Store at -20C

Phospho-Akt Substrate (RXRXXS*/T*) (23C8D2) Rabbit mAb



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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: Reactivity: Sensitivity: Source/Isotype: WB, E-P All Endogenous Rabbit

Product Usage Application Dilution Information Western Blotting 1:1000 Peptide ELISA (DELFIA) 1:1000

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than **Storage** 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

Phospho-Akt Substrate (RXRXXS*/T*) (23C8D2) Rabbit mAb recognizes endogenous proteins containing Specificity / Sensitivity

> phospho-Ser/Thr preceded by Arg at positions -5 and -3 in a manner largely independent of the surrounding amino acid sequence. Minor cross-reactivity is observed for proteins that contain phospho-Ser/Thr preceded by Arg at position -3 only. No cross-reactivity is observed with the corresponding

nonphosphorylated sequences or with other phospho-Ser/Thr-containing motifs.

Source / Purification Monoclonal antibody is produced by immunizing animals with an Akt substrate peptide library.

An important class of kinases, referred to as Arq-directed kinases or AGC-family kinases, includes cAMP-**Background**

dependent protein kinase (PKA), cGMP-dependent protein kinase (PKG), protein kinase C, Akt, and RSK. These kinases share a substrate specificity characterized by Arg at position -3 relative to the

phosphorylated Ser or Thr (1,2). Akt plays a central role in mediating critical cellular responses including cell growth and survival, angiogenesis, and transcriptional regulation (3-5). While a number of Akt substrates are known (such as GSK-3, Bad, and caspase-9) many important substrates await discovery. Akt phosphorylates substrates only at Ser/Thr in a conserved motif characterized by Arg at positions -5 and -3 (6). Phospho-Akt substrate-specific antibodies from Cell Signaling Technology are powerful tools for investigating the regulation of phosphorylation by Akt and other Arg-directed kinases, as well as for high

throughput kinase drug discovery.

1. Montminy, M. (1997) Annu Rev Biochem 66, 807-22. **Background References**

2. Pearson, R.B. and Kemp, B.E. (1991) Methods Enzymol 200, 62-81.

3. Marte, B.M. and Downward, J. (1997) Trends Biochem Sci 22, 355-8.

4. Jiang, B.H. et al. (2000) Proc Natl Acad Sci USA 97, 1749-53.

5. Scheid, M.P. and Woodgett, J.R. (2000) Curr Biol 10, R191-4.

6. Alessi, D.R. et al. (1996) FEBS Lett 399, 333-8.

Species Reactivity Species reactivity is determined by testing in at least one approved application (e.g., western blot).

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, **Western Blot Buffer**

0.1% Tween® 20 at 4°C with gentle shaking, overnight.

WB: Western Blotting E-P: Peptide ELISA (DELFIA) **Applications Key**

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster **Cross-Reactivity Key**

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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