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# **ProductInformation**

## DEQUALINIUM ANALOG, C<sub>14</sub>-LINKER

Product Number **D 2064** Storage Temperature –20 °C

CAS #: Not available

Synonyms: 1,1'-(1,14-Tetradecanediyl)bis[4-amino-2-methylquinolinium] diiodide, DECA-14, Dequalinium-14

$$H_{2}N$$
 $N^{+}(CH_{2})_{14}$ 
 $H_{3}C$ 
 $NH_{2}$ 

### **Product Description**

Molecular Formula: C<sub>34</sub> H<sub>48</sub> N<sub>4</sub> I<sub>2</sub> Molecular Weight: 766.16 (anhydrous)

Supplied as light brown solid

Purity: 94% (HPLC)
Melting Point: 258-260 °C

Dequalinium (DECA) is a neurotoxin and potent antitumor agent that is selectively accumulated by mitochondria. It is a mitochondrial toxin that depletes cell energy (ATP), reduces the mitochondrial membrane potential, and increases free-radical production. The mitochondria also serve as a depot from which DECA is slowly released into the cytoplasm.

The activity of DECA is also due in part to its noncompetitive inhibition of protein kinase  $C\alpha$  (PKC $\alpha$ ), an enzyme that is involved in cell motility and cell adhesion. When activated by exposure to UV light, DECA covalently modifies and irreversibly inhibits PKC $\alpha$  with an IC $_{50}$  = 7-18  $\mu$ M. PKC inactivation also inhibits the motility and invasiveness of both human and murine melanoma cells *in vitro*.  $^{2,3}$ 

DECA analogues with long, saturated linkers ( $C_{12}$ ,  $C_{14}$ , or  $C_{16}$ ) exhibit enhanced inhibition of PKC in cell-free systems, which reached a plateau with the  $C_{14}$  linker ( $IC_{50} = 2.6 \mu M$ ). Studies with cultured metastatic melanoma cells demonstrated that photoactivated  $C_{12}$ -,  $C_{14}$ -, or  $C_{16}$ -DECA also inactivates intracellular PKC $\alpha$ .

### **Preparation Instructions**

Dequalinium analog,  $C_{14}$ -linker is soluble in DMSO at 22 mg/ml. It is slightly soluble in water at 0.7 mg/ml.

#### Storage/Stability

Store tightly sealed at -20 °C, protected from light.

#### References

- Chan, C.F., and Lin-Shiau, S.Y., Suramin prevents cerebellar granule cell-death induced by dequalinium. Neurochem. Int., 38, 135-143 (2001).
- Rotenberg, S.A. and Sun, X-G., Photoinduced inactivation of protein kinase C by dequalinium identifies the RACK-1-binding domain as a recognition site. J. Biol. Chem., 273, 2390-2395 (1998).
- 3. Sullivan, R.M., et al., Photo-induced inactivation of protein kinase  $C\alpha$  by dequalinium inhibits motility of murine melanoma cells. Mol. Pharmacol., **58**, 729-739 (2000).
- Qin, D., et al., Inhibition of Protein Kinase C alpha by Dequalinium Analogues: Dependence on Linker Length and Geometry. J. Med. Chem., 43, 1413-1417 (2000).

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