

## # M071-1

Product Information	
Catalog Number:	M071-1
Clone / Isotype:	Sam.C1 / Rat (Wistar) IgG2b
Contents:	FITC-labeled immunoglobulin in 20 mM Tris buffer with 137 mM NaCl, 0.5% BSA and 0.09% (w/v) sodium azide
Size:	1.5 ml / 300 tests

For research use only, not for diagnostic or therapeutic use. This product is no medical device.

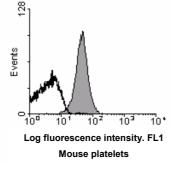
**Specificity:** The Sam.C1 antibody reacts with murine integrin  $\alpha$ 2 chain (CD49b), the 160-kDa transmembrane glycoprotein that non-covalently associates with the integrin  $\beta$ 1 subunit to form the integrin  $\alpha$ 2 $\beta$ 1 complex known as VLA-2. Integrin  $\alpha$ 2 $\beta$ 1, a receptor for collagen and laminin, is expressed on platelets, epithelial cells, and activated lymphocytes.

**Preparation and Storage:** The antibody was purified from hybridoma cell culture supernatant by Protein G-Sepharose chromatography. The antibody was conjugated with FITC under optimum conditions. The solution is free of unbound FITC. Store product undiluted at 4°C and avoid prolonged exposure to light. Stable for one year from date of shipment. Do not freeze.

**Usage:** The antibody preparation is optimized for flow cytometric applications: Use 5  $\mu$ l to stain ~10<sup>6</sup> platelets or ~0.5x10<sup>6</sup> cells in a recommended volume of 25  $\mu$ l. Incubate for 15 minutes at room temperature, stop reaction by addition of 400  $\mu$ l PBS and analyze samples within 30 minutes. For immunofluorescent staining of acetone-fixed frozen sections, the appropriate dilution must be determined individually.

**Caution:** Sodium azide is a reversible inhibitor of oxidative metabolism; therefore, antibody preparations containing this preservative agent must not be used in cell cultures nor injected into animals. Sodium azide may be removed by washing stained cells or plate-bound antibody or dialyzing soluble antibody in sodium azide-free buffer.

Detection of integrin  $\alpha$ 2 chain on mouse platelets Mouse blood was diluted 1:20 and 25 µl of this dilution were stained with 5 µl control IgG-FITC (emfret Analytics, P190-1, *black line*) or Sam.C1-FITC (*shaded area*) for 15 min at RT and analyzed directly. Platelets were gated by FSC/SSC characteristics.



## **References:**

1. Mendrick, DL and Kelly, DM (1993) Lab.Invest. 69:690-702.

- 2. Mendrick, DL, Kelly, DM, duMont, SS, and Sandstrom, DJ (1995) *Lab.Invest.* 72:367-375.
- 3. emfret Analytics. Unpublished results.