

## Technical Data Sheet

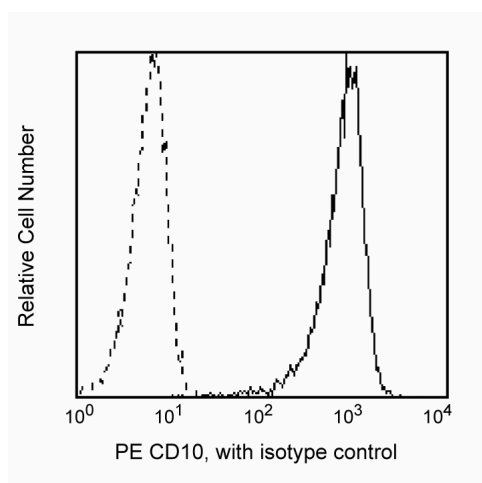
## PE Mouse Anti-Human CD10

## Product Information

|                         |  |
|-------------------------|--|
| <b>Material Number:</b> | 555375   |
| <b>Alternate Name:</b>  | MME; CALLA; EPN; NEP; neprilysin; SFE; atriopeptidase; enkephalinase               |
| <b>Size:</b>            | 100 Tests  |
| <b>Vol. per Test:</b>   | 20 µl  |
| <b>Clone:</b>           | HI10a  |
| <b>Immunogen:</b>       | Acute CALLA Leukemia Blast Cells   |
| <b>Isotype:</b>         | Mouse (BALB/c) IgG1, κ   |
| <b>Reactivity:</b>      | QC Testing: Human<br>Tested in Development: Rhesus, Cynomolgus, Baboon<br>V CD10.7 |
| <b>Workshop:</b>        | V CD10.7   |
| <b>Storage Buffer:</b>  | Aqueous buffered solution containing BSA and ≤0.09% sodium azide.                  |

## Description

The HI10a monoclonal antibody specifically binds to CD10 which is also known as Neutral endopeptidase (NEP), Enkephalinase, Atriopeptidase, and Neprilysin. CD10 is encoded by *MME* (membrane metallo-endopeptidase). CD10 is a 100 kDa type II transmembrane glycoprotein that has neutral endopeptidase activity and is otherwise known as the Common Acute Lymphoblastic Leukemia Antigen (CALLA). CD10 is expressed on a wide variety of normal and neoplastic cell types. Normal cells expressing CD10 include granulocytes, bone marrow stromal cells, a subset of B-cell progenitors, germinal center B cells and fibroblasts. This cell surface metalloendopeptidase inactivates a number of signaling molecules and serves as a major regulator in the nervous, immune and other systems.



**Flow cytometric analysis of CD10 expression on REH cell line.** REH cells were stained with either PE Mouse Anti-Human CD10 (Cat. No. 555375/561002; solid line histogram) or PE Mouse IgG1, κ Isotype Control (Cat. No. 555749; dashed line histogram). Fluorescence histograms depicting CD10 (or Ig isotype control) expression were derived from gated events with the side and forward light-scattering characteristics of viable cells. Flow cytometry was performed on a BD FACScan™ system.

## Preparation and Storage

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

The antibody was conjugated with R-PE under optimum conditions, and unconjugated antibody and free PE were removed.

## Application Notes

## Application

|                |                  |
|----------------|------------------|
| Flow cytometry | Routinely Tested |
|----------------|------------------|

## Suggested Companion Products

| Catalog Number | Name                             | Size      | Clone   |
|----------------|----------------------------------|-----------|---------|
| 555749         | PE Mouse IgG1, κ Isotype Control | 100 Tests | MOPC-21 |
| 561002         | PE Mouse Anti-Human CD10         | 25 Tests  | HI10a   |
| 554656         | Stain Buffer (FBS)               | 500 mL    | (none)  |
| 554657         | Stain Buffer (BSA)               | 500 mL    | (none)  |

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555375 Rev. 11



## Product Notices

1. This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use  $1 \times 10^6$  cells in a 100- $\mu$ l experimental sample (a test).
2. An isotype control should be used at the same concentration as the antibody of interest.
3. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
4. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
5. For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at [www.bdbiosciences.com/colors](http://www.bdbiosciences.com/colors).
6. Species testing during development may have been performed with a different format of the same clone. Selected applications have been tested for cross-reactivity.
7. Please refer to [www.bdbiosciences.com/pharming/protocols](http://www.bdbiosciences.com/pharming/protocols) for technical protocols.

## References

- Barclay NA, Brown MH, Birkeland ML, et al, ed. *The Leukocyte Antigen FactsBook*. San Diego, CA: Academic Press; 1997(Biology)
- Knapp W. W. Knapp .. et al., ed. *Leucocyte typing IV : white cell differentiation antigens*. Oxford New York: Oxford University Press; 1989:1-1182. (Biology)
- Letarte M, Vera S, Tran R, et al. Common acute lymphocytic leukemia antigen is identical to neutral endopeptidase. *J Exp Med*. 1988; 168(4):1247-1253. (Biology)