

For Research Use Only.
Not for use in diagnostic procedures.

Anti-GP2 (Glycoprotein 2) (Human) mAb -Alexa Fluor[®] 488

CODE No. D277-A48

CLONALITY Monoclonal
CLONE 3G7-H9
ISOTYPE Mouse IgG1 κ
QUANTITY 100 μ L, 1 mg/mL

SOURCE Purified IgG from hybridoma supernatant
IMMUNOGEN Human GP2, extracellular domain (recombinant, human Fc fusion protein)
FORMULATION PBS containing 1% BSA and 0.1% ProClin 150
STORAGE This antibody solution is stable for one year from the date of purchase when stored at 4°C.

APPLICATIONS-CONFIRMED

Flow cytometry 1 μ g/mL

SPECIES CROSS REACTIVITY

Species	Human	Mouse	Rat	Hamster
Tissues	Peyer's patches	Peyer's patches	Not tested	Not Tested
Reactivity	+	-		

Entrez Gene ID 2813 (Human)

REFERENCES 1) Hase, K., *et al.*, *Nature* **462**, 226-230 (2009)
This clone is used in this reference.

For more information, please visit our web site <https://ruo.mbl.co.jp/>

LABEL LICENSES:

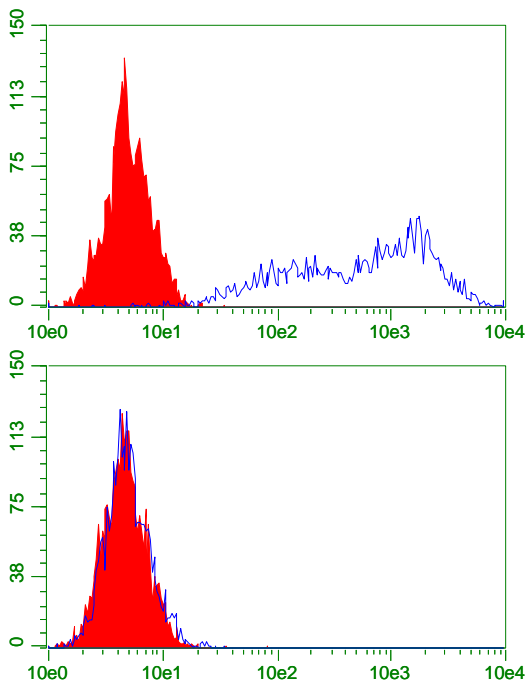
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Flow cytometric analysis

- 1) Wash the cells (5×10^5 cells/sample) 1 time with 1 mL of washing buffer [PBS containing 2% fetal calf serum (FCS)].
- 2) Add 20 μ L of Clear Back (human Fc receptor blocking reagent, MBL; code no. MTG-001) to the cell pellet after tapping. Mix well and incubate for 5 min. at room temperature.
- 3) Add 40 μ L of the primary antibody at the concentration as suggested in the **APPLICATION** diluted in the washing buffer. Mix well and incubate for 30 min. at room temperature. (Optimization of antibody concentration or incubation condition is recommended if necessary.)
- 4) Wash the cells 2 times with 1 mL of the washing buffer.
- 5) Resuspend the cells with 500 μ L of the washing buffer and analyze by a flow cytometer.

(Positive control for Flow cytometry; Transfectant)



Flow cytometric detection of GP2 (Human) in transfectant

Cell

Upper: Human GP2/293T

Lower: Parental cell (293T)

Antibody

Open: D277-A48

Closed: Isotype control (M075-A48)