

## Product datasheet for **CL049B**

### Galectin-3 Rat Monoclonal Antibody [Clone ID: M3/38]

#### Product data:

|                       |   |
|-----------------------|---|
| Product Type:         | Primary Antibodies  |
| Clone Name:           | M3/38   |
| Applications:         | ELISA, FC, IHC  |
| Recommended Dilution: | CL049B can be used for Indirect Immunofluorescence staining, including flow cytometric analysis of live cells. The addition of propidium iodide is optional; its use eliminates staining artifacts caused by dead cells (1). This clone is also suitable for frozen sections (4,5) and ELISA (6).   |
| Reactivity:           | Mouse   |
| Host:                 | Rat   |
| Isotype:              | IgG2a   |
| Clonality:            | Monoclonal  |
| Immunogen:            | Plasma membrane glycoproteins from C57BL/6 mouse thioglycollateelicited peritoneal exudate  |
| Specificity:          | Anti-Mac-2 monoclonal antibody specifically binds the mouse Mac-2 antigen. The antibody recognizes a 32,000 Dalton surface antigen found on a subpopulation of mouse macrophages. It reacts with peritoneal exudate macrophages where the exudate is provoked by thioglycollate, protease peptone (20%), macrophages of lymphoid and non-lymphoid tissues, interdigitating dendritic cells and Langerhans cells. Mac-2 is also expressed in the cytoplasm on non-elicited resident macrophages; 5% are strongly reactive, the remaining 95% show much weaker staining. The antibody does not react with peritoneal exudate macrophages where the exudate is provoked by <i>Listeria monocytogenes</i> , lipopolysaccharide or concanavalin A. It also does not react with peritoneal macrophages, splenic macrophages, granulocytes, thymocytes, peripheral lymph node cells and with 99% of bone marrow cells. |
| Formulation:          | PBS containing 0.02% Sodium Azide as preservative and EIA Grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml.<br>Label: Biotin<br>State: Liquid   |
| Concentration:        | lot specific  |
| Conjugation:          | Biotin  |



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| <b>Storage:</b>       | Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.<br>Avoid freeze/thaw cycles.   |
| <b>Stability:</b>     | Shelf life: One year from despatch.   |
| <b>Database Link:</b> | <a href="#">P16110</a>  |
| <b>Background:</b>    | <p>Galectins are a new family of animal lectins which appear to exhibit a variety of biological functions. Lectins, of either plant or animal origin, are carbohydrate binding proteins that interact with glycoprotein and glycolipids on the surface of animal cells. The Galectins are lectins that recognize and interact with beta-galactoside moieties.</p> <p>Galectin 3 is one of the more extensively studied members of this family and is a 30 kDa protein. Due to a C-terminal carbohydrate binding site, Galectin 3 is capable of binding IgE and mammalian cell surfaces only when homodimerized or homooligomerized. Galectin 3 is normally distributed in epithelia of many organs, in various inflammatory cells, including macrophages, as well as dendritic cells and Kupffer cells. The expression of this lectin is up-regulated during inflammation, cell proliferation, cell differentiation and through trans-activation by viral proteins.</p> |
| <b>Synonyms:</b>      | Mac-2, Lgals3, GAL3, GALBP, CBP35, L-31   |

Note: Protocol: **FLOW CYTOMETRY ANALYSIS:**

**Method:**

1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population with separation medium.
2. Wash 2 times.
3. Resuspend the cells to a concentration of  $2 \times 10^7$  cells/ml in media A. Add 50  $\mu$ l of this suspension to each tube (each tube will then contain  $1 \times 10^6$  cells, representing 1 test).
4. To each tube, add 2  $\mu$ g\* of CL049B per  $10^6$  cells.
5. Vortex the tubes to ensure thorough mixing of antibody and cells.
6. Incubate the tubes for 30 minutes at 4°C.
7. Wash 2 times at 4°C.
8. Add 100  $\mu$ l of secondary antibody (Streptavidin-FITC) at a dilution recommended by the manufacturer.
9. Incubate the tubes at 4°C for 30-60 minutes.  
(It is recommended that the tubes are protected from light since most fluorochromes are light sensitive).
10. Wash 2 times at 4°C in media B.
11. Resuspend the cell pellet in 50  $\mu$ l ice cold media B.
12. Transfer to suitable tubes for flow cytometric analysis containing 15  $\mu$ l of propidium iodide at 0.5 mg/ml in PBS. This stains dead cells by intercalating in DNA.

**Media:**

- A. Phosphate buffered saline (pH 7.2) + 5% normal serum of host species + sodium azide (100  $\mu$ l of 2M sodium azide in 100 mls).
- B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100  $\mu$ l of 2M sodium azide in 100 mls).