

Product datasheet for **AM26668RP-N**

CD95 (FAS) Mouse Monoclonal Antibody [Clone ID: UB2]

Product data:

Product Type:	Primary Antibodies
Clone Name:	UB2
Applications:	FC
Recommended Dilution:	Flow cytometry: 20 µl (ready for use). For details see protocol below.
Reactivity:	Human
Host:	Mouse
Isotype:	IgG1
Clonality:	Monoclonal
Immunogen:	Recombinant human Fas
Specificity:	This antibody recognizes human Fas antigen.
Formulation:	PBS Label: PE State: Liquid Ig fraction Stabilizer: 1% BSA Preservative: 0.09% NaN ₃
Purification:	Affinity chromatography on protein A
Conjugation:	PE
Storage:	Store at 2-8 °C.
Stability:	Shelf life: one year from despatch.
Gene Name:	Fas cell surface death receptor
Database Link:	Entrez Gene 355 Human P25445
Background:	It is now widely accepted that apoptosis plays an important role in the selection of immature thymocytes and Ag-primed peripheral T cells. Fas antigen is a cell-surface protein that mediates apoptosis. It is expressed in various tissues including the thymus and has structural homology to a number of cell-surface receptors, including tumor necrosis factor receptor and nerve growth factor receptor.



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Synonyms: FASLG receptor, Apo-1 antigen, APT1, FAS1, TNFRSF6

Note: This product was originally produced by MBL International.

Protocol: Flow cytometric analysis for floating cells

We usually use Fisher tubes or equivalents as reaction tubes for all steps described below.

1) Wash the cells 3 times with washing buffer [PBS containing 2% fetal calf serum (FCS) and 0.1% NaN₃].

2) Resuspend the cells with washing buffer (5×10^6 cells/mL).

3) Add 50 μ L of the cell suspension into each tube, and centrifuge at 500 x g for 1 minute at room temperature (20~25 o C). Remove supernatant by careful aspiration.

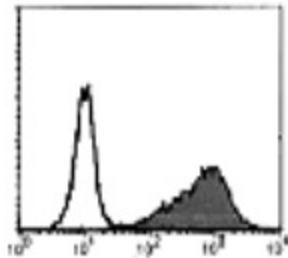
4) Add 10 μ L of normal goat serum containing 1 mg/mL normal human IgG and 0.1% NaN₃ to the cell pellet after tapping. Mix well and incubate for 5 minutes at room temperature.

5) Add 20 μ L of PE labeled anti-Fas monoclonal antibody (UB2). Mix well and incubate for 30 minutes at room temperature.

6) Add 1 mL of the washing buffer followed by centrifugation at 500 x g for 1 minute at room temperature. Remove supernatant by careful aspiration.

7) Resuspend the cells with 500 μ L of the washing buffer and analyze by a flow cytometer. (Positive control for Flow cytometry; transfectant)

Product images:



Flow cytometric analysis of human Fas expression on transfectant. Open histogram indicates the reaction of isotypic control to the cells. Shaded histogram indicates the reaction of AM26668RP-N to the cells.