

CD16/CD32 (2.4G2) Rat mAb (IF Formulated)

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Applications:	Reactivity:	Sensitivity:	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
IF-F, IF-IC	M	Endogenous	Rat IgG2b	#P08101-2, #P08508	14130, 14131

Product Usage Information**Application**

Immunofluorescence (Frozen)
Immunofluorescence (Immunocytochemistry)

Dilution

1:50 - 1:100
1:50 - 1:100

Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. *Do not aliquot the antibody.* This product is stable for 60 months when stored at -20°C.

Specificity/Sensitivity

CD16/CD32 (2.4G2) Rat mAb (IF Formulated) recognizes endogenous levels of total CD16/CD32 protein. This antibody detects an epitope within the extracellular domain and is expected to detect all isoforms of CD16 and CD32. Staining of both microglia and astrocytes was seen when staining was done in 5XFAD mouse brain.

Source / Purification

This monoclonal antibody was purified from tissue culture supernatant via affinity chromatography.

Background

CD64 (FcγRI), CD32 (FcγRII), and CD16 (FcγRIII) are three classes of the immunoglobulin superfamily. CD64 has a high affinity for IgG with three Ig-like domains while CD32 and CD16 have low affinities with two Ig-like domains. Two genes encode CD16-A and CD16-B resulting only in a 6 amino acid difference in their ectodomains. However, CD16-A has a transmembrane anchor versus CD16-B, which has a glycosylphosphatidylinositol (1). CD64, CD32, and CD16 are membrane glycoproteins that are expressed by all immunologically active cells and trigger various immune functions (activate B cells, phagocytosis, antibody-dependent cellular cytotoxicity, immune complex clearance, and enhancement of antigen presentation) (2). CD16 cross-linking induces tyrosine phosphorylation (Tyr394) of Lck in NK cells (3). CD32 has tyrosine-based activation motifs in the cytoplasmic domain in contrast to CD16, which associates with molecules possessing these motifs (1).

CD16 and CD32 are expressed on microglia within the brain (4).

Background References

1. Maenaka, K. et al. (2001) *J. Biol. Chem.* 276, 44898-44904.
2. Fridman, W. H. et al. (1992) *Immunol. Rev.* 125, 49-76.
3. Pignata, C. et al. (1993) *J. Immunol.* 151, 6794-6800.
4. Zhang, Y. et al. (2014) *J Neurosci* 34, 11929-47.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Applications Key

IF-F: Immunofluorescence (Frozen) **IF-IC:** Immunofluorescence (Immunocytochemistry)

Cross-Reactivity Key

M: Mouse

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