Inhibitors

Carubicin hydrochloride

Cat. No.: HY-B2171A CAS No.: 52794-97-5 Molecular Formula: $\mathsf{C}_{26}\mathsf{H}_{28}\mathsf{CINO}_{10}$

549.95 Molecular Weight: Target: **Apoptosis** Pathway: **Apoptosis**

Storage: 4°C, sealed storage, away from moisture

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

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 62.5 mg/mL (113.65 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.8183 mL	9.0917 mL	18.1835 mL
	5 mM	0.3637 mL	1.8183 mL	3.6367 mL
	10 mM	0.1818 mL	0.9092 mL	1.8183 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: ≥ 2.08 mg/mL (3.78 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (3.78 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Carubicin hydrochloride is a microbially-derived compound. Carubicin hydrochloride is an effective inhibitor of VHLdefective (VHL-/-) CCRCC cell proliferation. Carubicin hydrochloride also induces apoptosis by a mechanism independent of $p53 \ or \ hypoxia-inducible \ factor \ HIF2. \ Carubicin \ hydrochloride \ has \ the \ potential \ for \ the \ research \ of \ cancer \ diseases^{[1][2]}.$

REFERENCES

[1]. Woldemichael GM, et al. Carminomycin I is an apoptosis inducer that targets the Golgi complex in clear cell renal carcinoma cells. Cancer Res. 2011 Jan 1;71(1):134-42.

[2]. Tevyashova AN, et al. Carminomycin, 14-hydroxycarminomycin and its novel carbohydrate derivatives potently kill human tumor cells and their multidrug resistant variants. J Antibiot (Tokyo). 2004 Feb;57(2):143-50.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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