Technical Data Sheet

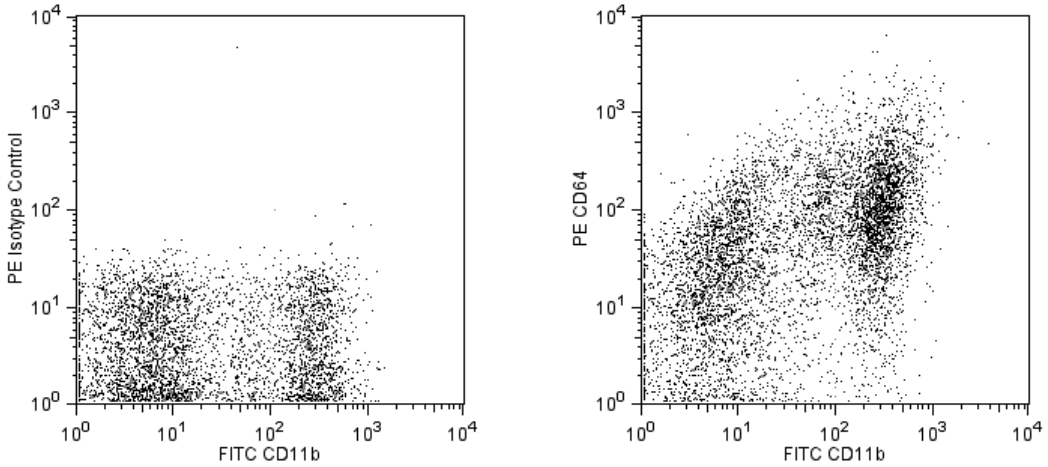
PE Mouse anti-Mouse CD64 a and b Alloantigens

Product Information

Material Number:	**558455**
Alternate Name:	FcgRI
Size:	0.1 mg
Concentration:	0.2 mg/ml
Clone:	X54-5/7.1
Immunogen:	Mouse CD64 a Alloantigen
Isotype:	Mouse (NOD/Lt) IgG1, κ
Reactivity:	confirmed postivie strains: BALB/c, C57BL/6 reported positive strains: 129, A, AKR, ALR, BUB, C3H, C57BL/10, C57BLKS, C57BR, C58, CBA, CE, DBA/2, HRS, MRL, NON, NZB, NZO, NZW, PL, SJL, ST, SWR reported negative strains: ABH, NOD
Storage Buffer:	Aqueous buffered solution containing ≤0.09% sodium azide.

Description

The monoclonal alloantibody X54-5/7.1 reacts with FcγRI (CD64) encoded by the more common Fcgr1a and Fcgr1b alleles. The alloantigens generated by the Fcgr1a and Fcgr1b alleles, have been confirmed positive in mouse strains BALB/c and C57BL/6 and reported positive in strains 129, A, AKR, ALR, BUB, C3H, C57BL/10, C57BLKS, C57BR, C58, CBA, CE, DBA/2, HRS, MRL, NON, NZB, NZO, NZW, PL, SJL, ST, SWR. The a and b alloantigens have been reported negative in mouse strains ABH, NOD. CD64, a key receptor in the development of immune responses, has a dual role as a low affinity receptor for IgG3 and high affinity receptor for IgG2a linking innate and adaptive immunities. CD64 mediates endocytosis, phagocytosis, antibody-dependent cellular toxicity, cytokine release and superoxide generation. CD64 is expressed largely on macrophages and dendritic cells. For more information regarding clone X54-5/7.1 and the alloantigens it recognizes, please refer to the reference by Tan et al listed below.



Flow cytometric analysis of PE-conjugated anti-mouse CD64 recognizing a and b Alloantigens on mouse bone marrow cells. Isolated murine bone marrow cells were preincubated with Mouse BD Fc Block™ purified anti-mouse CD16/CD32 mAb 2.4G2 (Cat. No. 553141/553142). The cells were then stained with FITC anti-CD11b (clone M1/70, catalog number No. 553310) and either PE anti-CD64 (clone X54-5/7.1, Cat. No. 558455, right panel) or a PE mouset IgG1 isotype control (catalog number 550617, left panel). Flow cytometry was performed on a BD FACSCalibur™ System and the dot plots were derived from the gated events based on light scattering characteristics of viable bone marrow cells.

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.

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Application Notes

Application

Flow cytometry	Routinely Tested

Recommended Assay Procedure:

For flow cytometry of cell suspensions from peripheral lymphoid tissues, it is recommended that the cells be pre-incubated with Mouse BD Fc Block™ purified anti-mouse CD16/CD32 mAb 2.4G2 (Cat. no.553141/553142).

Suggested Companion Products

Catalog Number	Name	Size	Clone
550617	PE Mouse IgG1, κ Isotype Control	0.1 mg	MOPC-31C

Product Notices

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

Kishimoto T, von dem Borne AEG, Goyert SM,et al., ed. *Leucocyte Typing VI: White Cell Differentiation Antigens.* London: Garland Publishing; 1997. (Biology)

Tan PS, Gavin AL, Barnes N, et al. Unique monoclonal antibodies define expression of Fc gamma RI on macrophages and mast cell lines and demonstrate heterogeneity among subcutaneous and other dendritic cells. *J Immunol.* 2003; 170(5):2549-2556. (Clone-specific)