

## Cromakalim

## Chemical Properties

CAS No. : 94470-67-4

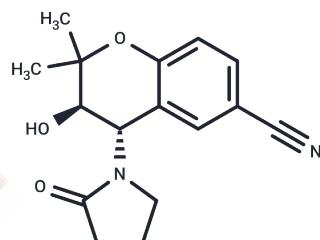
Formula: C16H18N2O3

Molecular Weight: 286.33

Appearance: Solid

Storage: keep away from direct sunlight, keep away from moisture

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year



## Biological Description

Description	Cromakalim (BRL 34915) is an ATP-dependent K(+) channel opener and a smooth muscle relaxant. Cromakalim has antiepileptic and anticonvulsant activity and may be useful in studies of asthma and disorders associated with vasodilation.
Targets(IC50)	Potassium Channel

## Solubility Information

Solubility	10% DMSO+40% PEG300+5% Tween 80+45% Saline: 2 mg/mL (6.98 mM), Sonication is recommended. H2O: Insoluble, DMSO: 50 mg/mL (174.62 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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## Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	3.4925 mL	17.4624 mL	34.9247 mL
5 mM	0.6985 mL	3.4925 mL	6.9849 mL
10 mM	0.3492 mL	1.7462 mL	3.4925 mL
50 mM	0.0698 mL	0.3492 mL	0.6985 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

Ostadhadi S, et al. Cromakalim, a Potassium Channel Opener, Ameliorates the Organophosphate and Carbamate-Induced Seizure in Mice. *Acta Med Iran.* 2018 Jan;56(1):14-20.

Sebille S, et al. Recent developments in the chemistry of potassium channel activators: the cromakalim analogs. *Curr Med Chem.* 2004 May;11(9):1213-22.

Shih CH, et al. Butyldenephthalide antagonizes cromakalim-induced systolic pressure reduction in conscious normotensive rats. *BMC Complement Altern Med.* 2015 Oct 5;15:344.

Spuler A, Lehmann-Horn F, Grafe P. Cromakalim (BRL 34915) restores in vitro the membrane potential of depolarized human skeletal muscle fibres. *Naunyn Schmiedebergs Arch Pharmacol.* 1989 Mar;339(3):327-31.

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