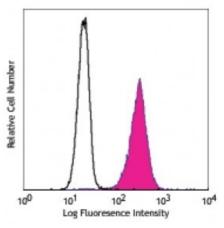
## **Product Data Sheet**

## Pacific Blue<sup>™</sup> anti-human CD11b

Catalog # / Size:	2106575 / 100 μg 2106580 / 25 μg
Clone:	ICRF44
Isotype:	Mouse IgG1, к
<b>Reactivity:</b>	Human
Preparation:	The antibody was purified by affinity chromatography, and conjugated with Pacific Blue <sup>™</sup> under optimal conditions. The solution is free of unconjugated Pacific Blue <sup>™</sup> .
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
Workshop Number:	IV M047
<b>Concentration:</b>	0.5



Human peripheral blood granulocytes stained with ICRF44 Pacific Blue™

## **Applications:**

Applications:	Flow Cytometry
Recommended Usage:	Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. The suggested use of this reagent is $\leq 2.0$ microg per 10 <sup>6</sup> cells in 100 microL volume or 100 microL of whole blood. It is highly recommended that the reagent be titrated for optimal performance for each application.
	* Pacific Blue <sup>™</sup> has a maximum emission of 455 nm when it is excited at 405 nm. Prior to using Pacific Blue <sup>™</sup> conjugate for flow cytometric analysis, please verify your flow cytometer's capability of exciting and detecting the fluorochrome.
Application Notes:	The ICRF44 antibody inhibits heterotypic adhesion of granulocytes in response to fMLP. Additional reported applications (for the relevant formats) include: immunohistochemical staining of acetone-fixed frozen tissue sections, immunofluorescence microscopy5, stimulation of monocytes3, blocking of heterotypic PMN aggregation <sup>8</sup> , and blocking of granulocyte activation <sup>12</sup> . This clone was tested in-house and does not work on formalin fixed paraffin-embedded (FFPE) tissue.
	The LEAF $^{\text{TM}}$ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 301312).
Application References:	<ol> <li>Knapp W. 1989. Leucocyte Typing IV. Oxford University Press New York.</li> <li>Barclay N, <i>et al.</i> 1997. The Leucocyte Antigen Facts Book. Academic Press Inc. San Diego.</li> <li>Rezzonico R, <i>et al.</i> 2001. <i>Blood</i> 97:2932. (Stim)</li> <li>Marsik C, <i>et al.</i> 2003. <i>Shock</i> 20:493. (FC)</li> <li>David A, <i>et al.</i> 2003. <i>J. Leukoc. Biol.</i> 74:551. (IF)</li> <li>Charles N, <i>et al.</i> 2010. <i>Nat. Med.</i> 16:701. (FC) <u>PubMed</u></li> <li>Thurlow LR, <i>et al.</i> 2010. <i>Infect. Immunol.</i> 128:1128. (FC) <u>PubMed</u></li> <li>Jadhav S, <i>et al.</i> 2001. <i>J. Immunol.</i> 167:5986. (Block)</li> <li>Yoshino N, <i>et al.</i> 2007. <i>Vet. Immunol.</i> 19:27. (FC)</li> <li>Sestak K, <i>et al.</i> 2014. <i>J Immunol.</i> 192:5481. (FC) <u>PubMed</u></li> </ol>

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Sprong T, *et al.* 2003. *Blood* 102:3702. (Block)
 Cash JL, *et al.* 2013. *EMBO Rep.* 14:999. (FC) <u>PubMed</u>
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Antigen 1. Stewart M, et al. 1995. Curr. Opin. Cell Biol. 7:690.

## **References:**