# Mouse Anti-Rat IgG2b Monoclonal Antibody, FITC Conjugated 

Mouse, Monoclonal (IgG2b FITC)
Cat. No. DMAB4858
Lot. No. (See product label)

## PRODUCT INFORMATION

Product Overview: Mab to $\lg _{2 \mathrm{~b}}$;
Mouse Monoclonal Antibody to Rat Immunoglobulin $\mathrm{G}_{2 \mathrm{~b}}$ $\left(\lg G_{2 b}\right),{ }_{v 2 b}$ heavy chain
Clone: 3C11A9
Ig Isotype: Mouse $\lg _{2 b} \mathrm{~K}$
Format: Fluores cein (FITC) Conjugate
Quality: 0.5 mg
Specificity: Reacts with the $\gamma 2 b$ heavy chain ( Fc ) of rat $\lg _{2 b}$
Applications: Identification and enumeration of $\lg _{2 b}{ }^{+}$cells by immunofluorescence; Second step reagent for rat $\lg \mathrm{G} 2 \mathrm{~b}$ monoclonal antibodies; ELISA
Characterization: To ensure lot-to-lot consistency, each batch of monoclonal antibody is tested as a second step reagent by flow cytometry and/or ELISA to conform to characteristics of a standard reference reagent. Representative data are included in this product insert.
Working Dilutions: Flow Cytometry: $\leq 0.3 \mu \mathrm{~g} / 10^{6}$ cells Other Applications: Since applications vary, each investigator should determine the optimum working dilutions of the product that is appropriate for their specific needs.
Handling And Storage: The fluores cein (FITC) conjugate is supplied as 0.5 mg in 1.0 mL of $\mathrm{PBS} / \mathrm{NaN} 3$. Store at $2-8^{\circ} \mathrm{C}$.
Warning: Reagents contain sodium azide. Sodium azide is very toxic if ingested or inhaled. Avoid contact with skin, eyes, or clothing. Wear eye or face protection when handling. If skin or eye contact occurs, wash with copious amounts of water. If ingested or inhaled, contact a physician immediately. Sodium azide yields toxic hydrazoic acid under acidic conditions. Dilute azide-containing compounds in running water before discarding to avoid accumulation of potentially explosive deposits in lead or copper plumbing.

## REFERENCES

1. Engberg K, Frank CW. Biomed Mater. 2011 Aug 26;6 (5):055006.
2. Larsen CJ. Bull Cancer. 2011 Jul;98(7):719-22. French.
3. Ueno E, Hisajima T, Nakano M, Goris RC, Funakoshi K. Histol Histopathol. 2011 Oct;26(10):1317-26.

## BACKGROUND

Introduction: Immunoglobulin G (lgG) are antibody molecules Each $\operatorname{lgG}$ is composed of four peptide chains - two heavy chains $\gamma$ and two light chains. Each $\operatorname{lgG}$ has two antigen binding sites. Other Immunoglobulins may be described in terms of polymers with the $\lg$ structure considered the monomer. $\lg$ G molecules are synthesized and secreted by plasma B cells. lgG antibodies are large molecules of about 150 kDa composed of 4 peptide chains. It contains 2 identical heavy chains of about 50 kDa and 2 identical light chains of about 25 kDa , thus a tetrameric quatemary structure. The two heavy chains are linked to each other and to a light chain each by disulfide bonds. The resulting tetramer has two identical halves, which together form the Y -like shape. Each end of the fork contains an identical antigen binding site. The Fc regions of IgGs bear a highly conserved N -glycosylation site. The N -glycans attached to this site are predominantly core-fucosylateddiantennary structures of the complextype. In addition, small amounts of these N -glycans also bear bisecting GIcNAc and $\alpha$ - 2,6 -linked sialic acid residues.
Keywords: $\lg \mathrm{G} 2 \mathrm{~b}$; Immunoglobulin G 2 b ; $\lg \mathrm{G}_{2} \mathrm{bv}_{2 \mathrm{~b}}$; Immunoglobulin $\mathrm{G}^{2 b} \mathrm{y}_{2 b}$; $\lg \mathrm{G} 2 \mathrm{~b}$ heavy chain; Immunoglobulin G 2 b heavy chain; $\lg \mathrm{G}^{2 b} \gamma_{2 b}$ heavy chain; Immunoglobulin $\mathrm{G}_{2} \mathrm{~b} \gamma_{2 b}$ heavy chain


IMMUNOFLUORESCENT STAINING
Amount Used: $0.3 \mu \mathrm{~g} / 10^{6}$ cells
BALB/c splenocytes were incubated with rat lgG2b isotype, rat anti-mouse CD22 (2D6, rat $\lg G 1$ ), rat anti-mouse CD8 (53-6.7, rat $\lg$ G2a), rat anti-mouse CD45(LCA) (I3/2.3, rat $\lg$ G2b), rat anti-mouse CD24 (30-F1, rat IgG2c), and rat anti-mouse CD45RC (GL24, rat $\lg \mathrm{M}$ ). After washing, the cells were stained with FITC-labeled Mouse Anti-Rat IgG2b. Small
lymphocytes were gated and analyzed on a FACScan ${ }^{\text {TM }}$ flow cytometer (BDIS, San Jose, CA).

