

# Human Fcγ RIIIA/CD16a Antibody

Monoclonal Mouse IgG<sub>2A</sub> Clone # 1041618 Catalog Number: MAB10751

DESCRIPTION		
Species Reactivity	Human	
Specificity	Detects human Fcγ RIIIA/CD16a in ELISA.	
Source	Monoclonal Mouse IgG <sub>2A</sub> Clone # 1041618	
Purification	Protein A or G purified from hybridoma culture supernatant	
Immunogen	Mouse myeloma cell line, NS0-derived human Fcγ RIIIA/CD16a Gly17-Gln208 Accession # P08637	
Formulation	Lyophilized from a 0.2 μm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 μm filtered solution in PBS.	

ΔF	PI	IC/	١TI	OI	NS

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
Immunocytochemistry	8-25 μg/mL	Immersion fixed human peripheral blood mononuclear cells (PBMCs)
Neutralization	In a functional ELISA binding assay, 0.5-5 μg/mL of this antibody will block 50% of the binding of 2 μg/mL of recombinant human fcgRIIIA (Catalog # 8894-FC) to Normal Human IgG, biotin at 2 ug/mL (100 μL/well). At 10 μg/mL, this antibody will block >90% of the binding.	

#### DATA

## Immunocytochemistry







Negative (A549 cells)

Fcv RIIIA/CD16a in Human PBMCs. Fcy RIIIA/CD16a was detected in immersion fixed human peripheral blood mononuclear cells (PBMCs) (positive staining) and A549 human lung carcinoma cell line (negative staining) using Mouse Anti-Human Fcγ RIIIA/CD16a Monoclonal Antibody (Catalog # MAB10751) at 8 µg/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557conjugated Anti-Rabbit IaG Secondary Antibody (red; Catalog # NL004) and counterstained with DAPI (blue). Specific staining was localized to plasma membrane. Staining was performed using our protocol for Fluorescent ICC Staining of Non-adherent Cells.

### PREPARATION AND STORAGE

**Reconstitution** Reconstitute at 0.5 mg/mL in sterile PBS.

Shipping

The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.

\*Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C

### 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 conditions after reconstitution.
- 6 months, -20 to -70 °C under sterile conditions after reconstitution.





# Human Fcy RIIIA/CD16a Antibody

Monoclonal Mouse IgG<sub>2A</sub> Clone # 1041618 Catalog Number: MAB10751

#### BACKGROUND

Fcγ RIlla is a low/intermediate affinity receptor for polyvalent immune-complexed IgG. It is involved in phagocytosis, secretion of enzymes and inflammatory mediators, antibody-dependent cytotoxicity and clearance of immune complexes (1, 2). In humans, it is a 50-70 kDa type I transmembrane activating receptor expressed by NK cells, T cells, monocytes, and macrophages (1). Fcγ RIllb is highly related, sharing 97% amino acid (aa) identity within the extracellular domain (ECD), but is a GPI-linked receptor expressed on human neutrophils and eosinophils (1, 2). The ECD of Fcγ RIlla shares 63%, 61%, 65%, 59% and 58% aa identity with mouse Fcγ RIV, rat Fcγ RIlla, feline CD16, bovine CD16 and porcine Fcγ RIllb paralogs, respectively. The Fcγ RIlla cDNA encodes 254 aa including a 16 aa signal sequence, 191 aa ECD with two C2-type Ig-like domains and five potential N-glycosylation sites, a 22 aa transmembrane (TM) sequence and a 25 aa cytoplasmic domain. In humans, a single nucleotide polymorphism creates high binding (176V) and low binding (176F) forms that, when homozygous, may influence susceptibility to autoimmune diseases or response to therapeutic IgG antibodies (3, 4). Catalog # 4325-FC is expressed as the 176V isoform of Fcγ RIlla. Fcγ RIlla surface expression requires interaction of an accessory chain, either the common γ-chain or CD3ζ (5, 6). Glycosylation patterns, electrophoretic mobility and binding affinity appear to differ between NK cell and monocyte Fcγ RIlla (7). The ECD of both Fcγ RIlla release (11). Soluble Fcγ RIll can be detected in normal plasma and is increased in rheumatoid arthritis and in coronary artery diseases (9, 10).

#### References:

- 1. Nimmerjahn, F. and J.V. Ravetch (2006) Immunity 24:19.
- 2. Ravetch, J.V. and B. Perussia (1989) J. Exp. Med. 170:481.
- 3. Wu, J. et al. (1997) J. Clin. Invest. 100:1059.
- 4. Dall'Ozzo, S. et al. (2004) Cancer Res. 64:4664.
- 5. Kim, M.-K. et al. (2003) Blood 101:4479.
- 6. Lanier, L.L. et al. (1989) Nature 342:803.
- 7. Edberg, J.C. and R.P. Kimberley (1997) J. Immunol. 159:3849.
- 8. Li, P. et al. (2007) J. Biol. Chem. 282:6210.
- 9. Masuda, M. et al. (2003) J. Rheumatol. 30:1911.
- 10. Masuda, M. et al. (2006) Atherosclerosis 188:377
- 11. Webster, N.L. et al. (2006) J. Leukoc. Biol. 79:294.