**Pirarubicin**

Cat. No.: HY-13725  
CAS No.: 72496-41-4  
Molecular Formula: C₃₂H₃₇NO₁₂  
Molecular Weight: 627.64  
Target: Topoisomerase; Autophagy; Bacterial; Antibiotic  
Pathway: Cell Cycle/DNA Damage; Autophagy; Anti-infection  
Storage: 4°C, protect from light  
* The compound is unstable in solutions, freshly prepared is recommended.

**SOLVENT & SOLUBILITY**

<table>
<thead>
<tr>
<th>Preparing Stock Solutions</th>
<th>Solvent Concentration</th>
<th>Mass</th>
<th>1 mg</th>
<th>5 mg</th>
<th>10 mg</th>
</tr>
</thead>
<tbody>
<tr>
<td>DMSO</td>
<td>1 mM</td>
<td>1.5933 mL</td>
<td>7.9664 mL</td>
<td>15.9327 mL</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5 mM</td>
<td>0.3187 mL</td>
<td>1.5933 mL</td>
<td>3.1865 mL</td>
<td></td>
</tr>
<tr>
<td></td>
<td>10 mM</td>
<td>0.1593 mL</td>
<td>0.7966 mL</td>
<td>1.5933 mL</td>
<td></td>
</tr>
</tbody>
</table>

In Vitro: DMSO : 50 mg/mL (79.66 mM; Need ultrasonic)

**BIOLOGICAL ACTIVITY**

**Description**  
Pirarubicin is an anthracycline antibiotics, acts as a topoisomerase II inhibitor, and is a widely used for treatment of various cancers, in particular, solid tumors.

**IC₅₀ & Target**  
Topoisomerase II

**In Vitro**  
Pirarubicin is a topoisomerase II inhibitor[^1]. Pirarubicin shows inhibitory activities against M5076 and Ehrlich cells, with IC₅₀ s of 0.366 and 0.078 μM, respectively. The cytotoxicity of Pirarubicin toward M5076 cells is lower than toward Ehrlich cells, and this is due to the much lower expression of topoisomerase II in M5076 cells than in Ehrlich cells[^2]. Pirarubicin (2.5, 5, 10 μg/mL) significantly induces autophagy in a dose dependent manner in bladder cancer (T24, EJ, 5637, J82 and UM-UC-3) cells. Furthermore, Pirarubicin (5 μg/mL) induces apoptosis through inhibition of mTOR/p70S6K/4E-BP1 in bladder cancer cells, and this effect is enhanced by inhibition of autophagy[^3].

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

**In Vivo**  
Pirarubicin (18 mg/kg, i.v.) significantly elevates serum level of BNP, CK-MB, CTnT, LDH, and MDA compared with those in

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[^1]: Reference 1
[^2]: Reference 2
[^3]: Reference 3
the control group in acute cardiac toxicity rats. Pirarubicin also lowers heart rate, and depresses R-wave voltage, and prolongation of QT intervals in the acute cardiac toxicity model[4].

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**PROTOCOL**

**Cell Assay**[3]

MTS is used to analyze cell survival. Briefly, cells are plated in 96-well plates in triplicate at $2 \times 10^3$ cells per well and cultured in growth medium. Then cells are treated with pirarubicin at different concentrations (2.5 μg/mL, 5 μg/mL, 10 μg/mL) for 24 h. MTS reagent (5 mg/mL) is added and incubated at 37°C for 4 h. The absorbance is monitored at 490 nm using a microplate reader[3].

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**Animal Administration**[4]

An acute cardiac toxicity model is established by a single dose of 18 mg/kg pirarubicin through the caudal vein injection. Thirty-six rats are randomized equally to six groups: normal control, cardiac injury (THP) model, dexrazoxane (180 mg/kg), low-dose rutin (25 mg/kg), middle-dose rutin (50 mg/kg), and high-dose rutin (100 mg/kg). Rats in the rutin-treated group are administered different doses of rutin and CMC-Na for 7 days by gavage and a single dose of 18 mg/kg pirarubicin through caudal vein injection. Rats in the dexrazoxane-treated group receive sodium carboxymethylcellulose (CMC-Na) by gavage for six days. 40 mg/kg dexrazoxane is then administered to rats by intraperitoneal injection and 18 mg/kg pirarubicin is administered by caudal vein injection on day 7. Rats in the THP model group receive CMC-Na by gavage for seven days, followed by pirarubicin 18 mg/kg through the caudal vein injection on day 7. Rats in the normal control group receive CMC-Na by gavage for seven days, followed by saline through caudal vein injection on day 7[4].

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**CUSTOMER VALIDATION**

- Cancer Drug Resist. 2020 Sep 12.

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**REFERENCES**


