

Catalog # 10-2621 Go-6983

CAS# 133053-19-7

3-[1-[3-(Dimethylamino)propyl]-5-methoxy-1H-indol-3-yl]-4-(1H-indol-3-yl)-1H-pyrrole-2,5-dione Lot # X101476

A potent and pan-specific, ATP-competitive protein kinase C inhibitor. PKC isotype IC₅₀s= 7, 7, 6, 10 and 60 nM for PKC α , β , γ , δ , ζ respectively. Suppresses stress induced HSP27 disassociation. Maintains rat embryonic stem cell pluripotency. Attenuates myocardial ischemia/reperfusion injury.

- 1) Gschwendt et al. (1996), Inhibition od protein kinas C mu by various inhibitors. Differentiation from protein kinas C isoenzymes; FEBS Lett., **392** 77
- 2) Kato et al. (2001), Protein kinase inhibitors can suppress stress-induced dissociation of Hsp27; Cell Stress Chaperones, **6** 16
- 3) Rajendran et al. (2013), Inhibition of protein kinas C signaling maintains rat embryonic stem cell pluripotency; J. Biol. Chem., **288** 24351
- 4) Young et al. (2005), Go 6983: a fast acting protein kinase C inhibitor that attenuates myocardial ischemia/reperfusion injury; Cardiovasc, Drug Rev., **23** 255

PHYSICAL DATA

Molecular Weight: 442.51

 $\begin{array}{ll} \text{Molecular Formula:} & C_{26}H_{26}N_4O_3 \\ \text{Purity:} & 98\% \text{ by HPLC} \end{array}$

NMR: (Conforms)

Solubility: Soluble in DMSO (up to 20 mg/ml)

Physical Description: Red-orange solid

Storage and Stability: Store as supplied, 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 month.

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