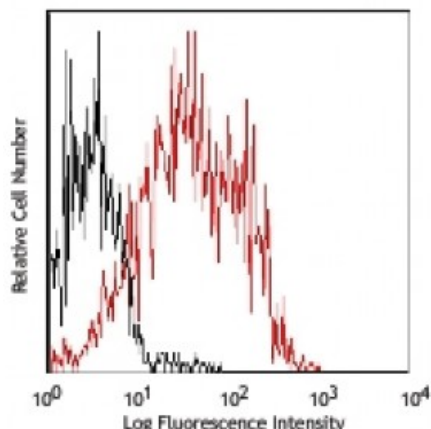


Alexa Fluor® 700 anti-mouse CD86

Catalog # / Size:	1125120 / 100 µg 1125115 / 25 µg
Clone:	GL-1
Isotype:	Rat IgG2a, κ
Immunogen:	LPS-activated CBA/Ca mouse splenic B cells
Reactivity:	Mouse
Preparation:	The antibody was purified by affinity chromatography, and conjugated with Alexa Fluor® 700 under optimal conditions.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
Concentration:	0.5



LPS-stimulated (3 days) C57BL/6 mouse splenocytes stained with GL-1 Alexa Fluor® 700

Applications:

Applications:	Flow Cytometry
Recommended Usage:	Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. The suggested use of this reagent is ≤1.0 microg per million cells in 100 microL volume. It is highly recommended that the reagent be titrated for optimal performance for each application.

* Alexa Fluor® 700 has a maximum emission of 719 nm when it is excited at 633 nm / 635 nm. Prior to using Alexa Fluor® 700 conjugate for flow cytometric analysis, please verify your flow cytometer's capability of exciting and detecting the fluorochrome.

Application Notes:	The GL-1 antibody can block the mixed lymphocyte reaction <i>in vitro</i> and has been shown to inhibit the priming of cytotoxic T lymphocytes <i>in vivo</i> (along with antibodies against B7-1). Additional reported applications (for the relevant formats) include: immunoprecipitation ¹ , immunohistochemical staining of acetone-fixed frozen sections ^{2,6} , immunofluorescence microscopy, and <i>in vivo</i> and <i>in vitro</i> blocking of T cell responses ¹⁻⁶ . GL-1 is not suitable for immunohistochemical staining of formalin-fixed paraffin sections. The LEAF™ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 105010).
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Application References:	<ol style="list-style-type: none"> Hathcock KS, <i>et al.</i> 1993. <i>Science</i> 262:905. (Block, IP) Inaba KM, <i>et al.</i> 1994. <i>J. Exp. Med.</i> 180:1849. (Block, IHC) Hathcock KS, <i>et al.</i> 1994. <i>J. Exp. Med.</i> 180:631. (Block) Krummel MF, <i>et al.</i> 1995. <i>J. Exp. Med.</i> 182:459. (Block) Liu Y, <i>et al.</i> 1997. <i>J. Exp. Med.</i> 185:251. (Block) Herold KC, <i>et al.</i> 1997. <i>J. Immunol.</i> 158:984. (Block, IHC) Shih FF, <i>et al.</i> 2006. <i>J. Immunol.</i> 176:3438. (FC) Lawson BR, <i>et al.</i> 2007. <i>J. Immunol.</i> 178:5366. Turnquist HR, <i>et al.</i> 2007. <i>J. Immunol.</i> 178:7018. Klinger MB, <i>et al.</i> 2007. <i>Am. J. Physiol. Regul. Integr. Comp. Physiol.</i> 293:R677. PubMed
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Description: CD86 is an 80 kD immunoglobulin superfamily member also known as B7-2, B70, and Ly-58. CD86 is expressed on activated B and T cells, macrophages, dendritic cells, and astrocytes. CD86, along with CD80, is a ligand of CD28 and CD152 (CTLA-4). CD86 is expressed earlier in the immune response than CD80. CD86 has also been shown to be involved in immunoglobulin class-switching and triggering of NK cell-mediated cytotoxicity. CD86 binds to CD28 to transduce co-stimulatory signals for T cell activation, proliferation, and cytokine production. CD86 can also bind to CD152, also known as CTLA-4, to deliver an inhibitory signal to T cells.

Antigen
References:

1. Barclay A, *et al.* 1997. The Leukocyte Antigen FactsBook Academic Press.
2. Hathcock KS, *et al.* 1993. *Science* 262:905.
3. Freeman GJ, *et al.* 1993. *Science* 262:907.
4. Carreno BM, *et a*