Product Data Sheet

Dequalinium Chloride

Cat. No.: HY-B0567 CAS No.: 522-51-0 Molecular Formula: $\mathsf{C}_{30}\mathsf{H}_{40}\mathsf{Cl}_2\mathsf{N}_4$

Molecular Weight: 527.57

Target: Potassium Channel; nAChR; Apoptosis; Bacterial; Parasite

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling; Apoptosis; Anti-infection

Storage: 4°C, sealed storage, away from moisture

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)

SOLVENT & SOLUBILITY

In Vitro

 $\rm H_2O$: 0.99 mg/mL (1.88 mM; ultrasonic and warming and adjust pH to 3 with 1 M HCL and heat to 60°C) DMSO: < 1 mg/mL (insoluble or slightly soluble)

| Preparing Stock Solutions | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg |
|------------------------------|-------------------------------|-------------|-----------|------------|
| | 1 mM | 1.8955 mL | 9.4774 mL | 18.9548 mL |
| | 5 mM | | | |
| | 10 mM | | | |

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

| Description | Dequalinium chloride is an Apamin (HY-P0256)-sensitive potassium channel selective blocker. Dequalinium chloride is a cationic, lipophilic mitochondrial poison. Dequalinium chloride is also an antagonist pf α 7 nAChR, and an anti-microbial antiseptic agent with a broad bactericidal and fungicidal activity ^{[1][2][3][4]} . |
|-------------|---|
| In Vitro | Dequalinium chloride blocks angiotensin II (100 nM)-evoked K ⁺ loss in guinea-pig hepatocytes, with an IC ₅₀ of 1.5 μ M ^[5] . Dequalinium (0-100 μ g/mL, 72 h) chloride inhibits cell growth in human Pca cell lines (PC3, DU145, LNCaP, MDA-PCA-2B), and induces cell apoptosis in PC3 cells (0.9 μ M, 4 h) ^[7] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |
| In Vivo | Dequalinium chloride shows a LD ₅₀ of 18.3 mg/kg in mice (i.p., a single time) ^[2] . Dequalinium chloride (2 mg/kg, i.p., daily for 10 days) inhibits the tumor growth of mouse bladder carcinoma MB49 ^[3] . Dequalinium chloride (2 mg/kg, s.c.) reduces Diisopropylfluorophosphate-induced tremors (organophosphate poisoning) in mice ^[6] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

REFERENCES

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Caution: Product has not been fully validated for medical applications. For research use only.

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