# Anti-SPARCL1 / SPARC-like 1 antibody

Catalog Number: 106267



#### Product name

Anti-SPARCL1 / SPARC-like 1 antibody

### **Specificity**

Human SPARCL1 / SPARC-like 1

# **Antibody description**

Mouse monoclonal to SPARCL1 / SPARC-like 1

## Preparation

This antibody was produced from a hybridoma resulting from the fusion of a mouse myeloma with B cells obtained from a mouse immunized with purified, recombinant Human SPARCL1 / SPARClike 1 (rh SPARCL1 / SPARC-like 1; NP\_004675.3; Met1-Phe664). The IgG fraction of the cell culture supernatant was purified by Protein A affinity chromatography.

#### **Formulation**

0.2 µm filtered solution in PBS with 5% trehalose

#### **Storage**

This antibody can be stored at 2°C-8°C for one month without detectable loss of activity. Antibody products are stable for twelve months from date of receipt when stored at -20°C to -80°C.

Preservative-Free.

Sodium azide is recommended to avoid contamination (final concentration 0.05%-0.1%). It is toxic to cells and should be disposed of properly. Avoid repeated freeze-thaw cycles.

#### **Clonality**

Monoclonal

#### Ig Type

Mouse IgG2b

#### **Applications**

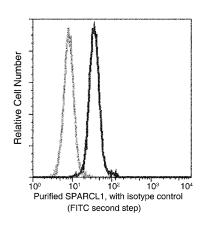
FCM, IF, ICC/IF

#### **Dilutions**

FCM: 0.5-2 µg/Test

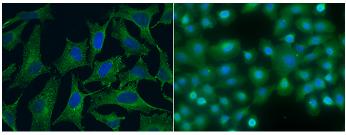
ICC/IF: 10-25 μg/mL

#### **Validations**



SPARCL1 / SPARC-like 1 Antibody, Mouse MAb, Flow cytometric analysis

Flow cytometric analysis of Human SPARCL1 expression on HL60 cells. The cells were treated according to manufacturer's manual (BD Pharmingen  $^{\text{TM}}$  Cat. No. 554714), stained with purified anti-Human SPARCL1, then a FITC-conjugated second step antibody. The fluorescence histograms were derived from gated events with the forward and side light-scatter characteristics of intact cells.



SPARCL1 / SPARC-like 1 Antibody, Mouse MAb, Immunofluorescence

Immunofluorescence staining of Human SPARCL1 in A431 or Hela cells. Cells were fixed with 4% PFA, permeabilzed with 1% Triton X-100 in PBS, blocked with 10% serum, and incubated with

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mouse anti-Human SPARCL1 monoclonal antibody (15  $\mu$ g/ml). Then cells were stained with the Alexa Fluor® 488-conjugated Goat Anti-mouse IgG secondary antibody (left panel, captured by laser

confocal scanning microscope; right panel, captured by fluorescence microscope), countstained with DAPI (blue). Positive staining was localized to cytoplasm.