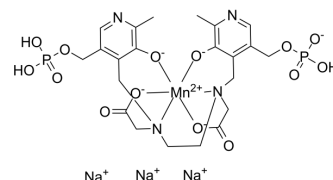


Mangafodipir trisodium

Cat. No.:	HY-B0993
CAS No.:	140678-14-4
Molecular Formula:	C ₂₂ H ₂₇ MnN ₄ Na ₃ O ₁₄ P ₂
Molecular Weight:	757.32
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)



SOLVENT & SOLUBILITY

In Vitro	H ₂ O : 33.33 mg/mL (44.01 mM; Need ultrasonic)																			
Preparing Stock Solutions	<table border="1"> <thead> <tr> <th rowspan="2">Solvent Concentration</th> <th colspan="3">Mass</th> </tr> <tr> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> </thead> <tbody> <tr> <td>1 mM</td> <td>1.3204 mL</td> <td>6.6022 mL</td> <td>13.2045 mL</td> </tr> <tr> <td>5 mM</td> <td>0.2641 mL</td> <td>1.3204 mL</td> <td>2.6409 mL</td> </tr> <tr> <td>10 mM</td> <td>0.1320 mL</td> <td>0.6602 mL</td> <td>1.3204 mL</td> </tr> </tbody> </table>	Solvent Concentration	Mass			1 mg	5 mg	10 mg	1 mM	1.3204 mL	6.6022 mL	13.2045 mL	5 mM	0.2641 mL	1.3204 mL	2.6409 mL	10 mM	0.1320 mL	0.6602 mL	1.3204 mL
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Please refer to the solubility information to select the appropriate solvent.																				
In Vivo	1. Add each solvent one by one: PBS Solubility: 100 mg/mL (132.04 mM); Clear solution; Need ultrasonic																			

BIOLOGICAL ACTIVITY

Description	Mangafodipir trisodium (MnDPDP), hepatocellular-specific contrast agent, is an efficacious inhibitor of CIPN (chemotherapy-induced peripheral neuropath) and other conditions caused by cellular oxidative stress. Mangafodipir trisodium shows no negative interference with the tumoricidal activity of chemotherapy ^{[1][2]} .
In Vitro	Mangafodipir trisodium (40, 200 and 1000 μM) inhibits the effect of HGrC cell viability caused by H ₂ O ₂ ^[3] . Mangafodipir trisodium (200 and 1000 μM) attenuates the apoptosis caused by cisplatin in HGrC cells ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	Mangafodipir trisodium (10 mg/kg; i.p., once) shows protective effects against ovarian damage caused by cisplatin and paclitaxel in mice ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Female CD-1(ICR) Mice with cisplatin- and paclitaxel-induced ovarian damage ^[3]
Dosage:	10 mg/kg
Administration:	Intraperitoneal injection; 10 mg/kg, once
Result:	Attenuated the loss of primordial follicles, reduction of secondary follicles and increase of antral follicles. Prevented the change of primary follicles caused by cisplatin and paclitaxel. Reduced the increasing of cleaved caspase-3 levels caused by cisplatin and paclitaxel.

REFERENCES

- [1]. Karlsson JOG, et al. Mangafodipir a Selective Cytoprotectant - with Special Reference to Oxaliplatin and Its Association to Chemotherapy-Induced Peripheral Neuropathy (CIPN). *Transl Oncol.* 2017 Aug;10(4):641-649.
- [2]. Wang C. Mangafodipir trisodium (MnDPDP)-enhanced magnetic resonance imaging of the liver and pancreas. *Acta Radiol Suppl.* 1998;415:1-31.
- [3]. Qin Y, et al. Protective effects of mangafodipir against chemotherapy-induced ovarian damage in mice. *Reprod Biol Endocrinol.* 2018 Oct 27;16(1):106.
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Caution: Product has not been fully validated for medical applications. For research use only.

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