

Akt Substrate-3 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 was generated for the 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: RXRXXT*
Molecular Weight: 912 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-Akt Substrate (RXRXXS/T) (110B7) Rabbit mAb #9614. Sample kinase assay protocol is attached.

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

Companion Products:

Serine/Threonine Kinase Substrate Screening Kit #7400 Phospho-Akt Substrate (RXRXXS/T) (110B7) Rabbit mAb

Mst1 Kinase #7364



Protocol for Serine/Threonine 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)

Bovine Serum Albumin (BSA)
 Stop Buffer: 50 mM EDTA pH 8

4. Kinase Buffer (10X) #9802

5. ATP (10 mM) #9804

 DELFIA® Europium-labeled Anti-rabbit Antibody (PerkinElmer Life Sciences #AD0105) or DELFIA® Europium-labeled Anti-mouse IgG (PerkinElmer Life Sciences #AD0124)

7. DELFIA® Enhancement Solution (PerkinElmer Life Sciences #1244-105)

 DELFIA® Streptavidin coated, 96-well, yellow plate (PerkinElmer Life Sciences AAAND-0005)

9. Active kinase (See companion products)

10. Primary antibody (See companion products)

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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 dH₂0 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.
- Microcentrifuge briefly at 4°C to bring liquid to the bottom of the vial. Return immediately to ice.
- 4. Add 1 ml 10X kinase buffer [250 mM Tris-HCl pH 7.5, 100 mM MgCl $_2$, 1 mM Na $_3$ VO $_4$, 50 mM β -glycerophosphate, 20 mM dithiothreitol (DTT)] to 1.5 ml dH $_3$ 0 to make 2.5 ml 4X reaction buffer.
- 5. Dilute enzyme in 1.25 ml of 4X reaction buffer to make 4X reaction cocktail ([enzyme]=0.8-8.0 ng/µl in 4X reaction cocktail).
- 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.
- Add 25 μI of 2X ATP/substrate cocktail to 25 μI/well preincubated reaction cocktail/compound.

Final Assay Conditions for a 50 µl Reaction

25 mM Tris-HCI (pH7.5)

10 mM MgCl₂

5 mM β-glycerophosphate

0.1 mM Na₃VO₄

2 mM DTT

200 uM 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 μl of each reaction to a 96-well streptavidin-coated plate containing 75 μl dH_•O/well and incubate at room temperature for 60 minutes.
- 11. Wash three times with 200 µl/well PBS/T.
- Dilute primary antibody in PBS/T with 1% BSA. *Add 100 µl/well primary antibody.
- 13. Incubate at 37°C for 120 minutes.
- 14. 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 µl/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 polyclonal antibody: 1:1000

**Secondary antibody:

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