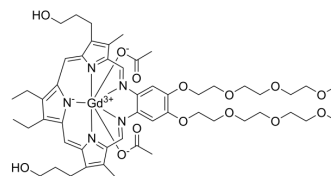


Motexafin gadolinium

Cat. No.:	HY-16336
CAS No.:	246252-06-2
Molecular Formula:	C ₅₂ H ₇₂ GdN ₅ O ₁₄
Molecular Weight:	1148.4
Target:	Others
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Motexafin gadolinium (PCI 0120) is a potent antitumor agent. Motexafin gadolinium is a specific texaphyrin in complex with gadolinium(III). Motexafin gadolinium increases intracellular ROS production. Motexafin gadolinium enhances sensitivity of tumor cells to ionizing radiation ^{[1][2][3]} .								
In Vitro	<p>Motexafin gadolinium (50 μM; 22 h) leads to dichlorofluorescein acetate (DCFA) oxidation in A549 cells^[1].</p> <p>Motexafin gadolinium (100 μM; 4, 12 h) increases intracellular ROS production and apoptosis (24 h) when combined with Zn (100 μM), and ascorbate (100 μM) in Ramos lymphoma cells^[2].</p> <p>Motexafin gadolinium (0.01-100 μM) dose-dependent decrease the expression of p53 protein in HF-1 cells when combined with Zn (100 μM), and ascorbate (100 μM)^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HF-1 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.01-100 μM, with Zn (100 μM), and ascorbate (100 μM)</td> </tr> <tr> <td>Incubation Time:</td> <td>5 h</td> </tr> <tr> <td>Result:</td> <td>Elicited a dose-dependent decrease in p53 in HF-1 cells, does not reduce p53 message in Ramos cells.</td> </tr> </table>	Cell Line:	HF-1 cells	Concentration:	0.01-100 μM, with Zn (100 μM), and ascorbate (100 μM)	Incubation Time:	5 h	Result:	Elicited a dose-dependent decrease in p53 in HF-1 cells, does not reduce p53 message in Ramos cells.
Cell Line:	HF-1 cells								
Concentration:	0.01-100 μM, with Zn (100 μM), and ascorbate (100 μM)								
Incubation Time:	5 h								
Result:	Elicited a dose-dependent decrease in p53 in HF-1 cells, does not reduce p53 message in Ramos cells.								

REFERENCES

- [1]. Magda D, et al. Motexafin gadolinium: a novel redox active drug for cancer therapy. *Semin Cancer Biol.* 2006 Dec;16(6):466-76.
- [2]. Singh AT, et al. Motexafin gadolinium enhances p53-Mdm2 interactions, reducing p53 and downstream targets in lymphoma cell lines. *Anticancer Res.* 2010 Apr;30(4):1131-6.
- [3]. Evens AM, et al. The novel expanded porphyrin, motexafin gadolinium, combined with [90Y]ibritumomab tiuxetan for relapsed/refractory non-Hodgkin's lymphoma: preclinical findings and results of a phase I trial. *Clin Cancer Res.* 2009 Oct 15;15(20):6462-71.

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

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