### Purified Mouse Anti-Human p56-Lck w/ Control

#### Product Information

<table>
<thead>
<tr>
<th>Material Number</th>
<th>Size</th>
<th>Reactivity</th>
</tr>
</thead>
<tbody>
<tr>
<td>551048</td>
<td>150 µg</td>
<td>QC Testing: Human Reported: Mouse</td>
</tr>
</tbody>
</table>

#### Component: 51-1495GR

- **Description:** Purified Mouse Anti-Human p56-Lck
- **Size:** 50 µg (3 ea)
- **Concentration:** 0.25 mg/ml
- **Clone Name:** MOL 171
- **Immunogen:** Human N-terminal Lck
- **Isotype:** Mouse (BALB/c) IgG1, κ
- **Storage Buffer:** Aqueous buffered solution containing BSA, glycerol, and ≤0.09% sodium azide.

#### Component: 51-16526N

- **Description:** Jurkat Cell Lysate
- **Size:** 50 µg (1 ea)
- **Concentration:** 1.0 mg/ml
- **Storage Buffer:** SDS-PAGE buffer (62mM Tris pH 6.8, 2% SDS, 0.9% b-mercaptoethanol, 0.003% bromophenol blue, 5% glycerol)

### Description

The T-cell receptor (TCR) is a multi-chain transmembrane receptor responsible for antigen recognition on the T-cell surface. T cells also express several other integral membrane proteins, including CD4 and CD8, which play significant roles in the functional responses of the TCR to antigen presentation. Following antigen or ligand binding to the TCR, a series of interrelated membrane and cytoplasmic activation or signaling events rapidly occurs. These events include tyrosine phosphorylation of membrane and cytoplasmic proteins, plasma membrane inositol phospholipid hydrolysis, increases in cytoplasmic calcium concentrations, and increases in protein kinase C activity. The earliest measurable biochemical changes are the appearance of newly phosphorylated tyrosine residues on a variety of cytoplasmic and membrane proteins. Phosphorylation of tyrosine residues is mediated by protein tyrosine kinases (PTKs). Several different PTKs have been implicated in early phosphorylation events of T cell activation, including lck. p56-lck is a member of the Src family, and its functional domains can be divided into two regions based on sequence comparison with p60-src. The N-terminal half, which is highly divergent, contains the membrane bound/substrate interactive domain. The C-terminal half, which has more conserved homology, contains the kinase domain. Lck is normally expressed exclusively in cells of lymphoid lineage, primarily T cells, and natural killer cells. Lck is generally expressed at detectable levels in T-cell lines, including CTLL-2 (ATCC TIB-214) and Jurkat. Lower levels of lck expression have been detected in B cells. Aberrant expression of lck has been described in human colon and lung carcinoma cell lines. Lck plays a role T-cell signal transduction through its physical association with the cytoplasmic tails of CD4 and CD8 expressed in helper and cytolytic T-cells, respectively. The importance of lck in T cell activation is supported by genetic studies. For example, mice with lck null mutations lack T cell development. Additionally, mice expressing only mutant CD4 that is unable to bind to lck, lack the ability to activate T cells through the TCR. MOL 171 reacts with human lck proteins (56-60 kDa). It cross-reacts with mouse lck proteins. A 25 amino acid synthetic peptide corresponding to the N-terminal region of the human lck sequence was used as immunogen.

### Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. Store undiluted at -20°C.
Western blot analysis of lck. Lysate from Jurkat cells was probed with anti-lck (clone MOL 171, Cat. No. 51-1495GR) at concentrations of 2.0 (lane 1), 1.0 (lane 2), and 0.5 µg/ml (lane 3). lck is identified as a band of 56 kDa.

Application Notes

<table>
<thead>
<tr>
<th>Application</th>
<th>Routinely Tested</th>
<th>Reported</th>
</tr>
</thead>
<tbody>
<tr>
<td>Western blot</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Immunoprecipitation</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Recommended Assay Procedure:
Applications include western blot analysis (0.5-2.0 µg/ml). Other applications not routinely tested at BD Biosciences Pharmingen include immunoprecipitation (2.0-5.0 µg/ml). lck is typically detected as one or multiple bands ranging between 56-60 kDa. Jurkat control lysate (50 µg (1 µg/µl)) is provided as a western blot positive control (Cat. No. 51-16526N; store lysate at -20°C). Additional control lysate (Cat. No. 611451) is sold separately. U937 human histiocytic lymphoma cells (ATCC CRL-1593) are suggested as a negative control. For western blot analysis, 5 µg total cell protein per lane is recommended for SDS/PAGE mini-gels. Non-specific bands may be detected if too much protein is loaded onto the gel.

Suggested Companion Products

<table>
<thead>
<tr>
<th>Catalog Number</th>
<th>Name</th>
<th>Size</th>
<th>Clone</th>
</tr>
</thead>
<tbody>
<tr>
<td>554002</td>
<td>HRP Goat Anti-Mouse Ig</td>
<td>1.0 ml</td>
<td>(none)</td>
</tr>
<tr>
<td>611451</td>
<td>Jurkat Cell Lysate</td>
<td>500 µg</td>
<td>(none)</td>
</tr>
</tbody>
</table>

Product Notices
1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
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