

**PE anti-mouse Galectin-9**

**Catalog # / Size:** 1280520 / 200 µg  
1280515 / 50 µg

**Clone:** RG9-35

**Isotype:** Rat IgG2a, κ

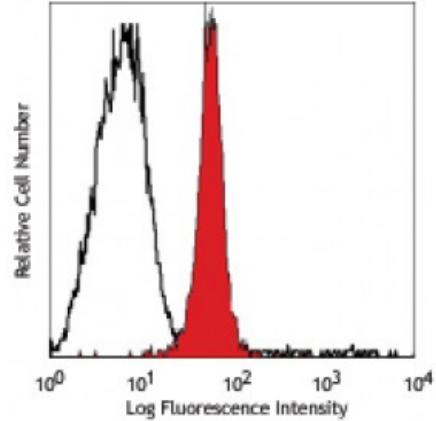
**Immunogen:** Recombinant mouse galectin-9

**Reactivity:** Mouse

**Preparation:** The antibody was purified by affinity chromatography, and conjugated with PE under optimal conditions. The solution is free of unconjugated PE and unconjugated antibody.

**Formulation:** Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.

**Concentration:** 0.2



Intracellular staining of mouse thymocytes with the RG9-35 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 ≤ 0.5 microg per 10<sup>6</sup> cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.

**Application References:** 1. Fukushima A, *et al.* 2008. *Int. Arch. Allergy Immunol.* 146:36. (FA)  
2. Hou H, *et al.* 2014. *PLoS One.* 9:110585. [PubMed](#)

**Description:** Galectin-9 is a mammalian lectin with a molecular weight of 40 kD that has two conserved carbohydrate recognition domains (CRDs) and forms homodimers. It recognizes N-acetyllactosamine (Galβ1-4GlcNAc) and T-antigen (Galβ1-3GalNAc). Tim-3 has been reported as its ligand. Galectin-9 is expressed by lymphocytes, dendritic cells, granulocytes, eosinophils, astrocytes, endothelial cells, fibroblasts, and thymus epithelial cells. It may be retained intracellularly or transported to the cell surface whereby cleavage generates a soluble form. Galectin-9 is involved in events such as cell aggregation, adhesion, chemotaxis, and apoptosis, and is important for the regulation of the immune response. Galectin-9 induces regulatory T cells, and suppresses Th1 and Th17 responses.

**Antigen References:** 1. Klibi J, *et al.* 2009. *Blood* 113:1957  
2. Seki M, *et al.* 2008. *Clin Immunol* 127:78  
3. Tsuboi Y, *et al.* 2007. *Clin Immunol* 124:221  
4. Zhu C, *et al.* 2005. *Nat Immunol*