# **Datasheet**

Mouse mAb to CD18 Clone 68-5A5 Isotype IgG2a- $\kappa$ 

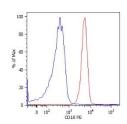


#### Source

A BALB/c mouse was immunized with stimulated human leucocytes. Fusion partner: NS-1.

## **Specifications**

It recognizes a transmembrane glycoprotein of 95 kDa, identified as CD18 or intergrin-2 (Workshop III). It complexes non-covalently with either L, M, or X integrin (CD11a, b, or c) to form the heterodimers. LFA-1, MAC-1, and p150,95, respectively. LFA-1 is the receptor for three members of the Ig supergene family of proteins, ICAM-1 (CD54, ICCAM-2 (CD102), and Mac-1 and p150,95 bind to ICAM-1, fibrinogen, and IC3b. ICAM-3 (CD50). CD18/CD11 heterodimeric molecules are involved with cell/cell and cell/extracellular adhesion in immune and inflammatory responses. This Mab blocks these cellular interactions. 68-5A5 was clustered at the III<sup>rd</sup> International Workshop on human leucocyte differentiation antigens.



**Figure 1:** Human granulocytes stained for CD18 (FACS).

## Species reactivity

Positive: human.

### **Applications**

68-5A5 can be used for staining of frozen tissues, in flow cytometry and for blocking cell-c ell interactions.

Flow cytometry	Frozen sections	Functional studies	Immunofluorescence	Paraffin sections	Western blot
+	+	+	+	-	+

#### **Format**

Produced in tissue culture, contains no host Ig. Antibodies are affinity purified and presented in PBS with 0,02% sodium azide.

Stored at 4°C-8°C, shelf life is at least 24 months after purchase.

#### Dilution advice

- Flow cytometry  $(0.5-1.0 \mu g/million cells in 0.1 ml)$ .
- Functional studies (0,02-2,0  $\mu$ g/ml without azide).
- $\triangleright$  Immunoblotting (1,0-2,0 µg/ml).
- Immunofluorescence (0,5-1,0 μg/ml).
- $\triangleright$  Immunohistology (1-2 µg/ml for 30 min at RT; an appropriate antigen retrieval method for staining formalin-fixed tissues has not been established to date).

#### **Positive control**

Human PBL and tonsil.

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# References

➤ McMichael A.J.et al., Leucocyte typing III, Oxford University Press, Oxford, (1987).