

PE Mouse anti-Mouse CD45.1 mAb

Catalog No.: A28894

Basic Information

Observed MW

Calculated MW

Category

Primary antibody

Applications

FC

Cross-Reactivity

Mouse

CloneNo number

AMC0922

Conjugate

PE. Ex:565nm. Em:574nm.

Background

Enables several functions, including heparan sulfate proteoglycan binding activity; heparin binding activity; and protein tyrosine phosphatase activity. Involved in several processes, including lymphocyte differentiation; positive regulation of macromolecule metabolic process; and regulation of signal transduction. Acts upstream of or within several processes, including lymphocyte differentiation; positive regulation of lymphocyte activation; and regulation of protein phosphorylation. Located in external side of plasma membrane; focal adhesion; and membrane raft. Is expressed in several structures, including 3rd branchial arch; alimentary system; cardiovascular system; hemolymphoid system; and placenta. Used to study systemic lupus erythematosus. Human ortholog(s) of this gene implicated in hepatitis C; multiple sclerosis; severe combined immunodeficiency; and severe combined immunodeficiency, autosomal recessive, T cell-negative, B cell-positive, Nk cell-positive. Orthologous to human PTPRC (protein tyrosine phosphatase receptor type C).

Recommended Dilutions

FC ≤0.25 µg per million cells
in 100 µl volume

Immunogen Information

Gene ID

19264

Swiss Prot

P06800

Immunogen

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

Synonyms

loc; B220; Cd45; L-CA; Ly-5; T200; CD45R; Lyt-4

Contact

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Product Information

Source

Mouse

Isotype

Mouse IgG2a, κ

Purification

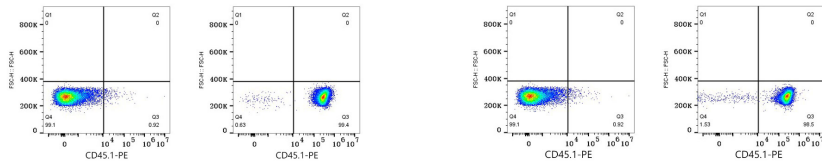
Affinity purification

Storage

Store at 2-8°C. Avoid freeze.

Buffer: PBS with 0.09% Sodium azide, 0.2% BSA, pH7.3.

Validation Data



Flow cytometry: 1×10^6 C57BL/6 mouse splenocytes (negative control, left) and B6-CD45.1 mouse splenocytes (right) were surface-stained with PE Mouse anti-Mouse CD45.1 mAb (A28894, 0.25 μ g). Cells in the lymphocyte gates were used for analysis.

Flow cytometry: 1×10^6 C57BL/6 mouse splenocytes (negative control, left) and SJL mouse splenocytes (right) were surface-stained with PE Mouse anti-Mouse CD45.1 mAb (A28894, 0.25 μ g). Cells in the lymphocyte gates were used for analysis.