

## Human Fcy RI/CD64 Biotinylated Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: BAF1257

DESCRIPTION			
Species Reactivity	Human		
Specificity	Detects human Fcy RI/CD64 in Western blots. In Western blots, approximately 50% cross-reactivity with recombinant mouse Fcy RI is observed and less than 1% cross-reactivity with recombinant human (rh) Fcy RIIA and rhFcy RIIB is observed.		
Source	Polyclonal Goat IgG		
Purification	Antigen Affinity-purified		
Immunogen	Mouse myeloma cell line NS0-derived recombinant human Fcγ RI/CD64 Gln16-Pro288 Accession # P12314.2		
Formulation	Lyophilized from a 0.2 µm filtered solution	in PBS with BSA as a carrier protein. See Certificate of Analysis for details.	
Please Note: Optimal diluti	ons should be determined by each laboratory for each app Recommended Concentration	olication. General Protocols are available in the Technical Information section on our website.  Sample	
Western Blot	0.1 μg/mL	Recombinant Human Fcy RI/CD64 (Catalog # 1257-FC)	
PREPARATION AND			
Reconstitution	Reconstitute at 0.2 mg/mL in sterile PBS.		
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.		
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles.		
	• 12 months from date of receipt, -20 to -70 °C as supplied.		
	1 month, 2 to 8 °C under sterile cond		
	<ul> <li>6 months, -20 to -70 °C under sterile conditions after reconstitution.</li> </ul>		

## 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 (~10<sup>-8</sup>-10<sup>-9</sup> M) capable of binding monomeric IgG, whereas the CD16 and CD32 proteins bind IgG with lower affinities (~10<sup>-6</sup>-10<sup>-7</sup> 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 ζ (3, 5). 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).

Three highly homologous genes (A, B, and C) sharing 98% identity at the nucleotide level have been identified for the human CD64 group (1). Fcy RI is transmembrane protein with three extracellular Ig-like domains, and it delivers an activating signal via the associated Fc Ry accessory chain. The genes for Fcy RIB and Fcy RIC contain stop codons within their membrane proximal Ig-like domains indicating possible secreted receptors (1, 6). An mRNA splice variant of Fcy RIB has a deletion of the membrane-proximal Ig-like domain and encodes a putative transmembrane receptor (6). The high affinity recognition of IgG by Fcy RI permits the triggering of effector responses at low IgG concentrations typical of early immune responses (2). Fcy 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
- 6. Ernst, L. et al. (1998) Mol Immunol. 35:943.

