Roscovitine

1 mg



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orders@cellsignal.com

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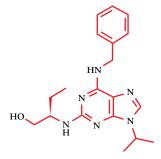
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For Research Use Only. Not For Use In Diagnostic Procedures.

Background: Roscovitine is a cell permeable reversible selective inhibitor of cyclin-dependent kinases CDK1 (cdc2), CDK2 and CDK5 (1). A purine analog, this drug competes for the binding site of ATP in the catalytic cleft. Treatment of cultured cells with roscovitine can cause cell cycle arrest or apoptosis (1-4). The IC₅₀ for cdc2 activity is 0.65 µM in vitro (1).

Molecular Formula: $C_{19}H_{28}N_6O$

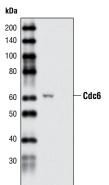


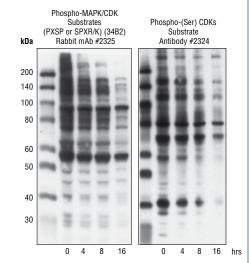
Molecular Weight: 354.46 g/mol

Directions for Use: Roscovitine [(R) stereoisomer] is supplied as 1 mg powder. Store at or below -20°C. Before use, dissolve powder in 143 µl DMSO or MeOH to make a 20 mM stock solution. The suggested working concentration is 20 μM in tissue culture medium. Treat cells for 4-24 hours depending on the individual experiment. Store solution at -20°C.

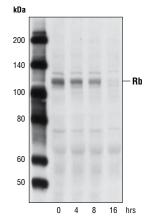
Background References:

- (1) Meijer, L. et al. (1997) Eur J Biochem 243, 527-36.
- (2) Whittaker, S.R. et al. (2007) Cell Cycle 6, 3114-31.
- (3) Dey, A. et al. (2008) Cell Death Differ 15, 263-73.
- (4) Wojciechowski, J. et al. (2003) Int J Cancer 106, 486-95.





Western blot analysis of extracts from HeLa cells treated with Roscovitine (20 µM) for the indicated times using Phospho-MAPK/CDK Substrates (PXSP or SPXR/K) (34B2) Rabbit mAb #2325 (left) or Phospho-(Ser) CDKs Substrate Antibody #2324



Western blot analysis of extracts from HeLa cells treated with Roscovitine (20 µM) for the indicated times using an antibody

■ Western blot analysis of extracts from U2OS cells, untreated or Roscovitine-treated (20 μM for 6 hours) using Cdc6 (C42F7)

Storage: Store lyophilized or in solution at -20°C, desiccated. In lyophilized form, the chemical is stable for 24 months. Once in solution, use within 3 months to prevent loss of potency. Aliquot to avoid multiple freeze/thaw cycles.

F—Flow cytometry E-P—ELISA-Peptide

Rabbit mAb #3387.