Anti-Human CD98 (FG1/8)



\ \	REF	Σ	\Diamond		[<u>•</u> 3
PURE	98PU	1 mg	1 mg/ml		
PURE	98PU-01MG	100 µg	1 μg/test	0,01 mg/ml	
FITC	98F-100T	100 test	20 µL/test	2 mg/ml	
PE	98PE-100T	100 test	20 μL/test	2 mg/ml	RUO
PerCP	98PP-100T	100 test	5 µL/test	0,05 mg/ml	KOO
APC	98A-100T	100 test	5 μL/test	0,05 mg/ml	
Biotin	98B-01MG	100 µg	l μg/test	0,01 mg/ml	
CF-Blue	98CFB-100T	100 test	5 µL/test	0,05 mg/ml	l

1. PRODUCT DESCRIPTION

Clone: FG1/8:

Isotype: IqG1;

Tested application: flow cytometry;

Immunogen: The anti-CD98 monoclonal antibody derives from T cells from leukemic HPB-ALL;

Species reactivity: Human;

Storage instruction: store in the dark at 2-8 °C:

Storage buffer: aqueous buffered solution containing protein stabilizer and 0.09% sodium azide (NaNa);

Recommended usage: Immunostep's CD98, clone FGI/8, is a monoclonal antibody intended for the identification and enumeration of human leukocyte lineages using flow cytometry. This reagent is effective for direct immunofluorescence staining of human tissue for flow cytometric analysis using I test for 106 cells;

Presentation: liquid;

Source: Supernatant proceeding from an in vitro cell culture of a cell hybridoma;

Purification: Affinity chromatography:

Other names: 4F2, FRP-1, RL-388, 4F2hc, 4F2 heavy chain antigen, Lymphocyte activation antigen 4F2 large subunit, Solute carrier family 3 member 2;

Gene ID: 6520:

Molecular weight: 125 kDa.

2. ANTIGEN DETAILS

Large description: This antibody reacts with the CD98 antigen, which is broadly expressed on peripheral blood lymphocytes, monocytes and granulocytes (low) .It is also expressed on nonhematopoietic cells. Reports suggest that CD98 may be involved in the regulation of cellular activation, CD98 is up-regulated on leukocytes and in inflammatory lesions, and strongly expressed by neoplastic cells. However, expression is not hematopoietic specific. CD98 is expressed in HUVECS and at low levels in resting peripheral blood T-lymphocytes and quiescent fibroblasts. Also it is expressed in fetal liver and in the astrocytic process of primary astrocytic gliomas, in retinal endothelial cells and in the intestinal epithelial cell line C2BBel.(1-3)

3. WARRANTY

Warranted only to conform to the quantity and contents stated on the label or in the product labelling at the time of delivery to the customer. Immunostep disclaims hereby other warranties.

Immunostep's sole liability is limited to either the replacement of the products or refund of the purchase price.

ADDITIONAL INFORMATION

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5. PROTOCOL

Direct Immunofluorescence Cell Surface Staining Protocol

- 1. Transfer 100 ul (106 cells/test) of the sample to a 12 x 75 mm polystyrene
- 2. Add the suggested volume indicated on the antibody vial to the 12x75 mm cytometer tube.
- 3. Mix well and incubate in the dark at room temperature at 4 °C for 30 minutes or at room temperature (20-25 °C) for 15 minutes.
- 4. After the incubation period, add 1,5 ml of an erythrocyte-lysing solution and mix. Incubate at room temperature in the darkness (the blood should be well mixed with the lysing solution).
- 5. Centrifuge tubes at 540xg for 5 minutes. The supernatant is removed with a Pasteur pipette or with a vacuum pump.
- 6. Resuspend and wash with 3-5 mL of PBS at 540xg for 5 min.
- 7. After removing the supernatant and resuspending the cell pellet, add 300 µL of PBS and adquire on the flow cytometer are recorded.
- 8. Analyse on a flow cytometer or store at 2-8 °C in the dark until analysis. Samples can be run up to 24 hours after lysis.

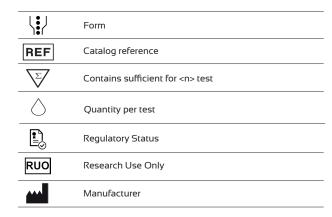
Indirect Immunofluorescence Cell Surface Staining Protocol

- 1. Transfer 100 ul (106 cells/test) of the sample to a 12 x 75 mm polystyrene test tube
- 2. Add purified reagent according to manufacturer's recommendation and mix gently with a vortex mixer
- 3. Incubate in the dark at room temperature at 4 °C for 30 minutes or at room temperature (20-25 °C) for 15 minutes.
- 4. Add 2 mL 0.01 mol/L PBS (It betters that it containing 2% bovine serum albumin) and resuspend the cells by using a vortex mixer. Centrifuge at 540xg for 5 min in order to remove the McAb not bound to its antigen.
- 5. Add a secondary conjugated reagent with some fluorochrome and mix. Incubate at room temperature for 15 min in the dark. The absence of light is necessary as the fluorochrome is photoinstability.
- 6. After the incubation period, add 1.5 ml of an erythrocyte-lysing solution and mix. Incubate at room temperature in the darkness (the blood should be well mixed with the lysing solution). Centrifuge at 540xg for 5 minutes. The supernatant is removed with a Pasteur pipette or with a vacuum pump.
- 7. Resuspend and a made a final wash with 3-5 mL of PBS at 540xg for 5 min.
- 8. After removing the supernatant and resuspending the cell pellet, add 300 µL of PBS and adquire on the flow cytometer are recorded.
- 9. Analyse on a flow cytometer or store at 2-8 °C in the dark until analysis. Samples can be run up to 24 hours after lysis.

REFERENCES

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7. **EXPLANATION OF SYMBOLS**



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