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Datasheet for ABIN2144866  
**anti-CD37 antibody (PE)**

1 Image

### Overview

|              |  |
|--------------|--|
| Quantity:    | 50 tests   |
| Target:      | CD37 (TSPAN26)   |
| Reactivity:  | Human  |
| Host:        | Mouse  |
| Clonality:   | Monoclonal   |
| Conjugate:   | This CD37 antibody is conjugated to PE   |
| Application: | Flow Cytometry (FACS), ELISA, Immunofluorescence (IF), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)) |

### Product Details

|                  |   |
|------------------|---|
| Clone:           | 4AIPO-24  |
| Isotype:         | IgG2b   |
| Characteristics: | 4AIPO-24 reacts with CD37 (a.k.a. gp52-40 ), a 40-52 kDa molecule, which is strongly expressed on B cells from the pre-B cell stage, but not on plasma cells. It is also present at low levels on some T cells, monocytes and granulocytes. CD37 is a stable marker for malignancies derived from mature B cells, such as B-CLL, HCL and all types of B-NHL. CD37 is involved in signal transduction. CD37 is stable marker for malignancies derived from mature B cells: B-CLL. HCL and B-NHL. |

### Target Details

|         |                |
|---------|----------------|
| Target: | CD37 (TSPAN26) |
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## Target Details

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Alternative Name: CD37 ([TSPAN26 Products](#))

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Pathways: [Positive Regulation of Immune Effector Process, Production of Molecular Mediator of Immune Response](#)

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## Application Details

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Application Notes: Optimal working dilution should be determined by the investigator.

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Assay Procedure:

- Take 100  $\mu$ L peripheral blood anticoagulated by EDTA and add to the bottom of 5 mL tube,
- Add 10  $\mu$ L labeled antibody to the bottom of flow tube mixing with the whole blood, incubate for 20 minutes at room temperature away from light,
- Add 2 mL  $1\times$ RBC lysis buffer, incubate for 10 minutes away from light after mixing, dissolve red blood cells (recommended: RBC lysing Solution  $10\times$ , Cat.: FXP001),
- Sample tube is set to 1000 rpm centrifugation for 5 minutes, discard the supernatant,
- Add 2 mL PBS wash buffer to resuspend the cells, then 1000 rpm centrifugation for 5 minutes, discard the supernatant,
- Add 0.5 mL PBS wash buffer to resuspend the cells and detect by flow cytometry (sample should be determined on the day on the machine and can also be added fixation overnight at 4 °C then measured).
- [PBS wash buffer: PBS +1 % FBS +0.1 % NaN<sub>3</sub>, Cat.: FXP005]
- [Cell fixation: 2 % formaldehyde solution]

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Restrictions: For Research Use only

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## Handling

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Format: Liquid

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Buffer: Phosphate-buffered solution, pH 7.4, containing and 0.2 % (w/v) BSA

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Preservative: Sodium azide

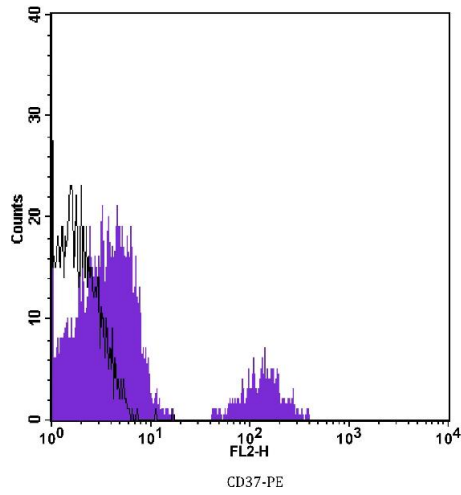
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Precaution of Use: This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

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Storage: 4 °C

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### Flow Cytometry

Image 1.