PKCpan (Tyr419 α) Biotinylated Peptide

🗹 1.25 ml at 6 µM



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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 Tyrosine 419 of PKC α . It was generated for use in CST's HTScan[®] kinase assay kits, but may also serve as a substrate in other heterogeneous or homogeneous kinase assays.

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

Molecular Weight: 2245 daltons

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

Directions for Use: The phosphorylated form of the peptide can be detected with the 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

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)
- DELFIA[®] Europium-labeled Anti-mouse IgG (PerkinElmer Life Sciences #AD0124)
- **10.** DELFIA[®] Enhancement Solution (PerkinElmer Life Sciences #1244-105)
- 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 6-12 μ M substrate peptide. Adjust the mixture with dH_20 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.
- Add 10 ml of DTT (1.25M) to 2.5 ml of 4X HTScan[®] tyrosine kinase buffer (240 mM HEPES pH 7.5, 20 mM MgCl₂, 20 mM MnCl₂, 12 mM Na₃VO₄) to make 4xDTT/Kinase buffer.
- Dilute enzyme in 1.25 ml of 4X DTT/Kinase buffer to make 4X reaction cocktail ([enzyme]=0.8-8.0 ng/µl in 4X DTT/Kinase buffer).
- Add 12.5 µl of the 4X reaction cocktail to 12.5 µl/well of prediluted compound of interest (usually around 10 µM) and incubate for 5 minutes at room temperature.
- **7.** Add 25 μl of 2X ATP/substrate cocktail to 25 μl/well preincubated reaction cocktail/compound.

Final Assay Conditions for a 50 µl Reaction

- 60 mM HEPES (pH7.5)
- 5 mM MgCl_2 5 mM MnCl_2
- 3 mM Na₃VO₄ 1.25 mM DTT
- 200 µM ATP
- 1.5-3 µM peptide
- 10-100 ng kinase
- **8.** Incubate reaction plate at room temperature for 30 minutes.
- 9. Add 50 µl/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 dH₂O/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.
- Wash three times with 200 μl/well PBS/T.
 Dilute Europium labeled secondary antibody in PBS/T with 1% BSA. **Add 100 μl/well diluted antibody.
- **16.** Incubate at room temperature for 30 minutes.
- **17.** Wash five times with $200 \,\mu$ /well PBS/T.
- **18.** Add 100 µl/well DELFIA® Enhancement Solution.
- **19.** Incubate at room temperature for 5 minutes.
- **20.** Detect 615 nm fluorescence emission with appropriate Time-Resolved Plate Reader.

*Recommended antibody dilution factor:

Primary antibody: Mouse mAb: 1:500 Rabbit mAb or poly-clone antibody: 1:1000

**Secondary antibody:

DELFIA® Europium-labeled Anti-mouse IgG: 1:500 DELFIA® Europium-labeled Anti-rabbit antibody: 1:1000