

Small Molecules

AKT Inhibitor VIII

PI3K/AKT pathway inhibitor; Inhibits AKT1, AKT2, and AKT3

Catalog # 72942
72944

1 mg
10 mg



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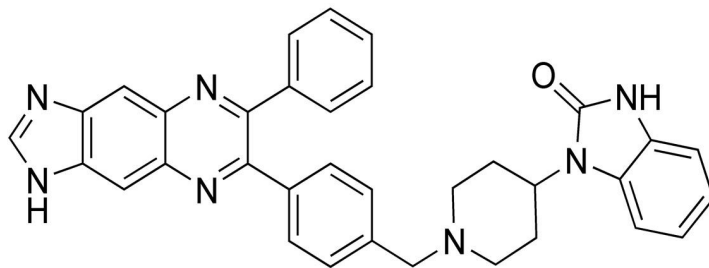
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Product Description

AKT Inhibitor VIII is a cell permeable, allosteric inhibitor of all three forms of the kinase AKT (AKT1, AKT2 and AKT3) with IC₅₀ values of 58, 210 and 2,200 nM respectively (Lindsley et al.; Calleja et al.). It displays good selectivity against a panel of 70 other kinases with micromolar inhibition against some kinases, for example, calcium/calmodulin-dependent protein kinase 1 and smooth muscle myosin light-chain kinase (Logie et al.).

Molecular Name:	AKT Inhibitor VIII
Alternative Names:	AKTi-1/2, AKT 1/2 inhibitor
CAS Number:	612847-09-3
Chemical Formula:	C ₃₄ H ₂₉ N ₇ O
Molecular Weight:	551.7 g/mol
Purity:	≥ 98%
Chemical Name:	3-[1-[[4-(7-phenyl-3H-imidazo[4,5-g]quinoxalin-6-yl)phenyl]methyl]piperidin-4-yl]-1H-benzimidazol-2-one
Structure:	



Properties

Physical Appearance:	A crystalline solid
Storage:	Product stable at -20°C as supplied. Protect from prolonged exposure to light. For product expiry date, please contact techsupport@stemcell.com .
Solubility:	· DMSO ≤ 25 mM For example, to prepare a 10 mM stock solution in DMSO, resuspend 1 mg in 181 µL of DMSO.

Prepare stock solution fresh before use. Information regarding stability of small molecules in solution has rarely been reported, however, as a general guide we recommend storage in DMSO at -20°C. Aliquot into working volumes to avoid repeated freeze-thaw cycles. The effect of storage of stock solution on compound performance should be tested for each application.

Compound has low solubility in aqueous media. For use as a cell culture supplement, stock solution should be diluted into culture medium immediately before use. Avoid final DMSO concentration above 0.1% due to potential cell toxicity.

Published Applications

METABOLISM

- Reduces insulin-dependent gene repression in liver cells leading to reduced insulin sensitivity (Logie et al.).

CANCER RESEARCH

- Sensitizes prostate tumor and cervical carcinoma cells to apoptotic stimuli (DeFeo-Jones et al.).
- Blocks mitosis and inhibits migration of HeLa cells (Jo et al.).

References

Calleja V et al. (2009) Role of a novel PH-kinase domain interface in PKB/Akt regulation: structural mechanism for allosteric inhibition. PLoS Biol 7(1): e17.

DeFeo-Jones D et al. (2005) Tumor cell sensitization to apoptotic stimuli by selective inhibition of specific Akt/PKB family members. Mol Cancer Ther 4(2): 271–279.

Jo H et al. (2011) Deactivation of Akt by a small molecule inhibitor targeting pleckstrin homology domain and facilitating Akt ubiquitination. Proc Natl Acad Sci U S A 108(16): 6486–91.

Lindsley CW et al. (2005) Allosteric Akt (PKB) inhibitors: discovery and SAR of isozyme selective inhibitors. Bioorg Med Chem Lett 15(3): 761–4.

Logie L et al. (2007) Characterization of a protein kinase B inhibitor in vitro and in insulin-treated liver cells. Diabetes 56(9): 2218–27.

Related Small Molecules

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This product is hazardous. Please refer to the Safety Data Sheet (SDS).

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