



LKT Laboratories, Inc.

Staurosporine

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Product Information

Product ID S7600

CAS No. 62996-74-1

Chemical Name

Synonym AM-2282

Formula $C_{28}H_{26}N_4O_3$

Formula Wt. 466.5

Melting Point 237-239°C

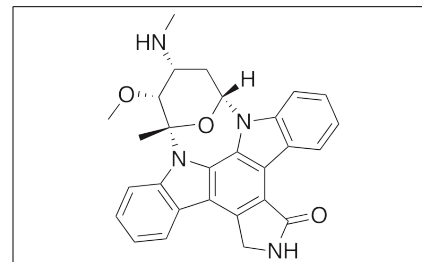
Purity ≥98%

Solubility Soluble in DMSO (50 mM),
ethanol, or methanol (2
mg/mL). Insoluble in water.

Store Temp -20°C

Ship Temp Ambient

Description Staurosporine is an alkaloid initially produced by *Streptomyces* that inhibits PKC and other protein kinases. Staurosporine is a precursor in the synthesis of kinase inhibitors K252c and PKC412. This compound exhibits anticancer, anti-parasitic, and anti-protozoan activities. Staurosporine induces apoptosis in hepatocarcinoma cells by decreasing levels of p38 MAPK, pERK, and DNA methyltransferase 3B (DNMT 3B) and increasing levels of pJNK. Staurosporine also causes cell death in *Trypanosoma*. This compound also inhibits mammalian RNA splicing.



Bulk quantities available upon request

Product ID	Size
S7600	1 mg
S7600	5 mg
S7600	25 mg
S7600	100 mg

References Zhao C, Yin P, Mei C, et al. Down-regulation of DNA methyltransferase 3B in staurosporine-induced apoptosis and its mechanism in human hepatocarcinoma cell lines. *Mol Cell Biochem.* 2013 Apr;376(1-2):111-9. PMID: 23397112.

Bruges G, Betancourt M, March M, et al. Apoptotic-like activity of staurosporine in axenic cultures of *Trypanosoma evansi*. *Rev Inst Med Trop Sao Paulo.* 2012 Mar-Apr;54(2):103-8. PMID: 22499424.

Aukema KG, Chohan KK, Plourde GL, et al. Small molecule inhibitors of yeast pre-mRNA splicing. *ACS Chem Biol.* 2009 Sep 18;4(9):759-68. PMID: 19634919.

Tanramluk D, Schreyer A, Pitt WR, et al. On the origins of enzyme inhibitor selectivity and promiscuity: a case study of protein kinase binding to staurosporine. *Chem Biol Drug Des.* 2009 Jul;74(1):16-24. PMID: 19519740.

Caution: This product is intended for laboratory and research use only. It is not for human or drug use.