Hu-CD3/CD19 2 Color FCM Reagent: *sc-3952*



BACKGROUND

Human CD3/CD19: sc-3952 is a direct immunofluorescence reagent formatted to identify and determine the percentage of human T lymphocytes and B lymphocytes in erythrocyte-lysed whole blood, based on cell-surface antigen expression. CD3 identifies T lymphocytes and non-covalently associates with either α/β or γ/δ TCR (1). NK lymphocytes identified as CD3⁻ and CD16⁺ and/or CD56⁺ mediate cytotoxicity against certain tumors and virus infected cells (2). CD19 is present on human B lymphocytes during all stages of B cell maturation, but is lost on plasma cells (3). The total population of T lymphocytes and B lymphocytes are used to characterize and monitor some forms of immunodeficiency and autoimmune disease (4,5).

Antigen Expression	Cell Type Identified	
CD3+	Mature T Lymphocytes	
CD3- CD19+	Total B Lymphocytes	

STORAGE

Store at 4° C. Do not freeze. Stable for one year from the date of shipment. Protect reagents from prolonged exposure to light.

PRODUCT

Supplied in 1.0 ml of PBS containing 0.1% azide and 0.1% gelatin. Sufficient for 50 tests. This product has been titrated for optimal performance. Recommended use is 20 uL per test $(1x10^6 \text{ cells})$. For research use only. Not for use in diagnostic procedures.

INSTRUMENT

Human CD3/CD19: sc-3952 is recommended for use with either a single or dual laser Flow Cytometer fitted with appropriate acquisition and analysis software, such as the FACSCalibur™ Flow Cytometer fitted with CellQuest[™] Software by Becton Dickinson.

The flow cytometer must be equipped with a 488 nm laser and must be capable of detecting light scatter (forward and side) and two-color fluorescence with emission detectable in two ranges: 515-545 nm, 562-607 nm.

Antigen	Clone	Isotype	Label*	Detection Range (nm)
CD3	UCH-T1	IgG ₁	FITC	515-545
CD19	SJ25C1	IgG ₁	PE	562-607

*Fluorescent labels include FITC: Fluorescein isothiocyanate; PE: phycoerythrin

ISOTYPE CONTROL

sc-3952 CON (IgG_1 FITC/ IgG_1 PE) is the isotype matched negative control for this system and is suitable for 50 tests.

REFERENCES

1. Exley, M., Terhorst, C., and Wileman, T. 1991. Structure, assembly and intracellular transport of the T cell receptor for antigen. Semin. Immunol. <u>3</u>: 283-297.

2. Fitzgerald-Bocarlsy, P., Herberman, R., Hercend, T., *et al.* 1989. A definition of natural killer cells. In: Ades, E., Lopez, C., eds. Natural Killer Cells and Host Defense. Fasel: Karger; 1.

3. Dörken, B., Möller, P., Pezzutto, A., Schwartz-Albiez, R., and Moldenhauer, G. B-cell antigens: CD19. In: Knapp, W., Dörken, B., Gilks, W.R. *et al.* eds. Leucocyte Typing IV: White Cell Differentiation Antigens. New York, NY: Oxford University Press; 1989: 34-36.

4. Foucar, K. and Goeken, J.A. 1982. Clinical Applications of immunologic techniques to the diagnosis of lymphoproliferative and immunodeficiency disorders. Lab. Med. <u>13</u>: 403-413.

5. Smolen, J.S., Chused, T.M., Leiserson, W.M., Reeves, J.P., Alling, D., and Steinberg, A.D. 1982. Heterogeneity of immunoregulatory T-cell subsets in systemic lupus erythematosus. Correlation with clinical features. Am. J. Med. <u>2</u>: 783-790.