

RH1

## Chemical Properties

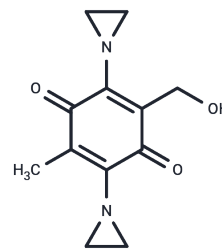
CAS No. : 221635-42-3

Formula: C<sub>12</sub>H<sub>14</sub>N<sub>2</sub>O<sub>3</sub>

Molecular Weight: 234.25

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

Actual storage temperature shall be subject to the COA.



## Biological Description

Description	RH1 (NSC 697726) is a bioreductive anticancer compound that exhibits dose-dependent biphasic effects in vitro, inducing apoptosis at higher doses and senescence at lower doses.
Targets(IC50)	Apoptosis,Others
In vitro	RH1 causes apoptosis in NQ16 cells in a time- and concentration-dependent manner. Treatment of NQ16 cells with RH1 (50 and 100 nM) for 60 and 120 minutes resulted in a significant (p< 0.05) increase in cross-linked DNA [1].
In vivo	RH1 exhibited dose-dependent antitumor activity against NQ16 tumors grown in thymus-free mice. A small dose of RH1 (0.1 mg/kg) also resulted in a significant reduction in tumor size in treated mice compared to controls. Treatment of mice with NQ16 tumors with RH1 (0.4 mg/kg and 0.2 mg/kg) resulted in a significant reduction in tumor size between the treated and control groups as early as 5 days after the end of the treatment period [2].

## Solubility Information

Solubility	DMSO: 1 mg/mL (4.27 mM),Sonication and heating are recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
------------	---

### Preparing Stock Solutions

---

	1mg	5mg	10mg
1 mM	4.2689 mL	21.3447 mL	42.6894 mL
5 mM	0.8538 mL	4.2689 mL	8.5379 mL
10 mM	0.4269 mL	2.1345 mL	4.2689 mL
50 mM	0.0854 mL	0.4269 mL	0.8538 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.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

### Reference

Park MT, et al. The anti-tumour compound, RH1, causes mitochondria-mediated apoptosis by activating c-Jun N-terminal kinase. Br J Pharmacol. 2011 Jun;163(3):567-85.

Dehn DL, et al. Development of a new isogenic cell-xenograft system for evaluation of NAD(P)H:quinone oxidoreductase-directed antitumor quinones: evaluation of the activity of RH1. Clin Cancer Res. 2004 May 1;10(9):3147-55.

**Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins**

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

Tel:781-999-4286 E\_mail:info@targetmol.com Address:34 Washington Street,Wellesley Hills,MA 02481