

Technical Data Sheet

PE Mouse anti-Mouse CD178.1

Product Information

Material Number:	553854
Alternate Name:	mFasL.1, CD95 Ligand
Size:	0.2 mg
Concentration:	0.2 mg/ml
Clone:	KAY-10
Immunogen:	Transfected Cell Line
Isotype:	Mouse (C57BL/6) IgG2b, κ
Reactivity:	QC Testing: Mouse
Storage Buffer:	Aqueous buffered solution containing $\leq 0.09\%$ sodium azide.

Description

The Kay-10 antibody reacts with CD178.1, the Fas Ligand alloantigen (mFasL.1, CD95 Ligand) expressed on activated T lymphocytes of selected strains of mice (eg, C57BL/6, C3H, MRL, NOD, NZB, NZW, and SJL). It does not react with similarly activated T blasts from BALB/c, DBA/1, or DBA/2 mice. It also reacts with Cos cells transfected with *mFasL* cDNA derived from C57BL/6 and C3H mice, but not with *mFasL* cDNA from BALB/c or DBA/2 mice. In addition, Kay-10 mAb efficiently blocks cytotoxic activity of a C3H T-cell line, but not a BALB/c T-cell line. Phenotypic and genotypic characterizations of mouse Fas Ligand reveal the existence of two alloantigens: mFasL.1, which is recognized by mAbs Kay-10, MFL3 (Cat. no. 555291), and MFL4 (Cat. no. 555022), and mFasL.2 (recognized by mAbs MFL3 and MFL4). Functional studies suggest that mFasL.2 has higher specific activity than mFasL.1. In the mouse, FasL is expressed on activated T cell lines and in spleen, testis, and eye. *FasL* mRNA has been demonstrated at various levels in bone marrow, thymus, spleen, lymph node, lung, small intestine, testis, and uterus. Moreover, T-cell activators, but not B-cell activators, enhanced the expression of *FasL* mRNA in splenocytes; and *FasL* mRNA was restricted to the T-cell lineage among a panel of cell lines from lymphoid tissues. Fas Ligand is not functional in mice homozygous for the *gld* (generalized lymphoproliferative disease) mutation; these mice cannot limit the expansion of activated lymphocytes and develop autoimmune disease. Fas Ligand is a member of the TNF/NGF family which binds to CD95 (Fas), inducing apoptotic cell death. This Fas/Fas Ligand interaction is believed to participate in T-cell development, the regulation of immune responses, and cell-mediated cytotoxic mechanisms. There is mounting evidence that Fas Ligand is also pro-inflammatory, mediating neutrophil extravasation and chemotaxis. Fas Ligand is released from the surface of transfectant cells by metalloproteinases, and the soluble Fas Ligand may block the activities of the membrane-bound molecule.

Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

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

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

Application Notes

Application

Flow cytometry	Routinely Tested
----------------	------------------

Recommended Assay Procedure:

We have found that enriched splenic T cells are induced to express Fas Ligand by 6-8-hour culture with plate-bound anti-mouse CD3 antibody mAb 17A2 (Cat. no. 555273), 145-2C11 (Cat. no. 557306/553058), or 500A2 (Cat. no. 553238).

Suggested Companion Products

Catalog Number	Name	Size	Clone
553238	Purified Hamster Anti-Mouse CD3e	0.5 mg	500A2
553046	FITC Rat Anti-Mouse CD4	0.1 mg	RM4-5
553237	PE Hamster Anti-Mouse CD69	0.2 mg	H1.2F3
559529	PE Mouse IgG2b, κ Isotype Control	0.1 mg	MPC-11

BD Biosciences

bdbiosciences.com

United States	Canada	Europe	Japan	Asia Pacific	Latin America/Caribbean
877.232.8995	888.259.0187	32.53.720.550	0120.8555.90	65.6861.0633	55.11.5185.9995

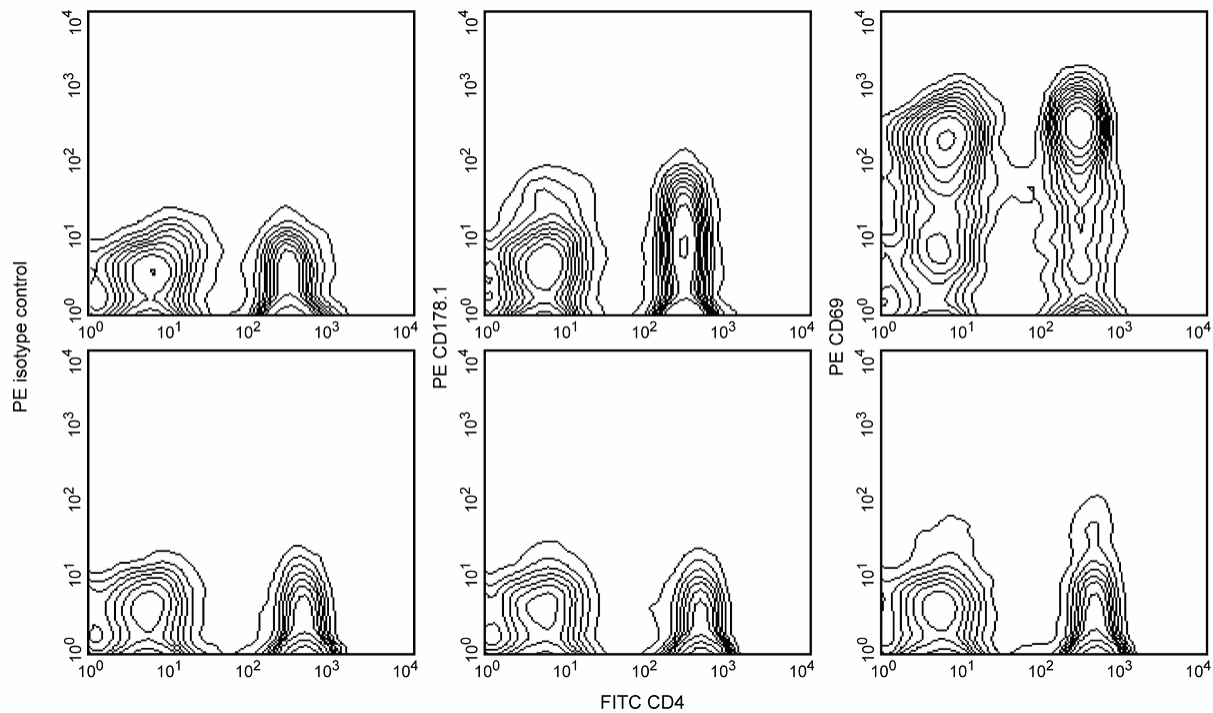
For country-specific contact information, visit bdbiosciences.com/how_to_order/

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





Expression of Fas Ligand on activated T lymphocytes. T lymphocytes from C57BL/6 spleen (mouse T Cell Enrichment Column, R&D Systems, Minneapolis, MN), cultured for 7 hours in the presence of plate-bound anti-mouse CD3e mAb 500A2 (Cat. no. 553238, Upper Panels or on uncoated plates Lower Panels), were simultaneously stained with FITC-conjugated antimouse CD4 mAb RM4-5 (Cat. no. 553046/553047) and PE-conjugated Mouse IgG2b, ϵ isotype control (Left Panels), PEconjugated mAb Kay-10 (Center Panels), or PE-conjugated anti-mouse CD69 mAb H1.2F3 (to demonstrate activation, Cat. no.553237, Right Panels). Flow cytometry was performed on a BD FACScan™ flow cytometry system.

Product Notices

1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
2. Please refer to www.bdbiosciences.com/pharming/protocols for technical protocols.
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

- Bellgrau D, Gold D, Selawry H, Moore J, Franzusoff A, Duke RC. A role for CD95 ligand in preventing graft rejection. *Nature*. 1995; 377(6550):630-632.(Biology)
- Brunner T, Mogil RJ, LaFace D, et al. Cell-autonomous Fas (CD95)/Fas-ligand interaction mediates activation-induced apoptosis in T-cell hybridomas. *Nature*. 1995; 373:441-444.(Biology)
- Griffith TS, Brunner T, Fletcher SM, Green DR, Ferguson TA. Fas ligand-induced apoptosis as a mechanism of immune privilege. *Science*. 1995; 270(5239):1189-1192.(Biology)
- Hohlbaum AM, Moe S, Marshak-Rothstein A. Opposing effects of transmembrane and soluble Fas ligand expression on inflammation and tumor cell survival. *J Exp Med*. 2000; 191(7):1209-1220.(Biology)
- Ju ST, Panka DJ, Cui H, et al. Fas(CD95)/FasL interactions required for programmed cell death after T-cell activation. *Nature*. 1995; 373(6513):444-448.(Biology)
- Kayagaki N, Yamaguchi N, Nagao F, et al. Polymorphism of murine Fas ligand that affects the biological activity. *Proc Natl Acad Sci U S A*. 1997; 94(8):3914-3919.(Immunogen)
- Kojima H, Shinohara N, Hanaoka S, et al. Two distinct pathways of specific killing revealed by perforin mutant cytotoxic T lymphocytes. *Immunity*. 1994; 1(5):357-364.(Biology)
- Lau HT, Yu M, Fontana A, Stoeckert CJ Jr. Prevention of islet allograft rejection with engineered myoblasts expressing FasL in mice. *Science*. 1996; 273(5271):109-112.(Biology)
- Lynch DH, Ramsdell F, Alderson MR. Fas and FasL in the homeostatic regulation of immune responses. *Immunol Today*. 1995; 16(12):569-574.(Biology)
- Pestano GA, Zhou Y, Trimble LA, Daley J, Weber GF, Cantor H. Inactivation of misselected CD8 T cells by CD8 gene methylation and cell death. *Science*. 1999 May; 284(5417):1187-1191.(Biology)
- Ramsdell F, Seaman MS, Miller RE, Picha KS, Kennedy MK, Lynch DH. Differential ability of Th1 and Th2 T cells to express Fas ligand and to undergo activation-induced cell death. *Int Immunol*. 1994; 6(10):1545-1553.(Biology)
- Schneider P, Holler N, Bodmer JL, et al. Conversion of membrane-bound Fas(CD95) ligand to its soluble form is associated with downregulation of its proapoptotic activity and loss of liver toxicity. *J Exp Med*. 1998; 187(8):1205-1213.(Biology)
- Smith CA, Farrah T, Goodwin RG. The TNF receptor superfamily of cellular and viral proteins: activation, costimulation, and death. *Cell*. 1994; 76(6):959-962.(Biology)
- Suda T, Okazaki T, Naito Y, et al. Expression of the Fas ligand in cells of T cell lineage. *J Immunol*. 1995; 154(8):3806-3813.(Biology)
- Takahashi T, Tanaka M, Brannan CI, Jenkins NA, Copeland NG, Suda T, and Nagata S. Generalized lymphoproliferative disease in mice, caused by a point mutation in the Fas ligand. *Cell*. 1994; 76(6):969-976.(Biology)
- Vignaux F, Vivier E, Malissen B, Depraetere V, Nagata S, Golstein P. TCR/CD3 coupling to Fas-based cytotoxicity. *J Exp Med*. 1995; 181(2):781-786.(Biology)