

Catalog # 10-2750 Toyocamycin

CAS# 606-58-6

7-Deaza-7-cyanoadenosine; 4-Amino-7-β-D-ribofuranosyl-7*H*-pyrrolo[2,3-*d*]pyrimidine-5-carbonitrile NSC 63701; NSC 99843; Neuro 000027; Unamycin B; Vengicide Lot # X106129

An adenosine analog which inhibits ribozyme self cleavage in mammalian cells, $EC_{50} = 0.4 \, \Box M$ (for expression of a luciferase reporter)¹. A potent inhibitor of ER stress-induced XBP1 mRNA splicing². It suppresses thapsigargin-, tunicamycin- and 2-deoxyglucose-induced XBP1 mRNA splicing in HeLa cells without affecting ATF6 and PERK activation. Although unable to inhibit IRE1 \Box phosphorylation, it prevented IRE1 \Box -induced XBP1 mRNA cleavage *in vitro*. It inhibits not only ER stress-induced but also constitutive activation of XBP1 expression in multiple myeloma cell lines as well as in primary patient samples². Displays synergistic effects with bortezomib. Inhibits unfolded protein response and induces apoptosis in pancreatic cancer cells³.

- 1) Yen et al. (2006), Identification of inhibitors of ribozyme self-cleavage in mammalian cells via high-throughput screening of chemical libraries; RNA, **12** 797
- 2) Ri et al. (2012), Identification of Toyocamycin, an agent cytotoxic for multiple myeloma cells, as a potent inhibitor of ER stress-induced XBP1 mRNA splicing; Blood Cancer J., 2 e79
- 3) Chien et al. (2014), Selective inhibition of unfolded protein response induces apoptosis in pancreatic cancer cells; Oncotarget, 5 4881

PHYSICAL DATA

Molecular Weight: 291.26 Molecular Formula: $C_{12}H_{13}N_5O_4$ Purity: 97% by TLC

NMR: (Conforms)

Solubility: DMSO (up to 25 mg/ml), moderately water soluble

Physical Description: White or off-white solid

Storage and Stability: Store as supplied, desiccated at -20°C for up to 2 years from the date of purchase. Solutions in

DMSO may be stored at -20°C for up to 3 months.

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