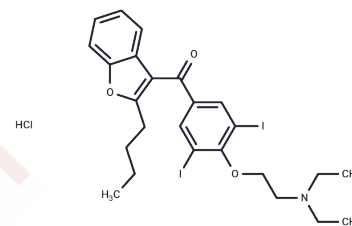


Amiodarone hydrochloride

Chemical Properties

CAS No. :	19774-82-4
Formula:	C ₂₅ H ₂₉ I ₂ NO ₃ ·HCl
Molecular Weight:	681.78
Appearance:	no data available
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year



Biological Description

Description	Amiodarone hydrochloride (Amiodarone HCl) is an antianginal and class III antiarrhythmic drug. It increases the duration of ventricular and atrial muscle action by inhibiting POTASSIUM CHANNELS and VOLTAGE-GATED SODIUM CHANNELS. There is a resulting decrease in heart rate and in vascular resistance.
Targets(IC50)	Adrenergic Receptor, Autophagy, Potassium Channel
In vitro	Amiodarone (AM) inhibits the intracellular conversion from thyroxine (T4) to triiodothyronine (T3) via 5'-deiodination (5'DI) without affecting the intracellular conversion from T4 to reverse T3 (rT3). 1.25-25 mg/kg Amiodarone in the AV node and in anaesthetized dogs resulted in a decrease in sinus rate, a prolongation of the AV node effective and functional occlusion of the AV node, as well as frequency-dependent conduction delays. 50 mg/kg daily for 3-4 weeks Amiodarone in rabbit ventricular myocytes resulted in a significant reduction in <i>i</i> K and <i>i</i> to current densities without affecting <i>I</i> CA and <i>I</i> K1 densities.
In vivo	Amiodarone penetrates deeply into the lipid matrix of the membrane and is released very slowly from cardiac tissue during washout. Amiodarone inhibits fast sodium channels as well as slow calcium channels. Amiodarone also has non-competitive antiarrhythmic effects and regulates thyroid function and phospholipid metabolism. 44-88 µM Amiodarone inhibits V _{max} of guinea pig papillary muscle without affecting normal hematopoietic stem cells. 44-88 µM Amiodarone inhibits V _{max} of guinea pig papillary muscle without affecting normal hematopoietic stem cells. Amiodarone inhibits V _{max} in papillary muscle without affecting the resting membrane potential, and this V _{max} inhibition is potentiated in a frequency- or use-dependent manner as is the case with class I antiarrhythmic drugs. 50-88 µM Amiodarone inhibits depolarization-induced spontaneous action potentials (aberrant automaticity) in ventricular myocardium and Purkinje fibers. 50-88 µM Amiodarone inhibits depolarization induced action potentials (aberrant automaticity) in human ventricular muscle.

Solubility Information

Solubility	DMSO: 7.5 mg/mL (11 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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A DRUG SCREENING EXPERT

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	1.4667 mL	7.3337 mL	14.6675 mL
5 mM	0.2933 mL	1.4667 mL	2.9335 mL
10 mM	0.1467 mL	0.7334 mL	1.4667 mL
50 mM	0.0293 mL	0.1467 mL	0.2933 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

Kodama I, et al. Cardiovasc Res,1997, 35(1), 13-29.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

This product is for Research Use Only· Not for Human or Veterinary or Therapeutic Use

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