Technical Data Sheet

PE Mouse anti-Lck

Product Information

 Catalog Number:
 558496

 Size:
 50 tests

 Volume per Test:
 20 µl

 Clone:
 MOL 171

Immunogen: Human N-terminal Lck Peptide

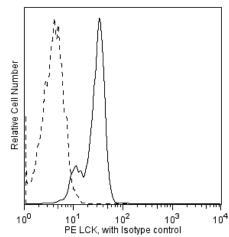
Isotype:Mouse IgG1Reactivity:Tested: HumanReported: Mouse

Storage Buffer: Aqueous buffered solution containing BSA and $\leq 0.09\%$ sodium azide

Description

Lck is a member of the Src family of cytoplasmic protein-tyrosine kinases (PTKs) that is normally expressed exclusively in lymphoid cells, primarily T lymphocytes and NK cells. A low level of expression has been detected in B lymphocytes, but its function in B cells is unknown. Its expression in other leukocytes is not well defined. Members of the Src family have several common features: 1) unique N-terminal domains, 2) attachment to cellular membranes through a myristylated N-terminus, and 3) homologous SH2, SH3, and catalytic domains. The unique N-terminal domain of Lck interacts with the cytoplasmic tails of the CD4 and CD8 cell-surface glycoproteins of T lymphocytes, which recognize antigen presenting cells via their surface MHC class II and class I molecules, respectively. The catalytic activity of Lck is regulated by both kinases and phosphatases that control the phosphorylation states of two tyrosine residues that have opposing effects. Repression of Lck's catalytic activity occurs via phosphorylation at tyrosine 505 (Y505), located near the carboxy terminus. Phosphorylation of this tyrosine site is mediated by the Csk family of PTKs, and its dephosphorylation is mediated by the protein tyrosine phosphatase, CD45. When Lck is phosphorylated at this site, it assumes a folded tertiary structure which is enzymatically inactive. When CD45 dephosphorylates it at Y505, Lck is able to autophosphorylate its Y394, which leads to conformational changes in the catalytic domain that induce kinase activity. However, it has been observed that the inhibitory effect of the phosphorylated Y505 can be overcome by direct engagement of Lck's SH3 domain and that both Y394 and Y505 are phosphorylated together in cells activated by hydrogen peroxide. Activated Lck phosphorylates the ITAMs (Immunoreceptor-based Tyrosine Activation Motifs) of the T cell receptor (TCR) and thus is critical for activation and development of T lymphocytes. The interactions of Lck, Csk, CD45, CD4 or CD8, and TCR are only a small part of a complex immunoregulatory cascade that involves additional substrates for Csk and CD45, other enzymes, adhesion molecules, adaptor proteins, and specialized membrane microdomains.

The MOL 171 monoclonal antibody recognizes the 56- and 60-kDa forms of human Lck protein, regardless of phosphorylation status. It cross reacts with mouse Lck.



Analysis of Lck in human peripheral blood lymphocytes. Human whole blood was lysed and fixed with 1X BD™ Phosflow Lyse/Fix Buffer (Cat. No. 558049) for 10-15 minutes at 37°C, then permeabilized (BD™ Phosflow Perm Buffer II, Cat. No. 558052) on ice for 30 minutes, and then stained with either PE Mouse anti-Lck (solid-line histogram) or Mouse IgG1, κ isotype control mAb MOPC-21 (Cat. No. 559320, dashed-line histogram). Flow cytometry was performed on a BD™ FACSCalibur flow cytometry system. The figure displays lymphocytes that were selected by their scatter profile.

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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 by gel filtration chromatography.

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

Application Notes

Application	Recommendation	Confirmation
Intracellular staining (flow cytometry)	Recommended	Tested

Suggested Companion Products

Catalog Number	Name	Size	Clone
558052	Perm Buffer II	125 ml	(none)
558049	Lyse/Fix Buffer 5X	250 ml	(none)
559320	PE Mouse IgG1κ Isotype Control	100 tests	MOPC-21

Product Notices

- 1. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- 2. This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use 1 X 10e6 cells in a 100-μl experimental sample (a test).
- 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.
- 4. Source of all serum proteins is from USDA inspected abattoirs located in the United States.

References

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Holdorf AD, Lee K-H, Burack WR, Allen PM, Shaw AS. Regulation of Lck activity by CD4 and CD28 in the immunological synapse. *Nat Immunol.* 2002; 3(3):259-264. Johnson KG, Bromley SK, Dustin ML, Thomas ML. A supramolecular basis for CD45 tyrosine phosphatase regulation in sustained T cell activation. *Proc Natl Acad Sci U S A.* 2000; 97:10138-10143.

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