# **FluoProbes**<sup>®</sup>



# **Staurosporine**

Potent and cell-permeable inhibitor of a wide variety of protein kinases

## **Product Description**

Name :	Staurosporine	H
	Synonyms: antibiotic AM-2282,	$ \subset \mathbb{P}^{\mathbb{N}} \neq 0 $
<b>Catalog Number :</b>	Staurosporine from Streptomyces staurosporeus FP-74146E 1 mg	
Structure :	C <sub>28</sub> H <sub>26</sub> N <sub>4</sub> O <sub>3</sub> CAS [62996-74-1]	
Molecular Weight :	MW= 466.53	
Melting point:	270°C	N N
Purity:	>99%	$\mathcal{V}^{\circ}\mathcal{A}$
Solubility:	Soluble in DMSO at 100 mg/mL; soluble in ethanol	H <sub>3</sub> C """ H
	at 2.5 mg/mL with warming; very poorly soluble in water; maximum solubility in plain water is estimated to be about 10-20 $\mu$ M; buffers, serum, or other additives may increase or decrease the aqueous solubility.	CH <sub>3</sub> O NHCH <sub>3</sub>

Storage: below -20 °C (M)

#### **Technical information**

• Biological activity

The staurosporine, isolated from *Streptomyces staurosporeus* is one of the most potent and widely used inhibitors of protein kinases. It is a potent inhibitor of protein kinase C (IC50 = 0.7 nM), but also protein kinase A (IC50 = 7-15 nM), and protein kinase G (IC50 = 8.5 nM). Other sources indicate for Protein Kinase C (IC<sub>50</sub> = 5 nM), PKA (IC50 = 15 nM), PKG (18 nM), CaMKII (20 nM), S6K (5 nM), MLCK (21 nM), SRC (6 nM), FGR (2 nM), LYN (20 nM) and SYK (16 nM).

In contrast it has a relatively low potency for ERK1 (1.5  $\mu$ M), CSK (2  $\mu$ M), IGF-IR (6.2  $\mu$ M), CK2 (19.5  $\mu$ M) and CK1 (>100 $\mu$ M).

It is cell permeable .



#### FT-74146E

Staurosporine induces apoptosis in human neuroblastoma cell lines and chick embryonic neurons<sup>1</sup>.

• Typical specifications:			
FORM:	Powder to crystalline solid or solid film at bottom of vial		
COLOR:	White to off-white to light yellow		
PURITY by HPLC:	>99%		
PURITY by TLC:	>99%		
ELEMENTAL ANALYSIS:	Element	Calculated	Found
$(C_{28}H_{26}N_4O_3 \bullet 1.0 C_4H_8)$	С	71.35%	71.29%
	Н	6.36%	6.29%
	Ν	10.40%	10.37%

#### **Guidelines for use**

#### • Protocols

Protocol may be found in the literature.

#### • Handling:

Toxic. May be carcinogenic. Wear gloves and mask when handling product. Protect from light.

#### References

Bijur GN, De Sarno P, Jope RS (2000) "Glycogen synthase kinase-3beta facilitates staurosporine- and heat shock-induced apoptosis. Protection by lithium" J. Biol. Chem. 275(11):7583-90.

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- Matsumoto, H. and Sasaki, Y. "Staurosporine, a protein kinase C inhibitor, interferes with proliferation of arterial smooth muscle cells." Biochem. Biophys. Res. Commun. 158: 105-109 (1989).

Meggio F, Donella Deana A, Ruzzene M et al. (1995) "Different susceptibility of protein kinases to staurosporine inhibition. Kinetic studies and molecular bases for the resistance of protein kinase CK2" Eur J Biochem. 234(1):317-22

Nakano H, Kobayashi E, Takahashi I, Tamaoki T, Kuzuu Y, Iba H (1987) "Staurosporine inhibits tyrosine-specific protein kinase activity of Rous sarcoma virus transforming protein p60" J. Antibiot. (Tokyo) 40(5):706-8

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Yamaki K, Ohuchi K (1999) "Participation of protein kinases in staurosporine-induced interleukin-6 production by rat peritoneal macrophages" Br J Pharmacol. 127(6):1309-16.

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#### **Related products**

- Protease inhibitors, <u>WT0900</u>
- Kinases substrates

- Live/Dead Mammalian Viability/Cytotoxicity Assay Kit, <u>BF4710</u>
- Caspase 8 Assay, BG4512

### **Ordering information**

Catalog size quantities and prices may be found at <u>http://www.interchim.com</u>. Please inquire for higher quantities (availability, shipment conditions).

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