

Technical Data Sheet

FITC Hamster Anti-Mouse CD54

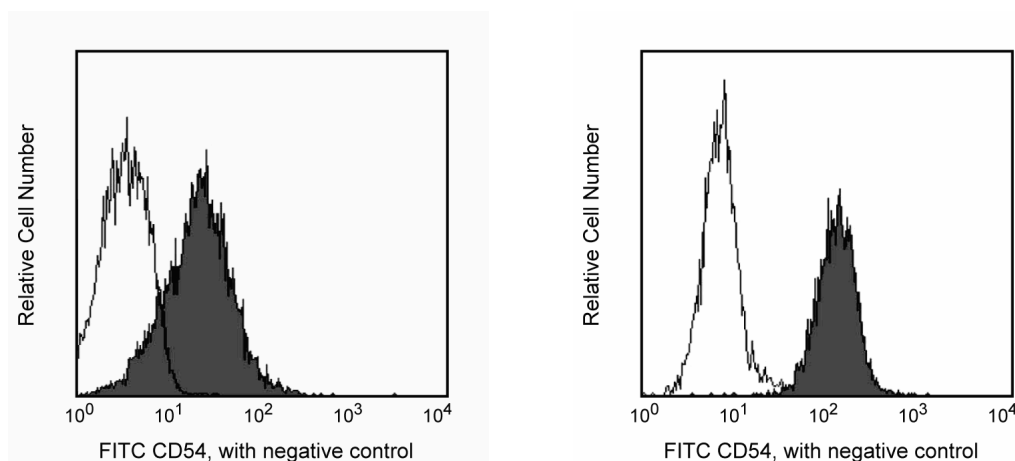
Product Information

Material Number:	553252
Alternate Name:	ICAM-1
Size:	0.5 mg
Concentration:	0.5 mg/ml
Clone:	3E2
Immunogen:	Not reported
Isotype:	Armenian Hamster IgG1, κ
Reactivity:	QC Testing: Mouse
Storage Buffer:	Aqueous buffered solution containing $\leq 0.09\%$ sodium azide.

Description

The 3E2 antibody reacts with CD54 (ICAM-1), a 95-kDa member of the Ig superfamily found on lymphocytes, vascular endothelium, high endothelial venules, epithelial cells, macrophages, and dendritic cells. ICAM-1 is a ligand for LFA1 (CD11a/CD18) and Mac-1 (CD11b/CD18). Its expression is upregulated upon stimulation by inflammatory mediators such as cytokines and LPS. Studies with mouse *Icam1*-transfected antigen-presenting cells, with CD54-blocking antibodies, and in CD54-deficient mice indicate that CD54 participates in inflammatory reactions and antigen-specific immune responses. In addition, there is evidence that CD54 is a receptor involved in MHC-non-restricted responses to weakly immunogenic tumor cells. The 3E2 antibody blocks in vitro and in vivo responses.

This antibody is routinely tested by flow cytometric analysis. Other applications were tested at BD Biosciences Pharmingen during antibody development only or reported in the literature.



Upregulation of CD54 expression on activated splenic B lymphocytes. Left panel: Naive BALB/c splenocytes were stained with FITC-conjugated 3E2 mAb (filled histogram) or unstained (open histogram). Viable resting lymphocytes were gated according to scatter profile and exclusion of 7-AAD (BD Via-Probe™, Cat. No. 555816/555815). The mean fluorescence intensity of the stained lymphocytes is about 8 times greater than that of the negative-control lymphocytes. Right panel: 2-day LPS-activated BALB/c splenocytes were stained with FITC-conjugated 3E2 mAb (filled histogram) or unstained (open histogram). Viable B-cell blasts were gated according to scatter profile and exclusion of 7-AAD. The mean fluorescence intensity of the stained blasts is about 17 times greater than that of the negative-control blasts. Flow cytometry was performed on a BD FACScan™ flow cytometry system.

Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. The antibody was conjugated with FITC under optimum conditions, and unreacted FITC was removed. Store undiluted at 4° C and protected from prolonged exposure to light. Do not freeze.

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Application Notes

Application

Flow cytometry	Routinely Tested
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Suggested Companion Products

Catalog Number	Name	Size	Clone
555816	Cell Viability Solution	100 tests	(none)
553971	FITC Hamster IgG1 κ Isotype Control	0.25 mg	A19-3

Product Notices

1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
2. Please refer to www.bdbiosciences.com/pharming/en/protocols for technical protocols.
3. Although hamster immunoglobulin isotypes have not been well defined, BD Biosciences Pharmingen has grouped Armenian and Syrian hamster IgG monoclonal antibodies according to their reactivity with a panel of mouse anti-hamster IgG mAbs. A table of the hamster IgG groups, Reactivity of Mouse Anti-Hamster Ig mAbs, may be viewed at http://www.bdbiosciences.com/pharming/hamster_chart_11x17.pdf.
4. 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

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