**Technical Data Sheet**

**APC-Cy™7 Rat Anti-Mouse CD3 Molecular Complex**

**Product Information**

**Material Number:** 560590

**Alternate Name:** CD3; CD3 epsilon; Cd3e; CD3ε; T3e

**Size:** 50 µg

**Concentration:** 0.2 mg/ml

**Clone:** 17A2

**Immunogen:** γδ TCR-positive T-T hybridoma D1

**Isotype:** Rat (SD) IgG2b, κ

**Reactivity:** Mouse

**Storage Buffer:** Aqueous buffered solution containing protein stabilizer and ≤0.09% sodium azide.

**Description**

The 17A2 monoclonal antibody specifically binds to the T-cell receptor-associated CD3 complex that is expressed on many thymocytes and mature T lymphocytes. Plate-bound 17A2 antibody has been reported to induce IL-2 production by cultured T cells in the absence of accessory cells. The binding of the 17A2 antibody to T cells can be blocked by the anti-CD3ε mAb 145-2C11. This suggests that the 17A2 antibody recognizes an epitope of the CD3 epsilon chain. In vivo treatment with 17A2 antibody has been reported to partially deplete T lymphocytes and temporarily down-modulates CD3 expression on T cells.

**Flow cytometric analysis of CD3 on mouse splenocytes.** Splenic leukocytes were preincubated with Purified Rat Anti-Mouse CD16/CD32 antibody (Mouse BD Fc Block™) (Cat. No. 553141/553142). **Left Panel:** Splenocytes from C57BL/6 mice were stained with either a APC-Cy™7 Rat IgG2b, κ isotype control (shaded; Cat. No. 552773) or with the APC-Cy7 Rat Anti-Mouse CD3 antibody (unshaded; Cat. No 560590). **Middle and Right Panels:** Splenocytes from C57BL/6 mice were stained with a FITC Rat Anti-Mouse CD19 antibody (Cat.No. 553785/557398/561740) in conjunction with either a APC-Cy7 Rat IgG2b, κ isotype control (middle panel) or the APC-Cy7 Rat Anti-Mouse CD3 antibody (right panel). Histograms and dot plots were derived from gated events based on light scattering characteristics for splenocytes. Flow cytometry was performed on a BD™ LSR II flow cytometry 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.

**Application Notes**

**Application**

Flow cytometry Routinely Tested

**Suggested Companion Products**

<table>
<thead>
<tr>
<th>Catalog Number</th>
<th>Name</th>
<th>Size</th>
<th>Clone</th>
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<tr>
<td>552773</td>
<td>APC-Cy™7 Rat IgG2b κ Isotype Control</td>
<td>0.1 mg</td>
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<td>553785</td>
<td>FITC Rat Anti-Mouse CD19</td>
<td>0.5 mg</td>
<td>ID3</td>
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<td>Purified Rat Anti-Mouse CD16/CD32 (Mouse BD Fc Block™)</td>
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</tbody>
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**BD Biosciences**

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Product Notices

1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
3. Warning: Some APC-Cy7 and PE-Cy7 conjugates show changes in their emission spectrum with prolonged exposure to formaldehyde. If you are unable to analyze fixed samples within four hours, we recommend that you use BD™ Stabilizing Fixative (Cat. No. 338036).
4. Please observe the following precautions: Absorption of visible light can significantly alter the energy transfer occurring in any tandem fluorochrome conjugate; therefore, we recommend that special precautions be taken (such as wrapping vials, tubes, or racks in aluminum foil) to prevent exposure of conjugated reagents, including cells stained with those reagents, to room illumination.
5. APC-Cy7 tandem fluorochrome emission is collected in a detector for fluorescence wavelengths of 750 nm and higher.
6. Cy is a trademark of Amersham Biosciences Limited.
7. APC-Cy7 is a tandem fluorochrome composed of Allophycocyanin (APC), which is excited by laser lines between 595 and 647 nm and serves as an energy donor, coupled to the cyanine dye Cy7™, which acts as an energy acceptor and fluoresces at 780 nm. BD Biosciences Pharmingen has maximized the fluorochrome energy transfer in APC-Cy7, thus maximizing its fluorescence emission intensity, minimizing residual emission from APC, and minimizing required electronic compensation in multilaser-laser flow cytometry systems. Note: Although every effort is made to minimize the lot-to-lot variation in residual emission from APC, it is strongly recommended that every lot be tested for differences in the amount of compensation required and that individual compensation controls are run for each APC-Cy7 conjugate.
8. 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.
9. For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at www.bdbiosciences.com/colors.
10. An isotype control should be used at the same concentration as the antibody of interest.

References