

#9273 Store at -20°C

Akt Control Cell Extracts



✓ Controls for 10 Western mini-blot

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For Research Use Only. Not For Use In Diagnostic Procedures.

Product Includes	Product #	Quantity
Akt Control Cell Extracts (Jurkat +Calyculin A)	48217	200 ul
Akt Control Cell Extracts (Jurkat + LY294002)	66376	200 ul

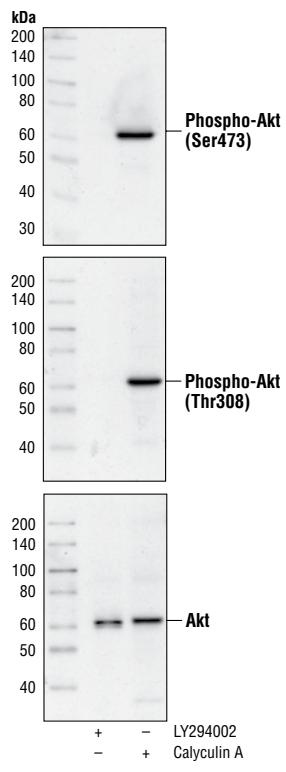
Background: Akt, also referred to as PKB or Rac, plays a critical role in controlling survival and apoptosis (1-3). This protein kinase is activated by insulin and various growth and survival factors to function in a wortmannin-sensitive pathway involving PI3 kinase (2,3). Akt is activated by phospholipid binding and activation loop phosphorylation at Thr308 by PDK1 (4) and by phosphorylation within the carboxy terminus at Ser473. The previously elusive PDK2 responsible for phosphorylation of Akt at Ser473 has been identified as mammalian target of rapamycin (mTor) in a rapamycin-insensitive complex with rictor and Sin1 (5,6). Akt promotes cell survival by inhibiting apoptosis by phosphorylating and inactivating several targets, including Bad (7), forkhead transcription factors (8), c-Raf (9) and caspase-9. PTEN phosphatase is a major negative regulator of the PI3 kinase/Akt signaling pathway (10). LY294002 is a specific PI3 kinase inhibitor (11).

Another essential Akt function is the regulation of glycogen synthesis through phosphorylation and inactivation of GSK-3 α and β (12,13). Akt may also play a role in insulin stimulation of glucose transport (12).

In addition to its role in survival and glycogen synthesis, Akt is involved in cell cycle regulation by preventing GSK-3 β mediated phosphorylation and degradation of cyclin D1 (14) and by negatively regulating the cyclin dependent kinase inhibitors p27 Kip (15) and p21 Waf1 (16). Akt also plays a critical role in cell growth by directly phosphorylating mTOR in a rapamycin-sensitive complex containing raptor (17). More importantly, Akt phosphorylates and inactivates tuberlin (TSC2), an inhibitor of mTOR within the mTOR-raptor complex (18). Inhibition of mTOR stops the protein synthesis machinery due to inactivation of its effector, p70 S6 kinase and activation of the eukaryotic initiation factor 4E binding protein 1 (4E-EP1), an inhibitor of translation (18,19).

Description: Phosphorylated Akt Cell Extracts: Total cell extracts from Jurkat cells, serum starved overnight and then treated with Calyculin A (CST #9902) to preserve their activated Akt state, serve as a positive control. Supplied in SDS Sample Buffer.

Nonphosphorylated Akt Cell Extracts: Total cell extracts from Jurkat cells, serum starved overnight and then treated with



Western blot analysis of extracts from Jurkat cells, Calyculin A or LY294002-treated, using Phospho-Akt (Ser473) Antibody #9271 (upper), Phospho-Akt (Thr308) Antibody # 9275 (middle) or Akt Antibody # 9272 (lower).

50 μ M LY294002 (CST #9901) for one hour, serve as a negative control. Supplied in SDS Sample Buffer.

Directions for Use: Boil for 3 minutes prior to use. Load 20 μ l of phosphorylated and nonphosphorylated Akt extracts per lane.

Storage: Supplied in SDS Sample Buffer: 62.5 mM Tris-HCl (pH 6.8 at 25°C), 2% w/v SDS, 10% glycerol, 50 mM DTT, 0.01% w/v bromophenol blue or phenol red. Store at -20°C, or at -80°C for long-term storage.

Please visit www.cellsignal.com for a complete listing of recommended companion products.

Background References:

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IMPORTANT: For Western blots, incubate membrane with diluted antibody in 5% w/v BSA, 1X TBS, 0.1% Tween-20 at 4°C with gentle shaking, overnight.

Applications Key: W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation IF—Immunofluorescence F—Flow cytometry E—ELISA E-P—ELISA Peptide
Species Cross-Reactivity Key: H—human M—mouse R—rat Hm—hamster Mk—monkey Mi—mink C—chicken Dm—D. melanogaster X—Xenopus Z—zebra fish B—bovine
 Dg—dog Pg—pig Sc—S. cerevisiae All—all species expected Species enclosed in parentheses are predicted to react based on 100% homology.