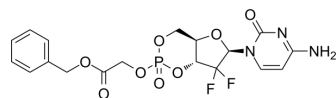


## Antitumor agent-91

Cat. No.:	HY-149927
Molecular Formula:	C <sub>18</sub> H <sub>18</sub> F <sub>2</sub> N <sub>3</sub> O <sub>8</sub> P
Molecular Weight:	473.32
Target:	Others
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Antitumor agent-91 is a cyclic phosphate ester derivative with anticancer and anti-proliferative activities. Antitumor agent-91 has potential application in human castration resistant prostate cancer and pancreatic cancer <sup>[1]</sup> .																
<b>In Vitro</b>	<p>Antitumor agent-91 (Compound 18c) has anti-tumor activity against MM1. S, BxPC-3, MP-2, 22Rv1, MV4-11, Jeko-1 and HUH7 cells with IC<sub>50</sub> values of 3.6 nM, 16.5 nM, 12.5 nM, 10.6 nM, 5.4 nM, 19.2 nM and 16.2 nM, respectively<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>BxPC-3 cells.</td> </tr> <tr> <td>Concentration:</td> <td>25 nM, 50 nM and 100 nM.</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h.</td> </tr> <tr> <td>Result:</td> <td>Increased the expression of P21.</td> </tr> </table> <p>Cell Cycle Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>BxPC-3 cells.</td> </tr> <tr> <td>Concentration:</td> <td>25 nM, 50 nM and 100 nM.</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h.</td> </tr> <tr> <td>Result:</td> <td>Induced cell S phase arrest.</td> </tr> </table>	Cell Line:	BxPC-3 cells.	Concentration:	25 nM, 50 nM and 100 nM.	Incubation Time:	24 h.	Result:	Increased the expression of P21.	Cell Line:	BxPC-3 cells.	Concentration:	25 nM, 50 nM and 100 nM.	Incubation Time:	24 h.	Result:	Induced cell S phase arrest.
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<b>In Vivo</b>	<p>Antitumor agent-91 (Compound 18c) (50 mg/kg; i.p.; twice a week for four weeks) significantly inhibits tumor growth in a castrated male Balb/c nude mouse model implanted with 22Rv1 tumor<sup>[1]</sup>.</p> <p>Pharmacokinetic (PK) parameters of Antitumor agent-91 in Beagle Dogs<sup>[1]</sup></p> <table border="1"> <thead> <tr> <th>Route</th> <th>Dose (mg/kg)</th> <th>AUC<sub>0-∞</sub> (h•ng/mL)</th> <th>T<sub>1/2</sub> (h)</th> <th>CL (mL/h/kg)</th> <th>C<sub>max</sub> (ng/mL)</th> </tr> </thead> <tbody> <tr> <td>Intravenous</td> <td>1</td> <td>520.24±79.49</td> <td>0.12±0.046</td> <td>1952±295</td> <td>3600.3±325.59</td> </tr> </tbody> </table>	Route	Dose (mg/kg)	AUC <sub>0-∞</sub> (h•ng/mL)	T <sub>1/2</sub> (h)	CL (mL/h/kg)	C <sub>max</sub> (ng/mL)	Intravenous	1	520.24±79.49	0.12±0.046	1952±295	3600.3±325.59				
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injection

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## REFERENCES

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[1]. Zhang L, et al. Design, Synthesis, and Anti-Cancer Evaluation of Novel Cyclic Phosphate Prodrug of Gemcitabine. J Med Chem. 2023 Mar 23;66(6):4150-4166.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA