PRODUCT INFORMATION SHEET

Monoclonal antibodies detecting human antigens

CD8 FITC **CD25** R-PE RUO REF IQP-213FR 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.

CD8

Clone MCD8

Isotype Murine IgG1

Specificity Clone MCD8 produces mouse IgG1 immunoglobulins directed against human CD8, and has been clustered as CD8 during the 6th Leukocyte Typing Workshop. The CD8 molecule is expressed as a heterodimer of CD8a (32-34 kD) and CD8b (32-34 kD) glycoproteins.
CD8 positive T cells, known as cytotoxic/suppressor cells (Tcyt), may be further sub-divided into Tc1 and Tc2 cells based on cytokine profile and functional activity. CD8 antibodies may help identify T cell leukemias, such as common T-ALL or mature T-ALL in combination with other markers such as TdT and cyCD3. It is also used to distinguish between chronic B and T cell lymphoid leukemias.

HLDA Workshop
6th International Workshop on Human Leukocyte Differentiation Antigens.

CD25

Clone B-B10

Isotype Murine IgG1

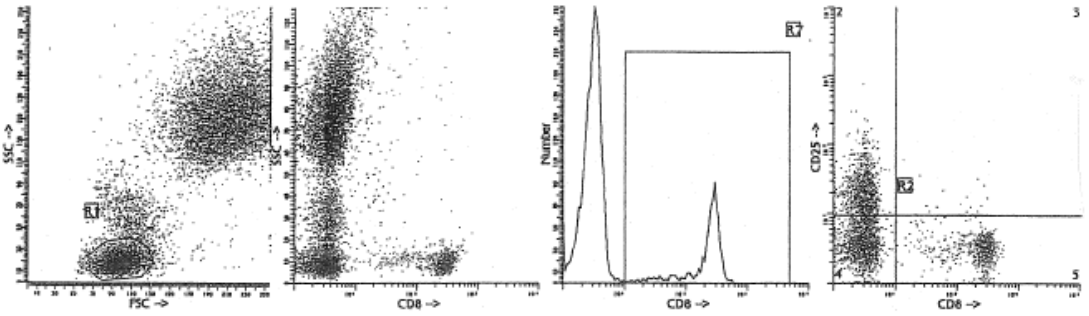
Specificity CD25 antibodies are used as a marker of cell activation in transplantation patients, for the detection of cells infected with the human T cell Leukemia viruses I and II, and for the immunophenotyping of lymphomas and leukemias. The IL-2 receptor is strongly expressed in Hodgkin's disease, 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/CD123 heterodimer binds IL-2 with intermediate affinity. Both the high and intermediate-affinity receptors are important for IL-2 signaling.

HLDA Workshop
4th International Workshop on Human Leukocyte Differentiation Antigens.

Usage All these reagents are effectively formulated for direct immunofluorescent staining of human tissue for flow cytometric analysis using 10 µl/10^6 leukocytes for singles and 20 µl/10^6 leukocytes in case of dual and triple combinations. Since applications vary, each investigator should titrate the reagent to obtain optimal results.

Representative Data
Staining with the dual CD8 FITC and CD25 R-PE was performed using 20 µl of the conjugated monoclonal antibody preparation and 100 µl of 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.

Warranty Products sold hereunder are warranted only to conform to the quantity and contents stated on the label at the time of delivery to the customer. There are no warranties, expressed or implied, which extend beyond the description on the label of the product. IQ Products is not liable for property damage, personal injury, or economic loss caused by the product.

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
REF	Catalogue number
	Sufficient for
IVD	In Vitro Diagnostic medical device
	Caution, consult accompanying document
	Keep away from (sun)light
	Biological risks
	Temperature limitation (°C)
RUO	For Research Use Only
LOT	Batch code
	Use by yyyy-mm-dd
	Manufacturer
EC REP	Authorized Representative in the European Community
CE	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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