

PRODUCT INSERT

RAT anti-MOUSE CD44

Product Code	Form	Volume	Antibody*	Excitation (nm)	Peak Emission (nm)	Matching Isotype Controls	
RM5700	Purified	1.0 ml	200 µg	N/A	N/A	Rat IgG2b Purified	Code R2b00
RM5715	Biotin	1.0 ml	100 µg	N/A	N/A	Rat IgG2b Biotin	Code R2b15
RM5715-3	Biotin	3.0 ml	300 µg				
RM5726	Alexa Fluor ^{®†} 405	1.0 ml	100 µg	405	421	Rat IgG2b Alexa Fluor 405	Code R2b26
RM5701	FITC	1.0 ml	100 µg	488	525	Rat IgG2b FITC	Code R2b01
RM5701-3	FITC	3.0 ml	300 µg				
RM5704	R-PE	0.5 ml	50 µg	488	575	Rat IgG2b R-PE	Code R2b04
RM5704-3	R-PE	3.0 ml	300 µg				

PRODUCT DESCRIPTION

Rat monoclonal antibody to mouse CD44 antigen

Clone: IM7.8.1

Isotype: Rat IgG2b

Immunogen: myeloid cell line M1 induced with 1x10⁻⁵ M dexamethazone¹

Lot No.: See label **Expiration:** See label

Buffer: Phosphate buffered saline (PBS)

Preservatives: 0.1% *sodium azide*. Sodium azide is an extremely toxic and dangerous compound particularly when combined with acids or metals. Solutions containing sodium azide should be disposed of properly.

Stabilizer: For conjugated products only, a highly purified grade of BSA has been added as a stabilizing protein.

STORAGE & HANDLING

Store reagents at 2-8°C. For fluorochrome conjugated antibodies, light exposure should be avoided. Use dim light during handling, incubation with cells and prior to analysis. It is recommended that cells be analyzed within 18 hours of staining. If the reagent is being diluted, it is recommended that only the quantity to be used within one week be diluted.

PRODUCT CHARACTERIZATION

Antigen Specificity: The IM7 monoclonal antibody (mAb) reacts with CD44¹ antigen. CD44 antigen is also known as Pgp or Ly-24, a polymorphic glycoprotein, which is broadly distributed on hematopoietic cells and a variety of non-hematopoietic cells^{1,2}.

CD44 antigen is a cell adhesion receptor and its primary ligand is hyaluronan³. The IM7 mAb recognizes both Ly-24.1 and Ly-24.2, as well as every isoform of CD44⁴ antigen. Applications of the IM7 mAb include immunoprecipitation, immunostaining for flow cytometry, detection of soluble CD44 antigen with ELISA, and complement-mediated depletion^{1,2,3}.

PRODUCT QUALITY CONTROL

Every lot is tested by flow cytometry using freshly harvested mouse splenocytes. From this testing it is recommended that between 0.1 and 0.25 µg of antibody be used per 1 x 10⁶ cells in a 100 µl staining volume. Because conditions may vary, it is recommended that each investigator determine the optimal amount of antibody to be used for each application.

REFERENCES:

1. Trowbridge, I. S., J. Lesley, R. Schulte, R. Hyman, and J. Trotter. 1982. Biochemical characterization and cellular distribution of polymorphic, murine cell-surface glycoprotein expressed on lymphoid tissues. *Immunogenetics* 15: 299–312.
2. Lesley, J., and I. S. Trowbridge. 1982. Genetic characterization of a polymorphic murine cell-surface glycoprotein. *Immunogenetics* 15: 313–320.
3. Katoh, S., J. B. McCarthy, and P. W. Kincade. 1994. Characterization of soluble CD44 in the circulation of mice. Levels are affected by immune activity and tumor growth. *J. Immunol.* 153: 3440–3449.
4. Lesley, J., R. Hyman, and P. W. Kincade. 1993. CD44 and its interaction with extracellular matrix. *Adv. Immunol.* 54: 271–335.

* The amount of antibody is determined by measuring the optical density using a spectrophotometer. The antibody titer is verified by immunofluorescent staining and flow cytometric analysis.

† The Alexa Fluor dye conjugates in this product are sold under license from Molecular Probes, Inc., and are covered by pending and issued patents.

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