
Product Data Sheet

Product Name: Flupirtine

Cat. No.: GC36060

Chemical Properties

Cas. No. 56995-20-1

SMILES O=C(NC1=CC=C(N=C1N)NCC2=CC=C(C=C2)F)OCCFormula $C_{15}H_{17}FN_4O_2$ M.Wt 304.32

Solubility Soluble in DMSO Storage Store at -20°C

General tips For obtaining a higher solubility, please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution: ship with blue ice. All other available size: ship with RT, or blue ice upon request.

Structure

Background

Flupirtine is an activator of voltage-gated potassium channel 7 (K_v7/KCNQ).^{1,2,3} It induces relaxation of precontracted pulmonary arteries isolated from wild-type and serotonin transporter-overexpressing (SERT⁺) mice.² Flupirtine (30 mg/kg per day) decreases mean right ventricular pressure and right ventricular hypertrophy in hypoxia-induced and SERT⁺ mouse models of pulmonary arterial hypertension. It increases the paw withdrawal threshold in a rat model of streptozotocin-induced diabetic neuropathy when administered at a dose of 10 mg/kg and increases paw withdrawal latency in a rat model of carrageenan-induced paw inflammation when used in combination with morphine.³ Flupirtine also indirectly antagonizes NMDA receptors *via* its effects on potassium channels.^{1,4}

1.Devulder, J. Flupirtine in pain management: Pharmacological properties and clinical use *CNS Drugs* 25(10)867-881(2010) 2.Morecroft, I., Murray, A., Nilsen, M., et al. Treatment with the Kv7 potassium channel activator flupirtine is beneficial in two independent mouse models of pulmonary hypertension *Br. J. Pharmacol.* 157(7)1241-

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

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- 1249(2009) 3.Goodchild, C.S., Kolosov, A., Tucker, A.P., et al. Combination therapy with flupirtine and opioid: Studies in rat pain models *Pain Med.* 9(7)928-938(2008)
- 4.Kornhuber, J., Bleich, S., Wiltfang, J., et al. Flupirtine shows functional NMDA receptor antagonism by enhancing Mg²⁺ block via activation of voltage independent potassium channels. *J. Neural Transm. (Vienna)* 106(9-10)857-867(1999)

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