do not freeze.

• 12 months from date of receipt, 2 to 8 °C as supplied.





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### **BACKGROUND**

Receptors for the Fc region of IgG (Fc  $\gamma$  Rs) are members of the Ig superfamily that function in the activation or inhibition of immune responses such as degranulation, phagocytosis, ADCC (antibody-dependent cellular toxicity), cytokine release, and B cell proliferation (1-3). The Fc  $\gamma$  Rs have been divided into three classes based on close relationships in their extracellular domains; these groups are designated Fc  $\gamma$  RI (also known as CD64), Fc  $\gamma$  RII (CD32), and Fc  $\gamma$  RIII (CD16). Each group may be encoded by multiple genes and exist in different isoforms depending on species and cell type. The CD64 proteins are high affinity receptors (~10-8-10-9 M) capable of binding monomeric IgG, whereas the CD16 and CD32 proteins bind IgG with lower affinities (~10-8-10-7 M) only recognizing IgG aggregates surrounding multivalent antigens (1, 4). Fc  $\gamma$  Rs that deliver an activating signal either have an intrinsic immunoreceptor tyrosine-based activation motif (ITAM) within their cytoplasmic domains or associate with one of the ITAM-bearing adapter subunits, Fc R $\gamma$  or  $\zeta$  (3, 5). The only inhibitory member in human and mouse, Fc  $\gamma$  RIIb, has an intrinsic cytoplasmic immunoreceptor tyrosine-based inhibitory motif (ITIM). The coordinated functioning of activating and inhibitory receptors is necessary for successful initiation, amplification, and termination of immune responses (5). Mouse CD16 is encoded by a single gene. The protein product is a type I transmembrane protein having two extracellular Ig-like domains. It is expressed on a variety of myeloid and lymphoid cells (4) and associates with Fc R $\gamma$  to deliver an activating signal upon ligand binding (5). Mouse CD32 is closely related to mouse CD16 throughout its extracellular domain (95% amino acid sequence identity), but has a divergent cytoplasmic domain and functions as an inhibitory receptor. Together these proteins constitute an activating/inhibiting receptor pair to regulate immune responses (5).

#### References:

- 1. van de Winkel, J. and P. Capes (1993) Immunol. Today 14:215.
- 2. Raghaven, M. and P. Bjorkman (1996) Annu. Rev. Cell Dev. Biol. 12:181.
- 3. Ravetch, J. and S. Bolland (2001) Annu. Rev. Immunol. 19:275.
- 4. Takai, T. (2002) Nature Rev. Immunol. 2:580.
- 5. Ravetch, J. and L. Lanier (2000) Science 290:84.

