

**DESCRIPTION**

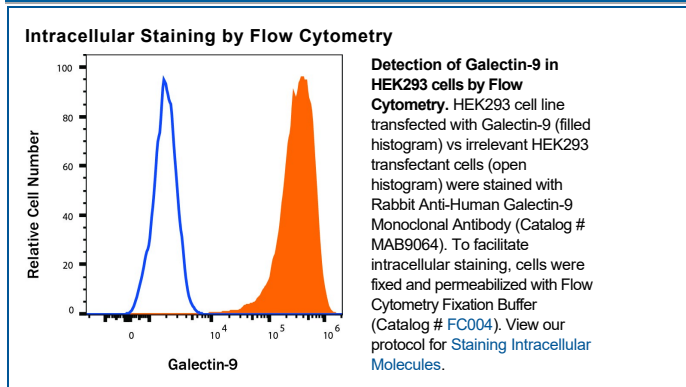
<b>Species Reactivity</b>	Human
<b>Specificity</b>	Detects Human Galectin-9 in direct ELISA.
<b>Source</b>	Recombinant Monoclonal Rabbit IgG Clone # 2315B
<b>Purification</b>	Protein A or G purified from hybridoma culture supernatant
<b>Immunogen</b>	Human embryonic kidney cell, HEK293 derived human Galectin-9 Met1-Thr323 Accession # O00182
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose.

**APPLICATIONS**

**Please Note:** Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

	<b>Recommended Concentration</b>	<b>Sample</b>
<b>Intracellular Staining by Flow Cytometry</b>	0.25 µg/10 <sup>6</sup> cells	HEK293 cell line transfected with Galectin-9 vs irrelevant HEK293 transfectant cells, fixed and permeabilized with Flow Cytometry Fixation Buffer (Catalog # FC004).

**DATA**



**PREPARATION AND STORAGE**

<b>Reconstitution</b>	Reconstitute at 0.5 mg/mL in sterile PBS.
<b>Shipping</b>	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.
<b>Stability &amp; Storage</b>	<p><b>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</b></p> <ul style="list-style-type: none"> <li>• 12 months from date of receipt, -20 to -70 °C as supplied.</li> <li>• 1 month, 2 to 8 °C under sterile conditions after reconstitution.</li> <li>• 6 months, -20 to -70 °C under sterile conditions after reconstitution.</li> </ul>

**BACKGROUND**

Galectins comprise a family of multifunctional carbohydrate-binding proteins with specificity for N-acetyl-lactosamine-containing glycoproteins. At least 14 mammalian Galectins share structural similarities in their carbohydrate recognition domains (CRD), forming three groups: prototype (one CRD), tandem-repeat (two CRDs), and chimeric (one CRD, unique N-terminus) (1, 2). Full length Galectin-9 is a widely expressed 39 kDa tandem-repeat Galectin that contains two CRDs connected by a linker region (3). Progressive deletion within the linker region generates a 36 kDa isoform, also known as Ecalectin or UAT, as well as a 35 kDa isoform (4). This recombinant protein corresponds to the Ecalectin isoform of human Galectin-9 and shares 70% and 73% aa sequence identity with the corresponding regions of mouse and rat Galectin-9, respectively. Galectin-9 exhibits a wide range of activities. All three isoforms function as eosinophil chemoattractants (5, 6). This activity is destroyed by thrombin-mediated cleavage within the linker region of the long isoform, although the Ecalectin isoform is resistant to thrombin (7). Galectin-9 binds to carbohydrate moieties of IgE, thereby preventing immune complex formation, mast cell degranulation, and asthmatic and cutaneous anaphylaxis reactions (8). Independent of its lectin properties, Galectin-9 induces the maturation of dendritic cells which promote Th1 polarization (9). Galectin-9 induces cellular apoptosis in part by direct binding to TIM-3 (10, 11). Its interaction with TIM-3 inhibits Th1 cell and CD8<sup>+</sup> cytotoxic T cell responses and also promotes regulatory T cell differentiation and activity (11, 12). Galectin-9 suppresses tumor cell metastasis by interfering with the associations between hyaluronic acid and CD44 and between VCAM-1 and Integrin α4β1 (13). The Ecalectin isoform (UAT; urate transporter) can also be expressed as an integral membrane protein and mediate the cellular efflux of urate (14).

**References:**

1. Heusschen, R. *et al.* (2013) *Biochim. Biophys. Acta* **1836**:177.
2. Elola, M. T. *et al.* (2007) *Cell. Mol. Life Sci.* **64**:1679.
3. Tureci, O. *et al.* (1997) *J. Biol. Chem.* **272**:6416.
4. Chabot, S. *et al.* (2002) *Glycobiology* **12**:111.
5. Matsumoto, R. *et al.* (2002) *J. Immunol.* **168**:1961.
6. Sato, M. *et al.* (2002) *Glycobiology* **12**:191.
7. Nishi, N. *et al.* (2006) *Glycobiology* **16**:15C.
8. Niki, T. *et al.* (2009) *J. Biol. Chem.* **284**:32344.
9. Dai, S.-Y. *et al.* (2005) *J. Immunol.* **175**:2974.
10. Seki, M. *et al.* (2007) *Arthritis Rheum.* **56**:3968.
11. Zhu, C. *et al.* (2005) *Nat. Immunol.* **6**:1245.
12. Sehrawat, S. *et al.* (2010) *PLoS Pathogens* **6**:e1000882.
13. Nobumoto, A. *et al.* (2008) *Glycobiology* **18**:735.
14. Leal-Pinto, E. *et al.* (2002) *Am. J. Physiol. Renal Physiol.* **283**:F150.