

DESCRIPTION

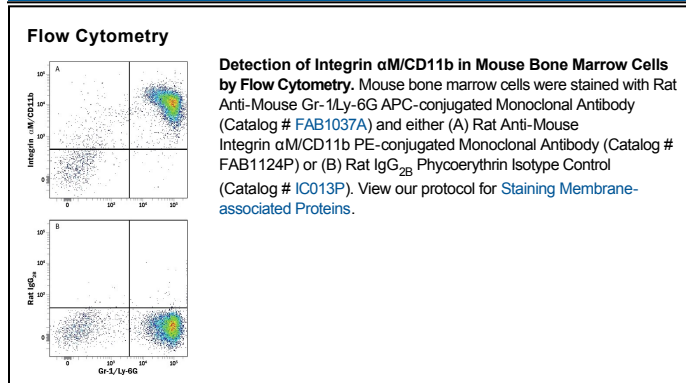
Species Reactivity	Mouse
Specificity	Detects mouse Integrin α M/CD11b. Cross-reaction with human Integrin α M has been reported (1, 2).
Source	Monoclonal Rat IgG _{2B} Clone # M1/70
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	Con A-activated C57BL/10 splenocytes
Conjugate	Phycoerythrin Excitation Wavelength: 488 nm Emission Wavelength: 565-605 nm
Formulation	Supplied in a saline solution containing BSA and Sodium Azide. See Certificate of Analysis for details. *Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Sheet (SDS) for additional information and handling instructions.

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.

	Recommended Concentration	Sample
Flow Cytometry	10 μ L/10 ⁶ cells	See Below

DATA



PREPARATION AND STORAGE

Shipping	The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.
Stability & Storage	Protect from light. Do not freeze. <ul style="list-style-type: none"> 12 months from date of receipt, 2 to 8 °C as supplied.

BACKGROUND

The Integrin family proteins are heterodimeric transmembrane receptors composed of an α and a β subunit. The Integrin α M subunit, also known as MAC-1 α subunit or CD11b, combines with the Integrin β 2 subunit (CD18) to form the non-covalent heterodimer Integrin α M/ β 2, also known as MAC-1 and complement receptor type 3 (CR3). Integrin α M/ β 2 is expressed on granulocytes, macrophages, dendritic cells and natural killer cells. Upon activation, α M/ β 2 can bind several ligands (including ICAM-1, fibrinogen, and the C3 complement fragment, C3bi) to mediate phagocyte adhesion, migration and ingestion of complement-opsonized particles.

References:

1. Beller, D.J. *et al.* (1982) J. Exp. Med. **156**:1000.
2. Ault, K.A. and T.A. Springer (1981) J. Immunol. **126**:359.