

产品名称 : Brompheniramine hydrogen maleate

同义词 : (±)-Brompheniramine

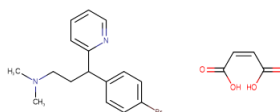
产品货号 : M16888

CAS Number : 980-71-2

分子式 : C₁₆H₁₉BrN₂·C₄H₄O₄

分子量 : 435.31

化学全名 : ——



产品描述 : Brompheniramine maleate is a histamine H1 receptors antagonist. (In Vitro):Brompheniramine (0.1-100 μM) blocks hERG K⁺ channels expressed in CHO cells in a concentration-dependent manner with an IC₅₀ of 0.90±0.14 μM, and reduced peak tail current amplitude measured at -60 mV (cells are depolarized for 2 s to +20 mV from a holding potential of -80 mV followed by a 3s repolarization back to -60 mV).Brompheniramine (1, 10 and 100 μM) significantly shortens the APD₅₀ and depresses the plateau phase on the action potential in guinea pig papillary muscle, as well as slightly prolongs the APD₉₀ in guinea pig papillary muscle at 10 and 100 μM.Brompheniramine (0.1-100 μM) inhibit the amplitude of the Ca²⁺ channel currents in rat ventricular myocytes by 14.1±1.1, 31.1±5.8, 38.0±3.8 and 90.2±3.7% at 0.1, 1, 10 and 100 μM, respectively.Brompheniramine blocks muscarinic cholinergic receptors in human chinese hamster ovary (CHO) cells.(In Vivo):Brompheniramine (0.3-3?μM; SC, single dosage) induces cutaneous analgesia in rats.

通路 : Endocrinology/Hormones

靶点 : 5-HT Receptor

受体 : HT

溶解度 : Ethanol: 18 mg/mL (41.34 mM); Water: 46 mg/mL warmed (105.67 mM); DMSO: 87 mg/mL (199.85 mM)

SMILES : CN(C)CCC(C1=CC=C(C=C1)Br)C2=CC=CC=N2.C(=C\C(=O)O)\C(=O)O

存储条件 : (-20℃)

稳定性 : ≥ 2 years

参考文献 :

1.Tardioli S, et al. J Phys Chem B. 2012 Jun 21;116(24):7033-9.