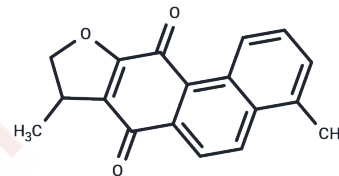


Dihydroisotanshinone I

Chemical Properties

| | |
|-------------------|--|
| CAS No. : | 20958-18-3 |
| Formula: | C ₁₈ H ₁₄ O ₃ |
| Molecular Weight: | 278.31 |
| Storage: | Store at low temperature Powder: -20°C for 3 years In solvent: -80°C for 1 year <small>Actual storage temperature shall be subject to the COA.</small> |



Biological Description

| | |
|---------------|---|
| Description | Dihydroisotanshinone I is a natural product isolated from the dried roots of <i>Salvia miltiorrhiza</i> , a traditional Chinese medicine, and exhibits various biological activities. Dihydroisotanshinone I alleviates menadione-induced liver injury through antioxidant effects such as scavenging free radicals and inhibiting lipid peroxidation; it also inhibits the migration of prostate cancer cells and induces cancer cell apoptosis. |
| Targets(IC50) | Apoptosis,Free radical scavengers,Lipid |
| In vitro | Dihydroisotanshinone I induces cell cycle G1 arrest by inhibiting the expression of cyclin D1, cyclin E1, CDK2, CDK4, CDK6, p-Rb, E2F1 and SKP2, and suppresses the proliferation of human osteosarcoma 143B cells in a dose- and time-dependent manner. Dihydroisotanshinone I induces apoptosis via activating caspase-3, caspase-8, caspase-9 and cleaving PARP, and inhibits cell migration by down-regulating VCAM-1 and ICAM-1. |
| In vivo | In xenograft tumor models, Dihydroisotanshinone I inhibits the proliferation of Hela human cervical cancer cells. It enhances radiation-induced apoptosis and significantly inhibits proliferation in Hela cells mainly by prolonging radiation-induced G2/M arrest and reducing the expression levels of Bcl-XL, Bcl-2 and HPV E6 protein. |

Solubility Information

| | |
|------------|---|
| Solubility | DMSO: 5.00 mg/mL (17.97 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble) |
|------------|---|

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|------------|------------|
| 1 mM | 3.5931 mL | 17.9656 mL | 35.9312 mL |
| 5 mM | 0.7186 mL | 3.5931 mL | 7.1862 mL |
| 10 mM | 0.3593 mL | 1.7966 mL | 3.5931 mL |
| 50 mM | 0.0719 mL | 0.3593 mL | 0.7186 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

Chen X, et al. 15,16-dihydrotanshinone I induces apoptosis and inhibits the proliferation, migration of human osteosarcoma cell line 143B in vitro[J]. *Anti-cancer agents in medicinal chemistry*. 2015, 15(999): 1234-1242.

F W, et al. Blockade of TNF- α -induced NF- κ B signaling pathway and anti-cancer therapeutic response of dihydrotanshinone I.[J]. *International Immunopharmacology*. 2015, 28(1): 764-772.

Ye Y, et al. Combination treatment with dihydrotanshinone I and irradiation enhances apoptotic effects in human cervical cancer by HPV E6 down-regulation and caspases activation[J]. *Molecular & Cellular Biochemistry*. 2012, 363 (1-2): 191-202.

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