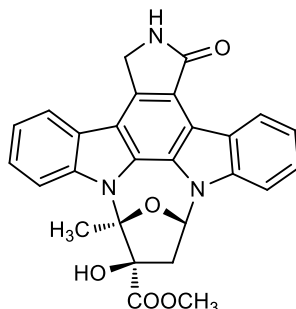


**Catalog # 10-2080**

**K-252a**

CAS# 99533-80-9

Lot # X101234



Potent pan-specific protein kinase inhibitor: PKA ( $K_i = 18$  nM), PKC ( $K_i = 25$  nM) PKG ( $K_i = 20$  nM)<sup>1</sup>. Inhibits CaMK ( $K_i = 1.8$  nM)<sup>2</sup> and phosphorylase kinase ( $IC_{50} = 1.7$  nM)<sup>3</sup> as well as other kinases<sup>4</sup>. Induces apoptosis<sup>5</sup>. Cell permeable.

- 1) Kase *et al.* (1987), *K-252 compounds, novel and potent inhibitors of protein kinase C and cyclic nucleotide-dependent protein kinases*; *Biochem. Biophys. Res. Commun.*, **142** 436
- 2) Hashimoto *et al.* (1991), *Potent and preferential inhibition of Ca<sup>2+</sup>/calmodulin-dependent protein kinase II by K252a and its derivative KT5926*; *Biochem. Biophys. Res. Commun.*, **181** 423
- 3) Berg *et al.* (1992), *K-252a inhibits nerve growth factor-induced trk proto-oncogene tyrosine phosphorylation and kinase activity*; *J. Biol. Chem.*, **267** 13
- 4) Ruegg *et al.* (1989), *Staurosporine, K-252 and UCN-01: potent but non-specific inhibitors of protein kinases*; *Trends Pharmacol. Sci.*, **10** 218
- 5) Mohri *et al.* (1999), *K252a induces cell cycle arrest and apoptosis by inhibiting Cdc2 and Cdc25c*; *Cancer Invest.*, **17** 391

**PHYSICAL DATA**

Molecular Weight:	467.49
Molecular Formula:	C <sub>27</sub> H <sub>21</sub> N <sub>3</sub> O <sub>5</sub>
Purity:	98% by TLC
	NMR: (Conforms)
Solubility:	DMSO (up to 100 mg/ml)
Physical Description:	White or off-white solid
Storage and Stability:	Store as supplied desiccated at -20°C for up to 1 year from the date of purchase. Solutions in DMSO may be stored at -20°C for up to 3 months.

Materials provided by Focus Biomolecules are for laboratory research use only and are not intended for human or veterinary applications.