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PII: S0022-1759(15)30042-9
DOI: doi: 10.1016/j.jim.2015.08.015
Reference: JIM 12085

To appear in: *Journal of Immunological Methods*

Received date: 4 May 2015
Revised date: 20 August 2015
Accepted date: 31 August 2015

Please cite this article as: Dong, Matthew B., Jubayer Rahman, M., Tarbell, Kristin V., Flow cytometric gating for spleen monocyte and DC subsets: differences in autoimmune NOD mice and with acute inflammation, *Journal of Immunological Methods* (2015), doi: 10.1016/j.jim.2015.08.015

# Flow cytometric gating for spleen monocyte and DC subsets: differences in autoimmune NOD mice and with acute inflammation

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Keywords: dendritic cells, immune homeostasis, inflammation, multicolor flow cytometry, non-obese diabetic mice, phosphoflow

Abbreviations: antigen presenting cells (APCs), B6.NOD-(*D11Mit167*) H2$^{g7}$/DvsJ (B6.g7), C57BL/6 (B6), conventional dendritic cells (cDCs), dendritic cells (DCs), fluorescence minus one (FMO), forward scatter (FSC-A), interferon (IFN), lineage (Lin), major histocompatibility complexes (MHC), mean fluorescence intensity (MFI), monocyte-derived dendritic cells (moDCs), non-obese diabetic (NOD), paraformaldehyde (PFA), plasmacytoid dendritic cells (pDCs), regulatory T cells (Tregs), room temperature (RT), side scatter (SSC), Signal Transducer and Activators of Transcription 1 phosphorylation (pSTAT1)

Highlights:

-Use of Siglec-H for pDC gating is more specific than BST2, even with inflammation

- DCIR2 (33D1) staining helps separate splenic CD11b$^{+}$ cDC2s from functionally distinct monocyte-derived DCs

- NOD mice have monocyte and DC subset alterations, including an increase in monocyte-