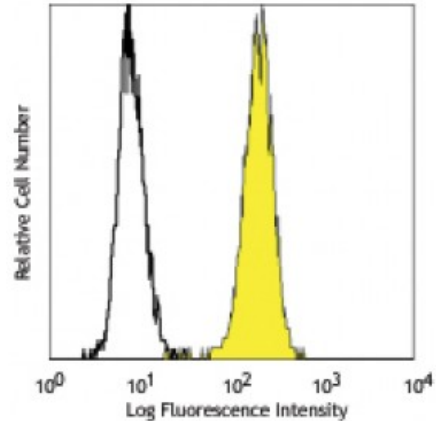


Purified anti-human Galectin-9

Catalog # / Size: 2344510 / 100 µg
Clone: 9M1-3
Isotype: Mouse IgG1, κ
Immunogen: Recombinant peptide from C-terminus of Galectin-9
Reactivity: Human
Preparation: The antibody was purified by affinity chromatography.
Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
Concentration: 0.5



Human acute lymphoblastic leukemia cell line MOLT-4 intracellularly stained with purified 9M1-3 conjugated with PE

Applications:

Applications: Flow Cytometry
Recommended Usage: Each lot of this antibody is quality control tested by intracellular immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is ≤1.0 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.
Application Notes: Additional reported applications (for the relevant formats) include: cell surface staining for flow cytometry¹ and blocking of TIM-3 binding to galectin-91.
Application References: 1. Klibi J, *et al.* 2009. *Blood* 113:1957. (FC, Block)
 2. Sada-Ovalle I, *et al.* 2012. *J. Immunol.* 189:5896. [PubMed](#)

Description: Galectin-9 is a mammalian lectin with a molecular weight around 50 kD. It is a member of the β-galactoside-binding family. With two conserved carbohydrate recognition domains (CRDs), galectin-9 binds small β-galactosides as well as complex glycoconjugates. HAVCR2/TIM3 has been reported as one of its ligands. Galectin-9 may be retained intracellularly or transported to the cell surface where it can be cleaved to generate a soluble form. Galectin-9 is expressed by lymphocytes, dendritic cells, granulocytes, eosinophils, astrocytes, endothelial cells, fibroblasts, and thymus epithelial cells. It can be induced by cytokines in various cell types and is involved in cell aggregation, adhesion, chemotaxis, and apoptosis; galectin-9 induces regulatory T cells and suppresses Th1 and Th17 responses.

Antigen References: 1. Seki M, *et al.* 2008. *Clin. Immunol.* 127:78.
 2. Tsuboi Y, *et al.* 2007. *Clin. Immunol.* 124:221.
 3. Zhu C, *et al.* 2005. *Nat. Immunol.* 6:1245.
 4. Dunphy JL, *et al.* 2002.