

PRODUCT INFORMATION SHEET

Monoclonal antibodies detecting human antigens

CD3 FITC **CD25** R-PE RUO REF IQP-257FR 50 tests

RUO *For Research Use Only*



Description

In the diagnosis of T-cells, the IL-2 receptor is expressed on activated cells including T cells. It is also present on a subset of thymocytes, HTLV-1 transformed T and B cells, EBV transformed B cells, myeloid precursors and oligodendrocytes. IL-2 induces the expression of the CD25 subunit on NK cells.

CD3

Clone UCHT-1

Isotype Murine IgG1

Specificity Clone UCHT1 produces mouse IgG1 immunoglobulins directed against an epitope expressed on the epsilon chain of the CD3/TcR complex (22-28 kD).

CD3 antibodies are used, in the characterization of various subtypes of chronic lymphoid leukemias. Examples of these chronic T cell leukemias are T-CLL (Sézary Syndrome) and the peripheral T cell lymphoma (ATLL) which co-express CD3, CD2, CD4 and CD5 antigens. The NK cell lymphoma or the intestinal T cell lymphoma, co-express CD3, CD2 and CD8.

HLDA Workshop

6th International Workshop on Human Leukocyte Differentiation Antigens

CD25

Clone B-B10

Isotype Murine IgG1

Specificity CD25 antibodies are used as a marker of cell activation in transplantation patients, for the detection of cells infected with the human T cell Leukemia viruses I and II, and for the immunophenotyping of lymphomas and leukemias. The IL-2 receptor is strongly expressed in Hodgkin's disease, hairy cell leukemia and anaplastic large cell lymphoma. The serum IL-2 receptor is a important parameter in transplantation, inflammatory and malignant disorders.

The functional high affinity IL-2R is composed of a non-covalently associated CD25/CD122/CD132 heterotrimer. The isolated CD25 subunit constitutes a low-affinity IL-2R, while the CD122/CD132 heterodimer binds IL-2 with intermediate affinity. Both the high and intermediate-affinity receptors are important for IL-2 signaling.

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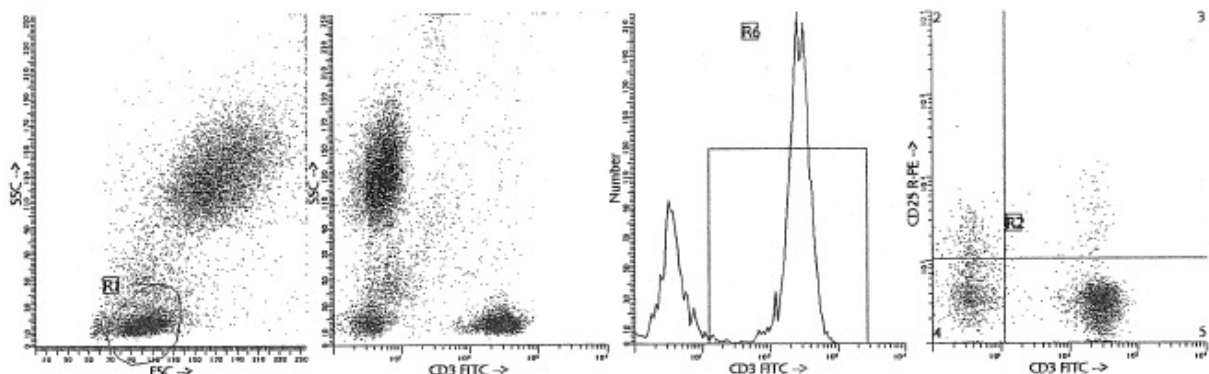
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Usage

All these reagents are effectively formulated for direct immunofluorescent staining of human tissue for flow cytometric analysis using 10 µl/10⁶ leukocytes for singles and 20 µl/10⁶ leukocytes in case of dual and triple combinations. Since applications vary, each investigator should titrate the reagent to obtain optimal results.

Representative Data

IQP-257FR (CD3/CD25) was analyzed by flow cytometry using a blood sample from a healthy volunteer. Direct staining was performed by adding 20 µl of this dual to 100 µl blood sample.



Immunofluorescence staining and lysing protocol

Flow cytometry method for use with dual and triple combinations

1. Add 100 µl of EDTA-treated blood (i.e. approx. 10^6 leukocytes) to a 5 ml reagent tube. The content of one tube is sufficient to perform one test.
For combinations with anti-kappa and/or anti-lambda Ig see application note below.
2. Add to each tube 20 µl of labeled monoclonal antibody combination*.
3. Vortex the tube to ensure thorough mixing of antibody and cells.
4. Incubate the tube for 15 minutes at room temperature in the dark.
5. Add 100 µl of IQ Lyse (IQP-199 ready-to-use) and mix immediately.
6. Incubate for 10 minutes at room temperature in the dark.
7. Add 2 ml of demineralized water and incubate for 10 minutes in the dark.
8. Centrifuge the labeled cell suspension for 2 minutes at 1000 x g.
9. Remove the supernatant and resuspend the cells in 200 µl of PBS**.
10. Analyze by flow cytometry within four hours (alternatively, the cells may be fixed by 0.05% of formaline in buffered saline for analysis the next day. Some antigens are readily destroyed upon fixation and this should be taken into account when using this alternative).

* Appropriate mouse Ig isotype control samples can be included in any labeling study

** PBS: Phosphate Buffered Saline, pH 7.2

Application note for anti-kappa and/or anti-lambda Ig combinations

Add 2 ml of PBS containing 0.001% (v/v) Heparin (**prewarmed to 37 °C**) to the cell suspension
Vortex, centrifuge (2 min at 300x g) and discard the supernatant
Repeat this step twice
Resuspend the pelleted blood cells in 100 µl PBS containing 0.001% (v/v) Heparin



Handling and Storage

Antibodies are supplied either as 100 tests per vial (1 ml) for singles or 50 tests per vial (1 ml) for dual and triple combinations. They are supplied in 0.01 M sodium phosphate, 0.15 M NaCl; pH 7.3, 0.2% BSA, 0.09% sodiumazide (NaN_3). Store the vials at 2-8 °C. Monoclonal antibodies should be protected from prolonged exposure to light. Reagents are stable for the period shown on the vial label when stored properly.








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Characterization

To ensure consistently high-quality reagents, each batch of monoclonal antibody is tested for conformance with characteristics of a standard reagent. Representative flow cytometric data is included in this data sheet.

Warning All products contain sodiumazide. This chemical is poisonous and hazardous. Handling should be done by trained staff only.

Explanation of used symbols

	Consult instructions for use
	Catalogue number
	Sufficient for
	In Vitro Diagnostic medical device
	Caution, consult accompanying document
	Keep away from (sun)light
	Biological risks
	Temperature limitation (°C)
	For Research Use Only
	Batch code
	Use by yyyy-mm-dd
	Manufacturer
	Authorized Representative in the European Community
	Conformité Européenne (European Conformity)

		Label - tandem	Ex -max (nm)	Em -max (nm)
P	PURE	purified material	-	-
F	FITC	FITC	488	519
R	R-PE	PE	488, 532	578
C	CyQ	PE-Cy5.18	488, 532	667
A	APC		595, 633, 635, 647	660
PC	PerCP		488, 532	678
PCC	PerCP-Cy5.5		488, 532	695



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