

Monoclonal Anti-human/mouse LAP (TGF- β 1)-PerCP

Catalog Number: FAB2463C

Lot Number: AATI03

100 Tests

Reagents Provided

Peridinin-Chlorophyll-Protein-Complex (PerCP)-conjugated mouse monoclonal anti-human/mouse LAP (TGF- β 1): Supplied as 25 μ g of antibody in 1 mL saline containing up to 0.5% BSA and 0.1% sodium azide.

Clone #: 27232

Isotype: mouse IgG₁

Reagents Not Provided

- Flow Cytometry Staining Buffer (Catalog # FC001) or other BSA-supplemented saline buffer.

Storage

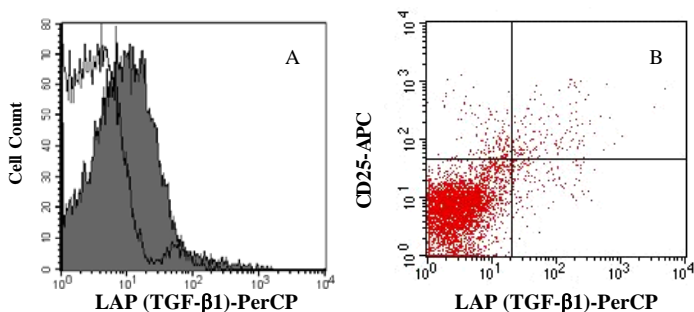
Reagents are stable for **twelve months** from date of receipt when stored in the dark at 2° - 8° C.

Intended Use

Designed to quantitatively determine the percentage of cells bearing LAP (TGF- β 1) within a population and qualitatively determine the density of LAP (TGF- β 1) on cell surfaces by flow cytometry.

Product Description

This antibody was produced from a hybridoma resulting from the fusion of a mouse myeloma with B cells obtained from a mouse immunized with Sf21-derived recombinant human latency associated peptide of TGF- β 1 (rhLAP (TGF- β 1); aa 30 - 278; Accession# P01137.2). The IgG fraction of the tissue culture supernatant was purified by Protein G affinity chromatography. The purified antibody was then conjugated to PerCP fluorochrome. Cell surface expression of LAP (TGF- β 1) is determined by flow cytometry. PerCP has a maximum absorption of 482 nm and 564 nm and a maximum emission of 675 nm.



(A) Human platelets or (B) mouse splenocytes were stained with PerCP-conjugated anti-human/mouse LAP (TGF- β 1) (Catalog # FAB2463C, filled histogram) or isotype control (Catalog # IC002C, open histogram). Splenocytes were costained with APC-conjugated anti-mouse CD25 (Catalog # FAB2438A), and quad marker was set based on isotype control staining.

Background Information

TGF- β is secreted by cells in the form of an inactive complex. This complex consists of TGF- β associated non-covalently with a protein designated the latency associated peptide (LAP). TGF- β and LAP represent components of a pro-peptide that is cleaved in a post-golgi compartment prior to secretion. LAP and TGF- β each consist of a disulfide-linked homodimer and the association of these two components renders TGF- β inactive and inaccessible to anti-TGF- β antibodies.

Flow Cytometry Validation

This antibody has been tested for flow cytometry using human platelets and mouse splenocytes.

- Cells may be Fc-blocked with 1 μ g of human or mouse IgG/ 10^5 cells for 15 minutes at room temperature. Do not wash excess blocking IgG from this reaction.
- After blocking, 10 μ L of conjugated antibody was added to 1 - 5 x 10^5 cells and incubated for 30 minutes at room temperature.
- Unbound antibody was removed by washing the cells twice in Flow Cytometry Staining Buffer (Catalog # FC001). Note that whole blood requires a RBC lysis step at this point using Flow Cytometry Human (Catalog# FC002) or Mouse (Catalog# FC003) Lyse Buffer.
- The cells were resuspended in Flow Cytometry Staining Buffer for final flow cytometric analysis. As a control for analysis, cells in a separate tube should be treated with PerCP-labeled mouse IgG₁ antibody. This procedure may need to be modified, depending upon cell type and final utilization. Individual users may need to titrate to determine optimal reagent amount for their specific use.

Warning: Contains sodium azide as a preservative - sodium azide may react with lead and copper plumbing to form explosive metal azides. Flush with large volumes of water during disposal.

FOR RESEARCH USE ONLY. NOT FOR USE IN HUMANS.

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