

FITC anti-human CD45RA

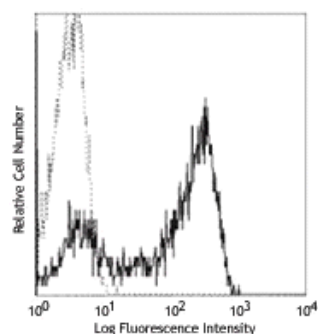
Analyte Specific Reagent. Analytical and performance characteristics are not established.

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|------------------------|--|
| Catalog# / Size | 983002 / 500 µL |
| Clone | HI100 |
| Workshop | IV N906 |
| Isotype | Mouse IgG2b, κ |
| Description | CD45RA is a 205-220 kD single chain type I glycoprotein. It is an exon 4 splice variant of the tyrosine phosphatase CD45. The CD45RA isoform is expressed on resting/naïve T cells, medullary thymocytes, B cells and monocytes. CD45RA enhances both T cell receptor and B cell receptor signaling. CD45 non-covalently associates with lymphocyte phosphatase-associated phosphoprotein (LPAP) on T and B lymphocytes. CD45 has been reported to be associated with several other cell surface antigens including CD1, CD2, CD3, and CD4. CD45 has also been reported to bind galectin-1. CD45 isoform expression can change in response to cytokines. |

Product Details








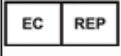


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|-------------------------------|---|
| Reactivity | Human |
| Formulation | Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA). |
| Preparation | The antibody was purified by affinity chromatography, and conjugated with FITC under optimal conditions. |
| Concentration | 200 µg/mL |
| Storage & Handling | The antibody solution should be stored undiluted between 2°C and 8°C, and protected from prolonged exposure to light. Do not freeze. |
| Application | Suggested for Flow Cytometry |
| Disclaimer | <p>WARNINGS AND PRECAUTIONS</p> <ol style="list-style-type: none"> 1. Use appropriate personal protective equipment and safety practices per universal precautions when working with this reagent. Refer to the reagent safety data sheet. 2. This antibody contains sodium azide. Follow federal, state and local regulations to dispose of this reagent. Sodium azide build-up in metal wastepipes may lead to explosive conditions; if disposing of reagent down wastepipes, flush with water after disposal. 3. All specimens, samples and any material coming in contact with them should be considered potentially infectious and should be disposed of with proper precautions and in accordance with federal, state and local regulations. 4. Do not use this reagent beyond the expiration date stated on the label. 5. Do not use this reagent if it appears cloudy or if there is any change in the appearance of the reagent as these may be an indication of possible deterioration. 6. Avoid prolonged exposure of the reagent or stained cells to light. |

Product Data



Typical results from human peripheral blood lymphocytes stained either with HI100 FITC used at 5 µL/test (solid line) or with an isotype control (dotted line).

Symbols Glossary*

| Symbol | Meaning | Symbol Title | Symbol No. | Symbol | Meaning | Symbol Title | Symbol No. |
|---|---|----------------------------|------------|---|--|---|------------|
|  | Catalog number | Catalogue number | 5.1.6 |  | Indicates the need for the user to consult the instructions for use. | Consult instructions for use | 5.4.3 |
|  | Indicates the temperature limits to which the medical device can be safely exposed. | Temperature limit | 5.3.7 |  | Indicates a medical device that needs protection from light sources. | Keep away from sunlight | 5.3.2 |
|  | Indicates the upper limit of temperature to which the medical device can be safely exposed. | Upper limit of temperature | 5.3.6 |  | Indicates the date after which the medical device is not to be used. | Use-by date | 5.1.4 |
|  | Indicates the medical device manufacturer. | Manufacturer | 5.1.1 |  | Indicates the authorized representative in the European Community. | Authorized representative in the European Community | 5.1.2 |
|  | Indicates the manufacturer's batch code so that the batch or lot can be identified. | Batch code | 5.1.5 |  | Indicates a medical device that is intended to be used as an in vitro diagnostic medical device. | <i>In vitro</i> diagnostic medical device | 5.5.1 |

* Symbol information is from EN ISO 15223-1:2016 Medical devices – Symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General requirements

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