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## 30-2546: Anti-Human CD8 APC (Clone: LT8)

Clonality: Monoclonal

**Clone Name:** LT8 Application: **FACS** Reactivity: Human Conjugate: APC Gene: CD8A Gene ID: 925 **Alternative Name:** p32, LEU2 Isotype: Mouse IgG1

Immunogen Information: human blood lymphocytes

## **Description**

The CD8 T cell coreceptor (monomer approx. 32-34 kDa) is expressed as alpha/beta heterodimer on majority of MHC I-restricted conventional T cells and thymocytes and as alpha/alpha homodimer on subsets of memory T cells, intraepithelial lymphocytes, NK cells and dendritic cells. Regulation of CD8 beta level on T cell surface seems to be an important mechanism to control their effector function. Assembly of CD8 alpha-beta but not alpha-alpha dimers is connected with formation or localization to the lipid rafts. Recruiting triggered TCR complexes to these membrane microdomains as well as affinity of TCR to MHC I is modulated by CD8, thereby affecting the functional diversity of the TCR signaling.

Specificity: The mouse monoclonal antibody LT8 recognizes an extracellular epitope of CD8, a cell surface glycoprotein found on most cytotoxic T lymphocytes that mediates efficient cell-cell interactions within the immune system. This antibody blocks Leu2 binding.

## **Product Info**

Amount: 100 tests

**Purification :** The purified antibody is conjugated with allophycocyanin (APC) under optimum conditions. The

conjugate is purified by size-exclusion chromatography.

**Content:** Formulation: Stabilizing phosphate buffered saline (PBS) solution containing 15 mM sodium

azide

**Storage condition:** Store in the dark at 2-8°C. Do not freeze. Avoid prolonged exposure to light.

## **Application Note**

Flow cytometry: The reagent is designed for analysis of human blood cells using 10  $\tilde{A} \square \hat{A} \mu l$  reagent / 100  $\tilde{A} \square \hat{A} \mu l$  of whole blood or 10<sup>6</sup> cells in a suspension. The content of a vial (1 ml) is sufficient for 100 tests.

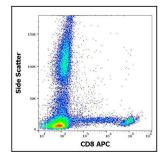


Figure 1 : Flow cytometry surface staining pattern of human peripheral whole blood stained using anti-human CD8 (LT8) APC antibody (4  $\mu$ l reagent / 100  $\mu$ l of peripheral whole blood)



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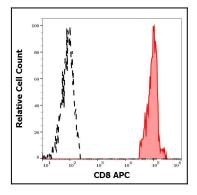


Figure 2 : Separation of human CD8 positive lymphocytes (red-filled) from CD8 negative lymphocytes (black-dashed) in flow cytometry analysis (surface staining) of human peripheral whole blood stained using anti-human CD8 (LT8) APC antibody (4  $\mu l$  reagent / 100  $\mu l$  of peripheral whole blood).