Amiodarone hydrochloride

Cat. No.: HY-14188

CAS No.: 19774-82-4

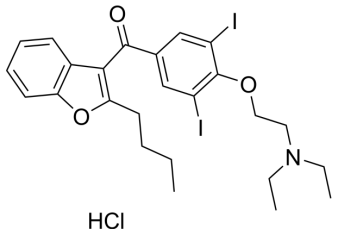
Molecular Formula: $C_{25}H_{30}ClI_2NO_3$

Molecular Weight: 681.77

Target: Potassium Channel; Autophagy

Pathway: Membrane Transporter/Ion Channel; Autophagy

Storage: 4°C, sealed storage, away from moisture and light
* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)



SOLVENT & SOLUBILITY

In Vitro

DMSO : 24.5 mg/mL (35.94 mM; Need ultrasonic and warming)

Preparing Stock Solutions	Mass / Solvent / Concentration	1 mg	5 mg	10 mg
	1 mM	1.4668 mL	7.3339 mL	14.6677 mL
	5 mM	0.2934 mL	1.4668 mL	2.9335 mL
	10 mM	0.1467 mL	0.7334 mL	1.4668 mL

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

In Vivo

1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
   Solubility: ≥ 2.5 mg/mL (3.67 mM); Clear solution
2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
   Solubility: ≥ 2.5 mg/mL (3.67 mM); Clear solution
3. Add each solvent one by one: 10% DMSO >> 90% corn oil
   Solubility: ≥ 2.5 mg/mL (3.67 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Amiodarone hydrochloride, a benzofuran-based Class III antiarrhythmic agent, inhibits WT outwardIhERG tails with an IC_{50} of -45 nM[1]. Amiodarone hydrochloride induces cell proliferation and myofibroblast differentiation via ERK1/2 and p38 MAPK signaling in fibroblasts[2]. Amiodarone hydrochloride can be used in the research of both supraventricular and ventricular arrhythmias[1].

In Vitro

Amiodarone blocks inward IhERG tails in a high K^+ external solution ($[K^+]e$) of 94 mM with an IC_{50} of 117.8 nM[1].
?Amiodarone (1 μM) blocks inwardIhERG by 68.8±6.1%, with concentration response data yielding IC_{50} and h values of 765.5±287.8 nM and 0.9±0.4 for T623A hERG[1].

?Amiodarone (1 μM) blocks inward IhERG with an IC_{50} and h values of 979.2±84.3 nM and 1.1±0.1 for S624A hERG[1].
?Amiodarone (1-6?μg/mL) induces human embryonic lung fibroblasts (HELFs) cell proliferation and PD98059 or SB203580 suppresses this effect[2].
?Amiodarone (1-6?ug/mL) does not induces HELFs cell apoptosis. Amiodarone (over 15?ug/mL) induces cell apoptosis[2].
?Amiodarone (1, 3 and 6?μg/mL;24?hours) induces α-SMA and vimentin mRNA and protein expression accompanied by increased phosphorylation of ERK1/2 and p38 MAPK[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Proliferation Assay[2]

Cell Line:	HELFs
Concentration:	1, 3 and 6 μg/mL
Incubation Time:	24 hours
Result:	Increased HELFs proliferation compared with the control group.

Western Blot Analysis[2]

Cell Line:	HELFs
Concentration:	1, 3 and 6 μg/mL
Incubation Time:	24 hours
Result:	α-SMA and vimentin were increased significantly in a dose-dependent manner.

RT-PCR[2]

Cell Line:	HELFs
Concentration:	1, 3 and 6 μg/mL
Incubation Time:	24 hours
Result:	Induced an increase of α-SMA and vimentin mRNA expression.

In Vivo

Amiodarone hydrochloride can be used in animal modeling to construct animal models of pulmonary fibrosis.

Long-term Amiodarone (90, and 180 mg/kg/day) treatment induces a dose-dependent remodeling of ion-channel expression that is correlated with the cardiac electrophysiologic effects of Amiodarone[3].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Ten-week-old male C57BL/6 mice[3]
Dosage:	30, 90, and 180 mg/kg/day
Administration:	Treated orally; for 6 weeks
Result:	Mice treated with 90 and 180 mg/kg/day had decreased body and heart weights, although their heart weight-to-body weight ratios were not significantly different from sham. 6-week treatment induced a decrease in plasma triiodothyronine and an increase in reverse triiodothyronine. This effect reached significance for the 90 and 180 but not for the 30 mg/kg/day dose groups.

CUSTOMER VALIDATION

- Cell. 2022 Dec 8;185(25):4801-4810.e13.
- Ecotoxicol Environ Saf. 2021 Apr 1;212:111991.
- Biotechnol Bioeng. 2021 Sep 3.
- Viruses. 2021 Jun 28;13(7):1255.
- Front Bioeng Biotechnol. 2022 Mar 17;10:826093.

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REFERENCES

[1]. Yihong Zhang,et al. Interactions between amiodarone and the hERG potassium channel pore determined with mutagenesis and in silico docking. Biochem Pharmacol. 2016 Aug 1;113:24-35.

[2]. Sabrina Le Bouter, et al. Long-term amiodarone administration remodels expression of ion channel transcripts in the mouse heart. Circulation. 2004 Nov 9;110(19):3028-35.

[3]. Jie Weng, et al. Amiodarone induces cell proliferation and myofibroblast differentiation via ERK1/2 and p38 MAPK signaling in fibroblasts. Biomed Pharmacother. 2019 Jul;115:108889.

Tel: 609-228-6898 Fax: 609-228-5909 E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA