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PRODUCT INFORMATION SHEET Monoclonal antibodies detecting human antigens									
CD2	FIT	C CD19	R-PE	RUO	REF	IQP-252FR	50 tests		
RUO	For R	esearch Use Only							
<u>[i]</u>		Description							
	CD2 reacts with all human peripheral T-lymphocytes and a fraction of the NK cell (large granular lymphocyte) population. It reacts specifically with a 45-50 kD single chain transmembrane glycoprotein, also known as the LFA-2, or the sheep erythrocyte receptor. CD2 antigen plays a role in T cell signaling and in lymphocyte adhesion. The major ligand for the extracellular portior of human CD2 is CD58 (LFA3). Detection of CD19 expressing cells is important in the diagnosis of leukemic precursor B cells (pre-B ALL), mature B cells (B ALL), and plasma cells. Distinction between subtypes of these (acute) leukemias can be performed using CD19 antibodies together with monoclonal antibodies to cytoplasmic or membrane Ig. A large number of B cell disorders can be effectively characterized by expression of CD19 and one or more additional antigens. One example is hairy cell leukemia (HCL), which shows specific expression of CD11c, CD19, CD20 and CD103. The combination for distinction between different leukemias using CD19 antibodies are CD5/CD19 (e.g. Chronic Lymphatic Leukemia); CD10/CD19 (common ALL and pre-B ALL) and CD19/cyCD79a (Acute B cell leukemia).								
		CD2							
Clone		B-E2							
Isotype		Murine IgG2b							
Specific	ity	Clone B-E2 pro transmembran				cific for a 45-50	kD single chain		
HLDA W	orksh	ор							
			I Workshop	on Human Le	ukocyte Differen	tiation Antigens			
		CD19							
Clone		HD37			ro	<u> </u>	Ctc.		
Isotype		Murine IgG1							
Specifici	ity	Monoclonal antibodies clustered as CD19 detect all peripheral blood B cells. In addition, CD19 is expressed on precursor B cells during maturation, but not on mature plasma cells. The function of the CD19 molecule is related to signal transfer and is involved in regulation of B cell proliferation. CD19 is considered to be a characteristic B cell marker and therefore commonly used in routine immunophenotyping. CD19 may also be expressed on follicular dendritic cells.							
HLDA W	orksh		ıl Workshor	o on Human Le	ukocyte Differen	tiation Antigens			
Usage		4 th International Workshop on Human Leukocyte Differentiation Antigens 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 cas of dual and triple combinations. Since applications vary, each investigator should titrate the reager to obtain optimal results.							
Represe	entativ	IQP-252FR (CD					mple from a healthy .00 μl blood sample.		
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Immunofluorescence staining and lysing protocol

Flow cytometry method for use with dual and triple combinations

- 1. Add 100 μ I of EDTA-treated blood (i.e. approx. 10⁶ 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 $1000 \times g$.
- 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

<u>∧ &/ * </u>

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 (NaN₃). 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

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<u>I</u> I	Consult instructions for use					
REF	Catalogue number					
$\overline{\mathbb{V}}$	Sufficient for					
IVD	In Vitro Diagnostic medical device					
\wedge	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)
Р	PURE	purified material	-	-
F	FITC	FITC	488	519
R	R-PE	PE	488, 532	578
С	CyQ	PE-Cy5.18	488, 532	667
Α	APC	·	595, 633, 635, 647	660
PC	PerCP		488, 532	678
PCC	PerCP-Cy5.5		488, 532	695

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4

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