

Phospho-PKC pan (α Tyr419) Biotinylated Peptide

1.25 ml at 6 μ M

New 09/07



Cell Signaling
TECHNOLOGY®

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

Description: This biotinylated peptide contains the residues surrounding Tyr419 of PKC α . The tyrosine residue in the peptide corresponding to Tyr419 has been chemically phosphorylated in the course of peptide synthesis. This phosphopeptide was generated for use as a positive control in kinase assay (DDR2 Kinase #7414), but it may also serve as a positive control in other heterogeneous or homogeneous kinase assays.

Peptide Core Sequence: V(M/I)EY*VNG

Molecular Weight: 2325 daltons

Quality Control: The quality of the biotinylated peptide was evaluated by reverse-phase HPLC and by mass spectrometry.

Notes on Use: This phosphorylated peptide can be detected with Phospho-Tyrosine mAb (P-Tyr-100) #9411. Sample kinase assay protocol is attached.

Storage: Supplied in 0.0001% DMSO. Store at -20°C.

Companion Products:

Tyrosine Kinase Substrate Screening Kit #7450

Phospho-Tyrosine Mouse mAb (P-Tyr-100) #9411

DDR2 Kinase #7414

PKC(pan) (α Tyr419) Biotinylated Peptide #1408

Protocol for Tyrosine Kinase Assay

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

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 mAb (P-Tyr-100) #9411
5. Kinase Buffer (4X) #9805
6. ATP (10 mM) #9804
7. DTT (1.25M)
8. Kinase (See companion products)

B Suggested Protocol for 100 Assays

1. Add 100 μ l 10 mM ATP to 1.25 ml 6-12 μ M substrate peptide. Adjust the mixture with dH_2O to 2.5 ml to make 2X ATP/substrate cocktail ([ATP]=400 μ M, [substrate] = 3-6 μ M).
2. 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 (1.25M) to 2.5 ml of 4X HTScan[®] tyrosine kinase buffer (240 mM HEPES pH 7.5, 20 mM MgCl_2 , 20 mM MnCl_2 , 12 μ M Na_3VO_4) to make 4xDTT/Kinase buffer.
5. Dilute enzyme in 1.25 ml of 4X DTT/Kinase buffer to make 4X reaction cocktail ([enzyme]=0.8-8.0 ng/ μ l in 4X DTT/Kinase buffer).
6. Add 12.5 μ l of the 4X reaction cocktail to 12.5 μ l/well of prediluted compound of interest (usually around 10 μ M) and incubate 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 (pH7.5)
5 mM MgCl_2
5 mM MnCl_2
3 μ M Na_3VO_4
1.25 mM DTT
200 μ M ATP
1.5-3 μ M peptide
10-100 ng 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 to a 96-well streptavidin-coated plate containing 75 μ l dH_2O /well and incubate at room temperature for 60 minutes.
11. Wash three times with 200 μ l/well PBS/T.
12. Dilute primary antibody (Phospho-Tyrosine mAb (P-Tyr-100) #9411) in PBS/T with 1% BSA. *Add 100 μ l/well primary antibody.
13. Incubate at 37°C for 120 minutes.
14. Wash three times with 200 μ l/well PBS/T.
15. For DELFIA[®] or Colorimetric ELISA detection methods please use the following protocols.

DELFIA[®] is a registered trademark of PerkinElmer Life Sciences

DELFIA[®] Assay

1. Prepare appropriate dilution of Europium labeled secondary antibody in PBS/T with 1% BSA (1:500 dilution for anti-mouse IgG or 1:1000 for anti-rabbit IgG).
 2. Add 100 μ l/well secondary antibody solution.
 3. Incubate at room temperature for 30 minutes.
 4. *Wash five times with 200 μ l/well PBS/T.
 5. Add 100 μ l/well DELFIA[®] Enhancement Solution.
 6. Incubate at room temperature for 5 minutes.
 7. Read plate using a Time Resolved Fluorescent plate reader using the following settings;
 - a. Excitation Filter: 340 nm
 - b. Emission Filter: 615 nm
 - c. Delay⁺⁺: 400 μ s
- ⁺⁺ Delay time is the delay from the excitation pulse to the beginning of the measurement.

Companion Products for DELFIA[®]

DELFIA[®] Europium-labeled Anti-mouse IgG (PerkinElmer Life Sciences #AD0124)
DELFIA[®] Europium-labeled Anti-rabbit IgG (PerkinElmer Life Sciences #AD0105)
DELFIA[®] Enhancement Solution (PerkinElmer Life Sciences #1244-105)
DELFIA[®] Streptavidin coated, 96-well, yellow plate (PerkinElmer Life Sciences AAAND-0005)

Colorimetric ELISA Assay

1. Prepare appropriate dilution of HRP labeled secondary antibody in PBS/T with 1% BSA (1:500 dilution for anti-mouse IgG or 1:1000 for anti-rabbit IgG).
2. Add 100 μ l/well secondary antibody solution.
3. Incubate at room temperature for 30 minutes.
4. *Wash five times with 200 μ l/well PBS/T.
5. Add 100 μ l/well TMB substrate.
6. Incubate at room temperature for 15 minutes.
7. Add 100 μ l/well of stop solution.
8. Mix well.
9. Read the absorbance at 450 nm with a microtiter plate reader.

Companion Products For Colorimetric ELISA Assay

Anti-mouse IgG, HRP Linked Antibody #7076
Anti-rabbit IgG, HRP Linked Antibody #7074
TMB Solution #7004
Stop Solution #7002

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

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