

# Mouse Fcγ RI/CD64 PE-conjugated Antibody

Monoclonal Rat IgG<sub>2A</sub> Clone # 290322

Catalog Number: FAB20741P 100 TESTS

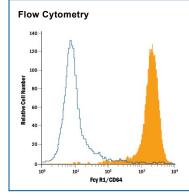
DESCRIPTION		
Species Reactivity	Mouse	
Specificity	Detects mouse Fcy RI/CD64 in direct ELISAs and Western blots. In Western blots, no cross-reactivity with recombinant human (rh) Fcy RIA, rhFcy RIIB, or rhFcy RIIIB is observed.	
Source	Monoclonal Rat IgG <sub>2A</sub> Clone # 290322	
Purification	Protein A or G purified from hybridoma culture supernatant	
Immunogen	NS0-derived recombinant mouse Fcγ RI/CD64 extracellular domain Glu25-Pro297 Accession # P26151	
Conjugate	Phycoerythrin Excitation Wavelength: 488 nm Emission Wavelength: 565-605 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 Sheet (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 Fcy RI/CD64 in RAW 264.7 Mouse Cell Line by Flow Cytometry. RAW 264.7 mouse monocyte/macrophage cell line was stained with Rat Anti-Mouse Fcy RI/CD64 PE-conjugated Monoclonal Antibody (Catalog # FAB20741P, filled histogram) or isotype control antibody (Catalog # IC006P, 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.

## BACKGROUND

Receptors for the Fc region of IgG (Fcγ 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 Fcγ Rs have been divided into three classes based on close relationships in their extracellular domains; these groups are designated Fcγ RI (also known as CD64), Fcγ RII (CD32), and Fcγ 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 ( $\sim 10^{-8} \cdot 10^{-9}$  M) capable of binding monomeric IgG, whereas the CD16 and CD32 proteins bind IgG with lower affinities ( $\sim 10^{-6} \cdot 10^{-7}$  M) only recognizing IgG aggregates surrounding multivalent antigens (1, 4). Fcγ 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 Rγ or  $\sim 10^{-2} \cdot 10^{-2}$  M; The only inhibitory member in human and mouse, Fcγ 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 Fcγ RI is transmembrane protein with three extracellular Ig-like domains, and it delivers an activating signal via the associated Fc Rγ accessory chain (1, 2). The high affinity recognition of IgG by Fcγ RI permits the triggering of effector responses at low IgG concentrations typical of early immune responses (2). Fcγ RI is expressed constitutively on monocytes and macrophages and can be induced on neutrophils and eosinophils (1, 4). Its expression is up-regulated during bacterial infections and sepsis.

## 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

Rev. 5/21/2014 Page 1 of 1

