

# Produktinformation



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### SZABO-SCANDIC HandelsgmbH

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Place your order with CEDARLANE<sup>®</sup> or your local distributor. Please contact CEDARLANE<sup>®</sup> for lot specific information.

## FITC Anti-Mouse Platelet Monoclonal Antibody

CL8960F LOT:

#### **DESCRIPTION**:

Cedarlane's anti-mouse platelet monoclonal antibody (clone AIP21) detects an unidentified antigen on mouse platelets that is not identical to CD9, GPIV or integrins. This antigen is also present on B16F10 and KN-3 cells, but not thymocytes or splenocytes. This antibody induces mouse platelet aggregation in the absence of plasma components via an FcR-independent mechanism. A dramatic increase in tyrosine phosphorylation of the 52 kDa Shc protein was observed during AIP21-mediated platelet aggregation.

This antibody is suitable for use in flow cytometry and functional assays.

#### **PRESENTATION:**

100  $\mu$ g (CL8960F) FITC conjugated Ig buffered in PBS, 0.02% NaN<sub>3</sub> and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml.

#### **STORAGE/STABILITY:**

Store at 4°C. For long term storage, aliquot and freeze unused portion at -20°C in volumes appropriate for single usage. Avoid freeze/thaw cycles. Avoid prolonged exposure to light.

#### SPECIFICATIONS:

<u>Clone</u>: AIP21
 <u>Specificity</u>: Mouse Platelets
 <u>Ig Class</u>: Rat IgM
 <u>Format</u>: FITC conjugated Ig buffered in PBS, 0.02% NaN3 and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml.
 <u>Antibody Concentration</u>: 0.1 mg/ml

Continued Overleaf...

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#### FLOW CYTOMETRY ANALYSIS:

#### Method:

- 1. Prepare a cell suspension of mouse platelets in media A.
- 2. Wash 2 times.
- 3. Resuspend the cells to a concentration of  $2x10^7$  cells/ml in media A. Add 50 µl of this suspension to each tube (each tube will then contain  $1x10^6$  cells, representing 1 test).
- 4. To each tube, add  $\sim 1.0 \,\mu g^*$  of **CL8960F**.
- 5. Vortex the tubes to ensure thorough mixing of antibody and cells.
- 6. Incubate the tubes for 30 minutes at  $4^{\circ}$ C.
- 7. Wash 2 times at  $4^{\circ}$ C.
  - (It is recommended that the tubes are protected from light since most fluorochromes are light sensitive).
- 8. Resuspend the cell pellet in 50  $\mu$ l ice cold media B.
- 9. Transfer to suitable tubes for flow cytometric analysis containing 15 μl of propidium iodide at 0.5 mg/ml in PBS. This stains dead cells by intercalating in DNA.

#### Media:

- A. Phosphate buffered saline (pH 7.2) + 5% normal serum of host species + sodium azide (100 µl of 2M sodium azide in 100 mls).
- B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100 μl of 2M sodium azide in 100 mls).

#### **REFERENCES:**

 Kato, Y., *et al.* 1998. A novel anti-platelet monoclonal antibody induces mouse platelet aggregation through an Fc receptor-independent mechanism. *Biochemical and Biophysical Research Communications*, 242: 250-255