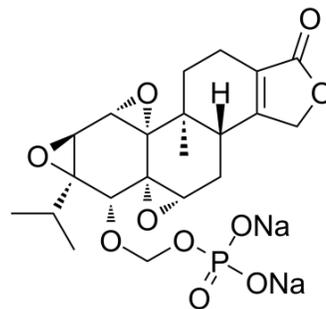


## Minnelide

Cat. No.:	HY-124584
CAS No.:	1254702-87-8
Molecular Formula:	C <sub>21</sub> H <sub>25</sub> Na <sub>2</sub> O <sub>10</sub> P
Molecular Weight:	514.37
Target:	Apoptosis
Pathway:	Apoptosis
Storage:	Please store the product under the recommended conditions in the COA.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Minnelide is a prodrug of triptolide that shows potent <b>antitumor</b> activity in a number of tumor types, particularly in pancreatic cancer. Minnelide causes <b>apoptotic</b> <sup>[1]</sup> .								
<b>In Vitro</b>	<p>Minnelide (0-200 nM; 48 hours) shows significantly decreased cell viability in pancreatic cancer cell lines after treatment in the presence, but not in the absence, of phosphatase<sup>[2]</sup>.</p> <p><b>Cell Viability Assay</b><sup>[2]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>Pancreatic cancer cell line: S2-013, MIA PaCa-2, S2-VP10, and Panc-1 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.100 nM, 200 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> <tr> <td>Result:</td> <td>Decreased cell viability of in vitro.</td> </tr> </table>	Cell Line:	Pancreatic cancer cell line: S2-013, MIA PaCa-2, S2-VP10, and Panc-1 cells	Concentration:	0.100 nM, 200 nM	Incubation Time:	48 hours	Result:	Decreased cell viability of in vitro.
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Concentration:	0.100 nM, 200 nM								
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Result:	Decreased cell viability of in vitro.								
<b>In Vivo</b>	<p>Minnelide (injection intraperitoneally; 0.1-0.6 mg/kg; once daily or twice daily) leads to a marked decrease in tumor weight and volume at the end of treatment and increases survival in orthotopic model of pancreatic cancer with MIA PaCa-2-derived human pancreatic tumors<sup>[2]</sup>.</p> <p>Minnelide (injection intraperitoneally; 0.42 mg/kg; once daily; 28 days) prevents locoregional spread and leads to a decrease in average tumor weight in a xenograft model of pancreatic cancer with metastatic S2-013 cells<sup>[2]</sup>.</p> <p>Minnelide (injection intraperitoneally; 0.42 mg/kg, 0.21 mg/kg; once daily) causes tumor regression and tumors from Minnelide-treated animals showed fibrosis and the presence of pyknotic nuclei in human pancreatic cancer xenografts in SCID mice<sup>[2]</sup>.</p> <table border="1"> <tr> <td><b>Animal Model:</b></td> <td>Orthotopic model of pancreatic cancer with MIA PaCa 2-derived human pancreatic tumors in athymic nude mice<sup>[2]</sup></td> </tr> <tr> <td><b>Dosage:</b></td> <td>0.1-0.6 mg/kg</td> </tr> <tr> <td><b>Administration:</b></td> <td>Injection intraperitoneally; 0.1-0.6 mg/kg; once daily or twice daily</td> </tr> <tr> <td><b>Result:</b></td> <td>Prevented pancreatic tumor growth in vivo.</td> </tr> </table>	<b>Animal Model:</b>	Orthotopic model of pancreatic cancer with MIA PaCa 2-derived human pancreatic tumors in athymic nude mice <sup>[2]</sup>	<b>Dosage:</b>	0.1-0.6 mg/kg	<b>Administration:</b>	Injection intraperitoneally; 0.1-0.6 mg/kg; once daily or twice daily	<b>Result:</b>	Prevented pancreatic tumor growth in vivo.
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<b>Animal Model:</b>	Xenograft model of pancreatic cancer with metastatic S2-013 cell line in athymic nude mice <sup>[2]</sup>
<b>Dosage:</b>	0.42 mg/kg
<b>Administration:</b>	Injection intraperitoneally; 0.42 mg/kg; once daily
<b>Result:</b>	Prevented extensive spread from the primary site of injection.
<b>Animal Model:</b>	Human pancreatic cancer xenografts in SCID mice <sup>[2]</sup>
<b>Dosage:</b>	0.21 mg/kg, 0.42 mg/kg
<b>Administration:</b>	Injection intraperitoneally; 0.42 mg/kg; once daily
<b>Result:</b>	Reduced tumor burden in human xenografts from patients.

## REFERENCES

- [1]. Noel P et al. Triptolide and Its Derivatives as Cancer Therapies. Trends Pharmacol Sci. 2019 May;40(5):327-341.
- [2]. Chugh R, et al. A preclinical evaluation of Minnelide as a therapeutic agent against pancreatic cancer. Sci Transl Med. 2012 Oct 17;4(156):156ra139.

**Caution: Product has not been fully validated for medical applications. For research use only.**

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