

BD CD8 (SK1)

Monoclonal Antibodies Detecting Human Antigens

Form	Catalog number	Form	Catalog number
Pure	346310	APC	340584
FITC	347313	APC-Cy7	348793
PerCP	347314	APC-H7	641400
PerCP-Cy5.5	341051	AmCyan	339188
PE-Cy7	335787	V500-C	647457

Product availability varies by region. Contact BD Biosciences Customer Support or your local sales representative for information.

Research Applications

Research applications include studies of:

- Enumeration and monitoring of the T-suppressor/cytotoxic subset in peripheral blood¹⁻³
- Analysis of cell-mediated cytotoxicity⁴
- Analysis of immunoregulation and T-lymphocyte-mediated suppression^{5,6}
- Natural killer (NK)-lymphocyte subsets⁷
- Immune function in cancer research⁸
- T-cell response to cytomegalovirus⁹⁻¹¹

Description

Specificity

The CD8 antibody recognizes the 32-kilodalton (kDa) α -subunit of a disulfide-linked bimolecular complex.^{12,13} The majority of peripheral blood CD8⁺ T lymphocytes express an α/β heterodimer (32, 30 kDa), while CD8⁺CD16⁺ NK lymphocytes and CD8⁺ T-cell receptor (TCR)- γ/δ ⁺ T lymphocytes express an α/α homodimer (30 kDa). CD8⁺TCR- α/β ⁺ T lymphocytes can express either an α/α homodimer or α/β heterodimer.^{12,13} The CD8 antigenic determinant binds to class I major histocompatibility complex (MHC) molecules, resulting in increased adhesion between the CD8⁺ T lymphocytes and target cells.¹⁴⁻¹⁶ Binding of the CD8 antigen to class I MHC molecules enhances the activation of resting T lymphocytes.¹⁴⁻¹⁷ The CD8 antigen is coupled to a protein tyrosine kinase, p56^{lck}. The CD8:p56^{lck} complex can play a role in T-lymphocyte activation through mediation of the interactions between the CD8 antigen and the CD3/TCR complex.^{16,17}

Antigen distribution

The CD8 antigen is present on the human suppressor/cytotoxic T-lymphocyte subset^{2-5,18,19} as well as on a subset of NK lymphocytes.⁶ The CD8 antigen is expressed on 19% to 48% of normal peripheral blood lymphocytes²⁰ and 60% to 85% of normal thymocytes.^{2,4} The CD8⁺ T- and NK-lymphocyte subsets can be further subdivided into the following groups: CD16⁺ NK lymphocytes that can express the CD8 antigen in low density;⁶ CD57⁺ T lymphocytes that express high-density CD8 antigen;⁶ and CD8⁺CD62L⁺ lymphocytes that collaborate with CD8⁺CD62L⁻ lymphocytes to generate suppression of B-lymphocyte function.¹ CD8 cross-reacts with lymphocytes of some nonhuman primate species.²¹

Clone

The CD8 antibody, clone SK1,²² is derived from hybridization of NS-1 mouse myeloma cells with spleen cells from BALB/c mice immunized with human peripheral blood T lymphocytes.

For Research Use Only. Not for use in diagnostic or therapeutic procedures.

Composition

The CD8 antibody is composed of mouse IgG₁ heavy chains and kappa light chains.

Product configuration

The following are supplied in buffer containing a stabilizer and a preservative.

Form	Number of tests	Volume per test (µL)	Amount provided (µg)	Total volume (mL)	Concentration (µg/mL)	Stabilizer	Preservative
Pure	200	20	50	4.0	12.5	Gelatin	0.1% Sodium azide
FITC	100	20	25	2.0	12.5	Gelatin	0.1% Sodium azide
PerCP	100	20	12	2.0	6	Gelatin	0.1% Sodium azide
PerCP-Cy5.5	50	20	5	1.0	5	Gelatin	0.1% Sodium azide
PE-Cy7	100	5	25	0.5	50	Gelatin	0.1% Sodium azide
APC	100	5	25	0.5	50	Gelatin	0.1% Sodium azide
APC-Cy7	100	5	25	0.5	50	Gelatin	0.1% Sodium azide
APC-H7	100	5	25	0.5	50	Gelatin	0.1% Sodium azide
AmCyan	100	5	50	0.5	100	BSA	0.1% Sodium azide
V500-C ^a	100	5	25	0.5	50	BSA	MIT

^a BD Horizon™ V500-C

CAUTION Some APC-Cy7 conjugates, and to a lesser extent PE-Cy7 and APC-H7 conjugates, show changes in their emission spectra with prolonged exposure to paraformaldehyde or light. For overnight storage of stained cells, wash and resuspend in buffer without paraformaldehyde after 1 hour of fixation.

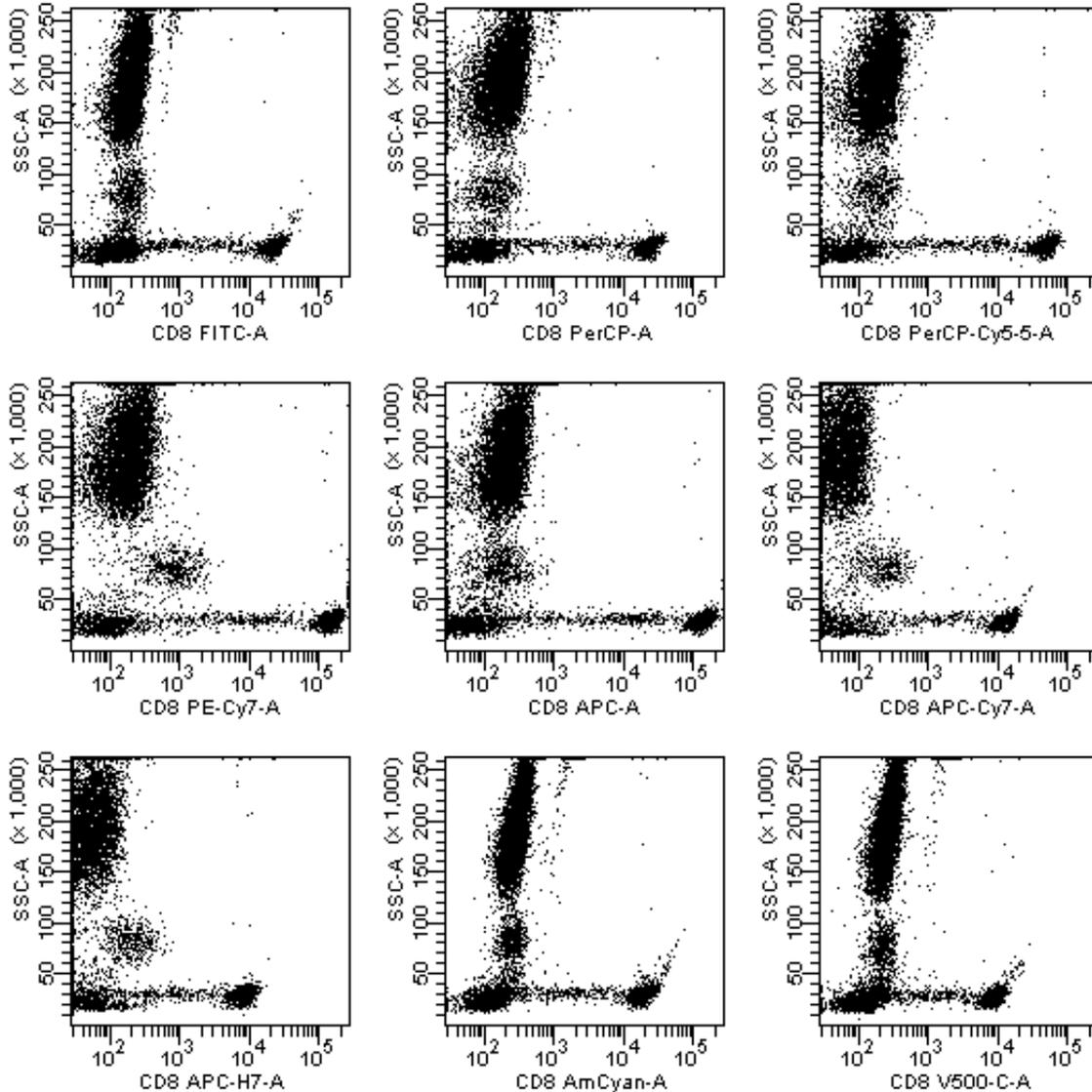
CAUTION Prolonged exposure of cells to paraformaldehyde can lead to increased autofluorescence in the violet channels. For overnight storage of stained cells, wash and resuspend in buffer without paraformaldehyde after 1 hour of fixation.

Procedure

Go to our website (bdbiosciences.com) or contact your local BD representative for the lyse/wash protocol for direct immunofluorescence.

Representative Data

Flow cytometric analysis was performed on whole blood stained with the indicated conjugated antibody. Laser excitation was at 405 nm, 488 nm, or 635 nm. Representative data analyzed with a BD flow cytometer is shown in the following plots.



Handling and Storage

Store vials at 2–8 °C. Conjugated forms should not be frozen. Protect from exposure to light. Each reagent is stable until the expiration date shown on the bottle label when stored as directed.

Warning

All biological specimens and materials coming in contact with them are considered biohazards. Handle as if capable of transmitting infection,^{23,24} and dispose of with proper precautions in accordance with federal, state, and local regulations. Never pipette by mouth. Wear suitable protective clothing, eyewear, and gloves.

The V500-C conjugate contains 0.008% 2-methyl-4-isothiazolin-3-one (MIT), CAS number 2682-20-4. The AmCyan conjugate contains 0.3958% ethylenediamine, ethoxylated and propoxylated, CAS number 26316-40-5. These reagents are classified as hazardous according to the Globally Harmonized System of Classification and Labelling of Chemicals (GHS).

	Warning
	H317: May cause an allergic skin reaction.
Prevention	P261: Avoid breathing dust/fume/gas/mist/vapors/spray. P272: Contaminated work clothing should not be allowed out of the workplace. P280: Wear protective gloves/protective clothing/eye protection/face protection.
Response	P302+P352: IF ON SKIN: Wash with plenty of water. P333+P313: If skin irritation or rash occurs: Get medical advice/attention. P362+P364: Take off contaminated clothing and wash it before reuse.
Disposal	P501: Dispose of contents/container to an approved facility in accordance with local, regional, national and international regulations.

Go to regdocs.bd.com to download the Safety Data Sheet.

Characterization

To ensure consistently high-quality reagents, each lot of antibody is tested for conformance with characteristics of a standard reagent.

Warranty

Unless otherwise indicated in any applicable BD general conditions of sale for non-US customers, the following warranty applies to the purchase of these products.

THE PRODUCTS SOLD HEREUNDER ARE WARRANTED ONLY TO CONFORM TO THE QUANTITY AND CONTENTS STATED ON THE LABEL OR IN THE PRODUCT LABELING AT THE TIME OF DELIVERY TO THE CUSTOMER. BD DISCLAIMS HEREBY ALL OTHER WARRANTIES, EXPRESSED OR IMPLIED, INCLUDING WARRANTIES OF MERCHANTABILITY AND FITNESS FOR ANY PARTICULAR PURPOSE AND NONINFRINGEMENT. BD'S SOLE LIABILITY IS LIMITED TO EITHER REPLACEMENT OF THE PRODUCTS OR REFUND OF THE PURCHASE PRICE. BD IS NOT LIABLE FOR PROPERTY DAMAGE OR ANY INCIDENTAL OR CONSEQUENTIAL DAMAGES, INCLUDING PERSONAL INJURY, OR ECONOMIC LOSS, CAUSED BY THE PRODUCT.

References

1. Gatenby PA, Kansas GS, Xian CY, Evans RL, Engleman EG. Dissection of immunoregulatory subpopulations of T lymphocytes within the helper and suppressor sublineages in man. *J Immunol.* 1982;129:1997-2000.
2. Ledbetter JA, Evans RL, Lipinski M, Cunningham-Rundles C, Good RA, Herzenberg LA. Evolutionary conservation of surface molecules that distinguish T lymphocyte helper/inducer and cytotoxic/suppressor subpopulations in mouse and man. *J Exp Med.* 1981;153:310-323.
3. Ledbetter JA, Frankel AE, Herzenberg LA, Herzenberg LA. Human Leu T-cell differentiation antigens: quantitative expression on normal lymphoid cells and cell lines. In: Hämmerling G, Hämmerling U, Kearney J, eds. *Monoclonal Antibodies and T-Cell Hybridomas: Perspectives and Technical Advances.* New York, NY: Elsevier/North Holland; 1981:16-22.

4. Evans RL, Wall DW, Platsoucas CD, et al. Thymus-dependent membrane antigens in man: inhibition of cell-mediated lympholysis by monoclonal antibodies to the T_{H2} antigen. *Proc Natl Acad Sci USA*. 1981;78:544-548.
5. Kotzin BL, Benike CJ, Engleman EG. Induction of immunoglobulin-secreting cells in the allogeneic mixed leukocyte reaction: regulation by helper and suppressor lymphocyte subsets in man. *J Immunol*. 1981;127:931-935.
6. Lanier LL, Le AM, Phillips JH, Warner NL, Babcock GF. Subpopulations of human natural killer cells defined by expression of the Leu-7 (HNK-1) and Leu-11 (NK-15) antigens. *J Immunol*. 1983;131:1789-1796.
7. Suni MA, Ghanekar SA, Houck DW, et al. CD4⁺CD8^{dim} T lymphocytes exhibit enhanced cytokine expression, proliferation and cytotoxic activity in response to HCMV and HIV-1 antigens. *Eur J Immunol*. 2001;31:2512-2520.
8. Campbell MJ, Scott J, Maecker HT, Park JW, Esserman LJ. Immune dysfunction and micrometastases in women with breast cancer. *Breast Cancer Res Treat*. 2005;91:163-171.
9. Jacobson MA, Maecker HT, Orr PL, et al. Results of a cytomegalovirus (CMV)-specific CD8⁺/interferon- γ ⁺ cytokine flow cytometry assay correlate with clinical evidence of protective immunity in patients with AIDS and CMV retinitis. *J Infect Dis*. 2004;189:1362-1373.
10. Maecker HT, Maino VC. Analyzing T-cell responses to cytomegalovirus by cytokine flow cytometry. *Hum Immunol*. 2004;65:493-499.
11. Tu W, Chen S, Sharp M, et al. Persistent and selective deficiency of CD4⁺ T cell immunity to cytomegalovirus in immunocompetent young children. *J Immunol*. 2004;172:3260-3267.
12. Moebius U. Cluster report: CD8. In: Knapp W, Dörken B, Gilks WR, et al., eds. *Leucocyte Typing IV: White Cell Differentiation Antigens*. New York, NY: Oxford University Press; 1989:342-343.
13. Terry LA, DiSanto JP, Small TN, Flomenberg N. Differential expression of the CD8 and Lyt-3 antigens on a subset of human T-cell receptor γ/δ -bearing lymphocytes. In: Knapp W, Dörken B, Gilks WR, et al., eds. *Leucocyte Typing IV: White Cell Differentiation Antigens*. New York, NY: Oxford University Press; 1989:345-346.
14. Anderson P, Blue M-L, Morimoto C, Schlossman SF. Cross-linking of T3 (CD3) with T4 (CD4) enhances the proliferation of resting T lymphocytes. *J Immunol*. 1987;139:678-682.
15. Eichmann K, Jönsson J-I, Falk I, Emmrich F. Effective activation of resting mouse T lymphocytes by cross-linking submitogenic concentrations of the T cell antigen receptor with either Lyt-2 or L3T4. *Eur J Immunol*. 1987;17:643-650.
16. Gallagher PF, Fazekas de St. Groth B, Miller JFAP. CD4 and CD8 molecules can physically associate with the same T-cell receptor. *Proc Natl Acad Sci USA*. 1989;86:10044-10048.
17. Rudd CE, Burgess KE, Barber EK, Schlossman SF. Monoclonal antibodies to the CD4 and CD8 antigens precipitate variable amounts of CD4/CD8-associated p56^{lck} activity. In: Knapp W, Dörken B, Gilks WR, et al., eds. *Leucocyte Typing IV: White Cell Differentiation Antigens*. New York, NY: Oxford University Press; 1989:326-327.
18. Engleman EG, Benike CJ, Evans RL. Circulating antigen-specific suppressor T cells in a healthy woman: mechanism of action and isolation with a monoclonal antibody. *Clin Res*. 1981;29:365A.

19. Engleman EG, Benike CJ, Glickman E, Evans RL. Antibodies to membrane structures that distinguish suppressor/cytotoxic and helper T lymphocyte subpopulations block the mixed leukocyte reaction in man. *J Exp Med*. 1981;154:193-198.
20. Reichert T, DeBruyere M, Deneys V, et al. Lymphocyte subset reference ranges in adult Caucasians. *Clin Immunol Immunopathol*. Aug 1991;60(2):190-208.
21. Cosimi AB. Anti-T-cell monoclonal antibodies in transplantation therapy. *Trans Proc*. 1983;XV:1889-1892.
22. Bernard A, Boumsell L, Hill C. Joint report of the first international workshop on human leucocyte differentiation antigens by the investigators of the participating laboratories. In: Bernard A, Boumsell L, Dausset J, Milstein C, Schlossman SF, eds. *Leucocyte Typing*. New York, NY: Springer-Verlag; 1984:9-108.
23. *Protection of Laboratory Workers from Occupationally Acquired Infections; Approved Guideline—Fourth Edition*. Wayne, PA: Clinical and Laboratory Standards Institute; 2014. CLSI document M29-A4.
24. Centers for Disease Control and Prevention. 2007 Guideline for Isolation Precautions: Preventing Transmission of Infectious Agents in Healthcare Settings. <https://www.cdc.gov/infectioncontrol/guidelines/isolation/index.html>. Accessed March 12, 2019.

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