Technical Data Sheet

Rat Activation Compensation Set

Product Information

558512 **Material Number:** 20 tests Size:

51-9004354 **Component:**

APC Mouse anti-Rat CD3 **Description:**

Size: 20 tests (1 ea) 20 µl Vol. per Test: Clone Name:

Mouse (BALB/c) IgM, κ Isotype:

Aqueous buffered solution containing BSA and ≤0.09% sodium azide. Storage Buffer:

Component:

FITC Mouse anti-Rat RT1B **Description:**

20 tests (1 ea) Size: 20 μl Vol. per Test: OX-6 Clone Name:

Mouse (BALB/c) IgG1, κ Isotype:

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

Component:

PE Mouse anti-Rat CD25 (IL-2R α Chain) **Description:**

Size: 20 tests (1 ea) 20 µl Vol. per Test: OX-39 Clone Name:

Mouse (BALB/c) IgG1, κ Isotype:

Aqueous buffered solution containing BSA and ≤0.09% sodium azide. Storage Buffer:

Description

Description Clone Clone 1F4 APC Mouse anti-Rat CD3 Clone OX-6 FITC Mouse anti-Rat RT1B Clone OX-39 PE Mouse anti-Rat CD25

The Rat Activation Compensation Set includes each specificity contained in the Rat Activated T Lymphocyte Cocktail (Cat. no. 558494) bottled separately, each in a 20 test size. Compensation is the process by which the effect of spectral overlap among flurochromes can be corrected minimizing fluorescence signal spillover in neighboring channels. Compensation values are entered into software programs allowing for the adjustment among the overlapping flurochrome emissions. These values can be achieved using single color controls, and verified using fluorescence-minus-one controls, or ad-mixtures of single color controls. Use this set to properly adjust your FITC/PE/APC compensation matrix for the Rat Activated T Lymphocyte Cocktail.

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 to APC under optimum conditions, and unconjugated antibody and free APC were removed.

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

The antibody was conjugated with FITC under optimum conditions, and unreacted FITC was removed.

Application Notes

Application

Flow cytometry Routinely Tested

Suggested Companion Products

BD Biosciences

bdbiosciences.com

United States Asia Pacific Canada Latin America/Caribbean Europe Japan 877.232.8995 888.259.0187 32.53.720.550 0120.8555.90 65.6861.0633

For country-specific contact information, visit bdbiosciences.com/how_to_order/

Conditions: The information disclosed herein is not to be construed as a recommendation to use the above product in violation of any patents. BD Biosciences will not be held responsible for patent infringement or other violations that may occur with the use of our products. Purchase does not include or carry any right to resell or transfer this product either as a stand-alone product or as a component of another product. Any use of this product other than the permitted use without the express written authorization of Becton Dickinson and Company is strictly prohibited. For Research Use Only. Not for use in diagnostic or therapeutic procedures. Not for resale. BD, BD Logo and all other trademarks are the property of Becton, Dickinson and Company. ©2008 BD



558512 Rev. 1

Catalog Number	Name	Size	Clone
558494	Rat Activated T Lymphocyte Cocktail	50 tests	(none)

Product Notices

- 1. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- 2. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- 3. 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.

References

Shapiro, HM. Practical Flow Cytometry. New York: Wiley-Liss, Inc; 1995:18-19.(Biology)

558512 Rev. 1 Page 2 of 2