

## Purified anti-human CD64 Antibody

<b>Catalog# / Size</b>	305002 / 100 µg
<b>Clone</b>	10.1
<b>Regulatory Status</b>	RUO
<b>Workshop</b>	VI MA36
<b>Other Names</b>	FcγRI, FcR I
<b>Isotype</b>	Mouse IgG1, κ
<b>Description</b>	CD64 is a 72 kD single chain type I glycoprotein also known as FcγRI and FcR I. CD64 is a member of the immunoglobulin superfamily and is expressed on monocytes/macrophages, dendritic cells, and activated granulocytes. The expression can be upregulated by IFN-γ stimulation. CD64 binds IgG immune complex. It plays a role in antigen capture, phagocytosis of IgG/antigen complexes, and antibody-dependent cellular cytotoxicity (ADCC).

### Product Details

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<b>Verified Reactivity</b>	Human, Cynomolgus, Rhesus
<b>Reported Reactivity</b>	Baboon, Capuchin Monkey, Chimpanzee, Squirrel Monkey
<b>Antibody Type</b>	Monoclonal
<b>Host Species</b>	Mouse
<b>Immunogen</b>	Human rheumatoid synovial fluid cells and fibronectin-purified monocytes.
<b>Formulation</b>	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
<b>Preparation</b>	The antibody was purified by affinity chromatography.
<b>Concentration</b>	0.5 mg/ml
<b>Storage &amp; Handling</b>	The antibody solution should be stored undiluted between 2°C and 8°C.
<b>Application</b>	<a href="#">FC - Quality tested</a> <a href="#">IHC-F, Block - Reported in the literature, not verified in house</a>
<b>Recommended Usage</b>	Each lot of this antibody is quality control tested by <a href="#">immunofluorescent staining with flow cytometric analysis</a> . For flow cytometric staining, the suggested use of this reagent is ≤ 1.0 µg per 10 <sup>6</sup> cells in 100 µl volume or 100 µl of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.
<b>Application Notes</b>	Clone 10.1 recognizes the EC3 epitope of CD64. While both contain the EC3 domain, in-house testing suggests that clone 10.1 preferentially binds to CD64A (FcγRIA), but not CD64B (FcγRIB). Additional reported applications (for the relevant formats) include: blocking of human IgG3 and murine IgG2a binding to FcγRI <sup>2,5,6,11</sup> and immunohistochemical staining of acetone-fixed frozen tissue sections <sup>12</sup> .
<b>Application References</b>	<ol style="list-style-type: none"> <li>1. McMichael A, <i>et al.</i> Eds. 1987. Leucocyte Typing III. Oxford University Press. New York.</li> <li>2. Schlossman S, <i>et al.</i> Eds. 1995. Leucocyte Typing V. Oxford University Press. New York. p. 874.</li> <li>3. Kishimoto T, <i>et al.</i> Eds. 1997. Leucocyte Typing VI. Garland Publishing Inc. London.</li> <li>4. Holl V, <i>et al.</i> 2004. <i>J. Immunol.</i> 173:6274.</li> <li>5. Hober D, <i>et al.</i> 2002. <i>J. Gen. Virol.</i> 83:2169.</li> <li>6. Cho HJ, <i>et al.</i> 2007. <i>Physiol Genomics</i> 149:60.</li> <li>7. van Tits L, <i>et al.</i> 2005. <i>Arterioscler Thromb Vasc Biol.</i> 25:717. <a href="#">PubMed</a></li> <li>8. Bruhns P, <i>et al.</i> 2008. <i>Blood</i> 113:3716. <a href="#">PubMed</a></li> <li>9. Yoshino N, <i>et al.</i> 2000. <i>Exp. Anim. (Tokyo)</i> 49:97. (FC)</li> <li>10. Carter DL, <i>et al.</i> 1999. <i>Cytometry</i> 37:41. (FC)</li> <li>11. Dougherty GJ, <i>et al.</i> 1987. <i>Eur. J. Immunol.</i> 17:1453.</li> <li>12. Blom AB, <i>et al.</i> 2003. <i>Arthritis Rheum.</i> 48(4):1002-14. (IHC)</li> </ol>
<b>(PubMed link indicates BioLegend citation)</b>	

## Product Citations

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## RRID

AB\_314486 (BioLegend Cat. No. 305002)

## Antigen Details

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<b>Structure</b>	Ig superfamily, type I glycoprotein, 72 kD
<b>Distribution</b>	Monocytes, macrophages, dendritic cells, activated granulocytes
<b>Function</b>	Phagocytosis, ADCC
<b>Ligand/Receptor</b>	IgG receptor
<b>Cell Type</b>	Dendritic cells, Granulocytes, Macrophages, Monocytes
<b>Biology Area</b>	Immunology, Innate Immunity
<b>Molecular Family</b>	CD Molecules, Fc Receptors
<b>Antigen References</b>	1. Hulet M, <i>et al.</i> 1994. <i>Adv. Immunol.</i> 57:1. 2. van de Winkel J, <i>et al.</i> 1993. <i>Immunol. Today</i> 14:215.
<b>Gene ID</b>	<a href="#">2209</a>

## Related Protocols

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[Cell Surface Flow Cytometry Staining Protocol](#)

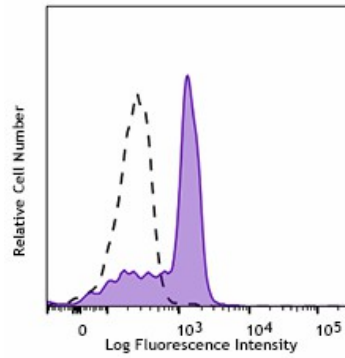
## Other Formats

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Biotin anti-human CD64, FITC anti-human CD64, PE anti-human CD64, Purified anti-human CD64, Alexa Fluor® 488 anti-human CD64, Alexa Fluor® 647 anti-human CD64, APC anti-human CD64, Pacific Blue™ anti-human CD64, Brilliant Violet 421™ anti-human CD64, PE/Cyanine7 anti-human CD64, PerCP/Cyanine5.5 anti-human CD64, APC/Cyanine7 anti-human CD64, Brilliant Violet 510™ anti-human CD64, Purified anti-human CD64 (Maxpar® Ready), PE/Dazzle™ 594 anti-human CD64, Brilliant Violet 605™ anti-human CD64, APC/Fire™ 750 anti-human CD64, TotalSeq™-A0162 anti-human CD64, Brilliant Violet 711™ anti-human CD64, Alexa Fluor® 700 anti-human CD64, Brilliant Violet 785™ anti-human CD64, TotalSeq™-C0162 anti-human CD64, Ultra-LEAF™ Purified anti-human CD64, TotalSeq™-B0162 anti-human CD64, TotalSeq™-D0162 anti-human CD64, GMP PE anti-human CD64, GMP FITC anti-human CD64

## Product Data

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Human peripheral blood monocytes were stained with CD64 (clone 10.1) Purified (filled histogram) or Purified Mouse IgG1,  $\kappa$  isotype control (open histogram) followed by anti-mouse IgG FITC

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