Technical Data Sheet

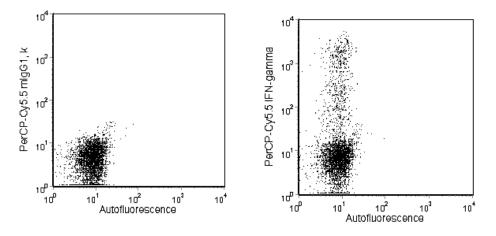
PerCP-Cy™5.5 Mouse Anti-Human IFN-γ

Product Information

| Material Number: | 560742 | |
|------------------|---|--|
| Size: | 50 tests | |
| Vol. per Test: | 5 μl | |
| Clone: | 4S.B3 | |
| Immunogen: | Partially purified human IFN-y from supernatants of human PBMC stimulated | |
| | with Staphylococcus aureus | |
| Isotype: | Mouse IgG1, κ | |
| Reactivity: | Human | |
| | QC Testing: Rhesus or Baboon or Cynomolgus | |
| Storage Buffer: | Aqueous buffered solution containing BSA and $\leq 0.09\%$ sodium azide. | |

Description

The 4S.B3 antibody reacts with human interferon- γ (IFN- γ). The immunogen used to generate this hybridoma was partially purified human IFN- γ obtained from supernatants of human PBMC stimulated with *Staphylococcus aureus*.



Flow cvtometric analysis for IFN-v in stimulated Rhesus macaque peripheral blood mononuclear cells (PBMC). PBMC from Rhesus macaque were stimulated for 6 hours with 50 ng/mL PMA (Sigma-Aldrich Cat. No. P-8139) and 500 ng/mL calcium ionophore A23187 (Sigma-Aldrich Cat. No. C-9275) in the presence of BD GolgiStop™ (Cat. No. 554724). Cells were then fixed and permeabilized using BD Cytofix/Cytoperm™ (Cat. No. 554714) followed by staining with either a PerCP-Cy™5.5 Mouse IgG1, κ isotype control (left panel) or with the PerCP-Cy™5.5 Mouse Anti-Human IFN-γ antibody (right panel). Dot plots were derived from gated events based on light scattering characteristics for lymphocytes. Flow cytometry was performed on a BD™ LSR II flow cytometry system.

(none)

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 PerCP-Cy5.5 under optimum conditions, and unconjugated antibody and free PerCP-Cy5.5 were removed. Storage of PerCP-Cy5.5 conjugates in unoptimized diluent is not recommended and may result in loss of signal intensity.

Application Notes

| Application | | | | | | | |
|---|---|---------------|---------|--|--|--|--|
| Intracellular staining (flow cytometry) | | tinely Tested | | | | | |
| Suggested Companion Products | | | | | | | |
| Catalog Number | Name | Size | Clone | | | | |
| 552834 | PerCP-Cy [™] 5.5 Mouse IgG1 κ Isotype Control | 50 tests | MOPC-21 | | | | |
| 550795 | PerCP-Cy [™] 5.5 Mouse IgG1 κ Isotype Control | 0.1 mg | MOPC-21 | | | | |
| 555899 | Lysing Buffer | 100 ml | (none) | | | | |
| 554714 | BD Cytofix/Cytoperm [™] Fixation/Permeablization Kit | 250 tests | (none) | | | | |

0.7 ml

Product Notices

554724

- 1. This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use 1×10^{6} cells in a 100-µl experimental sample (a test).
- 2. An isotype control should be used at the same concentration as the antibody of interest.

Protein Transport Inhibitor (Containing Monensin)

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- 3. 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.
- 4. This PerCP-conjugated product is sold under license to the following patent: US Patent No. 4,876,190.
- 5. Cy is a trademark of Amersham Biosciences Limited. This conjugated product is sold under license to the following patents: US Patent Nos. 5,486,616; 5,569,587; 5,569,766; 5,627,027.
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- 7. PerCP-Cy5.5 is optimized for use with a single argon ion laser emitting 488-nm light. Because of the broad absorption spectrum of the tandem fluorochrome, extra care must be taken when using dual-laser cytometers, which may directly excite both PerCP and Cy5.5[™]. We recommend the use of cross-beam compensation during data acquisition or software compensation during data analysis.
- 8. PerCP-Cy5.5–labelled antibodies can be used with FITC- and R-PE–labelled reagents in single-laser flow cytometers with no significant spectral overlap of PerCP-Cy5.5, FITC, and R-PE fluorescence.
- 9. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- 10. 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.
- 11. For fluorochrome spectra and suitable instrument settings, please refer to our Fluorochrome Web Page at www.bdbiosciences.com/colors.
- 12. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.

References

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