

Lck Kinase

5 µg

#7757

This product is for *in vitro* research use only and is not intended for use in humans or animals.

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Description: Purified recombinant human Lck kinase (Met1-Pro509), supplied as a GST fusion protein.

Background: The Src family of protein tyrosine kinases (including Src, Lyn, Fyn, Yes, Lck, Blk, Hck, etc.) is important in the regulation of growth and differentiation of eukaryotic cells (1). Src activity is regulated by tyrosine phosphorylation at two sites with opposing effects. Phosphorylation of Tyr416 in the activation loop of the kinase domain upregulates enzyme activity. Phosphorylation of Tyr527 in the carboxy-terminal tail by Csk renders the enzyme less active (2).

Source/Purification: The GST-Lck Kinase fusion protein was produced using a baculovirus expression system with a construct expressing human Lck (Met1-Pro509) (GenBank accession No. NM_005356) with an amino-terminal GST tag. The protein was purified by one-step affinity chromatography using glutathione-agarose.

Quality Control: The theoretical molecular weight of the GST-Lck fusion protein is 93 kDa. The purified kinase was quality controlled for purity using SDS-PAGE followed by Silver stain and Western blot [Fig.1]. Lck kinase activity was determined using a radiometric assay [Fig.2]. A kinase dose dependency assay was performed to measure Lck activity using HTScan™ Lck Kinase Assay Kit #7758 [Fig.3].

Background References:

- (1) Thomas, S.M. and Brugge, J.S. (1997) *Annu. Rev. Cell Dev. Biol.* 13, 513-609.
- (2) Hunter, T. (1987) *Cell* 49, 1-4.

Storage: Enzymes are supplied in 50 mM Tris-HCl (pH 8.0), 100 mM NaCl, 5 mM DTT, 15 mM reduced glutathione and 20% glycerol. Store at -80°C.

Keep enzymes on ice during use.

Avoid repeated freeze-thaw cycles.

Companion Products:

- HTScan™ LCK Kinase Assay Kit #7758
- FHIT (Tyr114) Biotinylated Peptide #1365
- Phospho-Tyrosine Mouse mAb (P-Tyr-100) #9411
- HTScan™ Tyrosine Kinase Buffer (4X) #9805
- ATP (10 mM) #9804
- Staurosporine #9953
- Tyrosine Kinase Substrate Screening Kit #7450

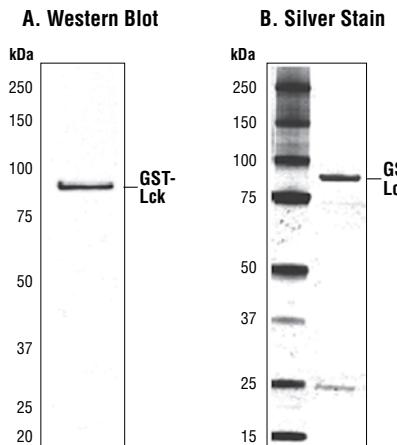


Figure 1. The purity of the GST-Lck fusion protein was analyzed using SDS/PAGE followed by anti-Lck Western blot (A) or Silver stain (B).

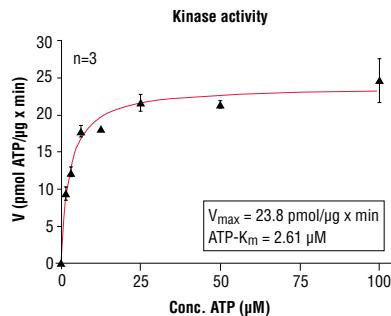


Figure 2. Lck kinase activity was measured in a radiometric assay using the following reaction conditions: 60 mM HEPES-NaOH, pH 7.5, 3 mM MgCl₂, 3 mM MnCl₂, 3 μM Na-orthovanadate, 1.2 mM DTT, ATP (variable), 2.5 μg/50 μl PEG20,000, Substrate: PolyEY, 2 μg/50 μl and 200 ng/50 μl Recombinant Lck.

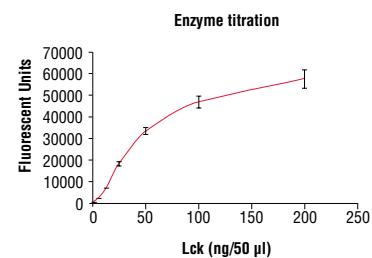


Figure 3. Dose dependence curve of Lck kinase activity: DELFIA® data generated using Phospho-Tyrosine mAb (P-Tyr-100) #9411 to detect phosphorylation of substrate peptide (#1365) by Lck kinase. In a 50 μl reaction, increasing amounts of Lck and 1.5 μM substrate peptide were used per reaction at room temperature for 30 minutes. (DELFIA® is a registered trademark of PerkinElmer, Inc.)

Protocol for Lck Kinase Assay

***IMPORTANT:** Use of an automated microplate washer as well as centrifugation of plates when appropriate, greatly improves reproducibility.

Kinase

Note: Lot-specific information for this kinase is provided on the enzyme vial. Optimal assay incubation times and enzyme concentrations must be determined empirically for each lot of kinase under specified conditions.

A Additional Solutions and Reagents (Not included)

1. **Wash Buffer:** 1X PBS, 0.05% Tween-20 (PBS/T)
2. Bovine Serum Albumin (BSA)
3. **Stop Buffer:** 50 mM EDTA pH 8
4. Phospho-Tyrosine Mouse mAb (P-Tyr-100) #9411
5. HTScan™ Kinase Buffer (10X) #9805
6. ATP (10 mM) #9804
7. FHIT (Tyr114) Biotinylated Peptide #1365
8. DTT (1000X, 1.25 M)
9. DELFIA® Europium-labeled Anti-mouse IgG (PerkinElmer Life Sciences #AD0124)
10. DELFIA® Enhancement Solution (PerkinElmer Life Sciences #1244-105)
11. DELFIA® Streptavidin coated, 96-well, yellow plate (PerkinElmer Life Sciences AAAND-0005)

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B Suggested Protocol for 100 Assays

1. Add 100 μ l 10 mM ATP to 1.25 ml 6 μ M substrate peptide. Dilute the mixture with dH₂O to 2.5 ml to make 2X ATP/substrate cocktail ([ATP]=400 μ M, [substrate] = 3 μ M).
2. Immediately transfer enzyme from -80°C to ice. Allow enzyme to thaw on ice.
3. **Microcentrifuge briefly at 4°C to bring liquid to the bottom of the vial. Return immediately to ice.**
4. Add 10 μ l of DTT (1.25 M) to 2.5 ml of 4X HTScan™ Tyrosine Kinase Buffer (240 mM HEPES pH 7.5, 20 mM MgCl₂, 20 mM MnCl₂, 12 μ M Na₃VO₄) to make DTT/Kinase buffer.
5. Transfer 1.25 ml of DTT/Kinase buffer to enzyme tube to make 4X reaction cocktail ([enzyme]=4.0 ng/ μ L in 4X reaction cocktail).
6. Incubate 12.5 μ l of the 4X reaction cocktail with 12.5 μ l/well of prediluted compound of interest (usually around 10 μ M) for 5 minutes at room temperature.
7. Add 25 μ l of 2X ATP/substrate cocktail to 25 μ l/well preincubated reaction cocktail/compound.

Final Assay Conditions for a 50 μ l Reaction

60 mM HEPES pH 7.5
5 mM MgCl₂
5 mM MnCl₂
3 μ M Na₃VO₄
1.25 mM DTT
200 μ M ATP
1.5 μ M peptide
50 ng Lck Kinase

8. Incubate reaction plate at room temperature for 30 minutes.
9. Add 50 μ l/well Stop Buffer (50 mM EDTA, pH 8) to stop the reaction.
10. Transfer 25 μ l of each reaction and 75 μ l dH₂O/well to a 96-well streptavidin-coated plate and incubate at room temperature for 60 minutes.
11. *Wash three times with 200 μ l/well PBS/T.
12. Dilute primary antibody in PBS/T with 1% BSA. Add 100 μ l/well primary antibody.
- Please note:** This protocol was validated using a FHIT (Tyr114) Biotinylated Peptide and Phospho-Tyrosine Mouse mAb (P-Tyr-100) diluted 1:1000 (see additional reagents). Primary antibody chosen should be specific to the substrate used.
13. Incubate at room temperature for 120 minutes.
14. *Wash three times with 200 μ l/well PBS/T.
15. Dilute Europium labeled anti-mouse IgG 1:500 in PBS/T with 1% BSA. Add 100 μ l/well diluted antibody.
16. Incubate at room temperature for 30 minutes.
17. *Wash five times with 200 μ l/well PBS/T.
18. Add 100 μ l/well DELFIA® Enhancement Solution.
19. Incubate at room temperature for 5 minutes.
20. Detect 615 nm fluorescence emission with appropriate Time-Resolved Plate Reader.

Please contact Cell Signaling Technology for HTS-ready antibodies (PBS formulated and carrier-free), and detailed peptide substrate sequence information.
Email: drugdiscovery@cellsignal.com