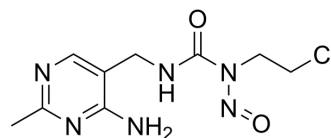


Nimustine

Cat. No.:	HY-13703
CAS No.:	42471-28-3
Molecular Formula:	C ₉ H ₁₃ ClN ₆ O ₂
Molecular Weight:	272.69
Target:	Apoptosis
Pathway:	Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Nimustine is a nitrosourea alkylating agent. Nimustine induces cell apoptosis, and activates DNA damage response and MAPK signaling. Nimustine shows anti-cancer effects, it can be used for the research of cancer ^{[1][2]} .																
In Vitro	<p>Nimustine (50 μM; 72-120 h) causes cell death by inducing cell apoptosis^[1]. Nimustine (50 μM; 24-96 h) activates the DNA damage response pathway^[1]. Nimustine (50 μM; 24-120 h) activates MAPK signaling in glioma cells^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Apoptosis Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>LN-229 cell line</td> </tr> <tr> <td>Concentration:</td> <td>50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72-120 hours</td> </tr> <tr> <td>Result:</td> <td>Time-dependently induced apoptosis.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>LN-229 and glioma cell lines</td> </tr> <tr> <td>Concentration:</td> <td>50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24-120 hours</td> </tr> <tr> <td>Result:</td> <td>Induced cleavage of caspase-8 and -9 and the effector caspase-3. Increased phosphorylation of ERK kinase and H2AX. Decreased phosphorylation of JNK</td> </tr> </table>	Cell Line:	LN-229 cell line	Concentration:	50 μM	Incubation Time:	72-120 hours	Result:	Time-dependently induced apoptosis.	Cell Line:	LN-229 and glioma cell lines	Concentration:	50 μM	Incubation Time:	24-120 hours	Result:	Induced cleavage of caspase-8 and -9 and the effector caspase-3. Increased phosphorylation of ERK kinase and H2AX. Decreased phosphorylation of JNK
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In Vivo	<p>Nimustine (15 four times a week and 30 mg/kg twice with an interval of 2 weeks; i.v.) effectively inhibits tumor growth and the higher dose is more effective^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Female C3H/HeN mice with solid FM3A tumors^[2]</td> </tr> </table>	Animal Model:	Female C3H/HeN mice with solid FM3A tumors ^[2]														
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Dosage:	15 and 30 mg/kg
Administration:	Intravenous injection; 15 (4 times a week) and 30 mg/kg (twice with an interval of 2 weeks)
Result:	The intermittent large-dose injections resulted in better inhibition of tumor growth than did the fractionated small-dose injections.

REFERENCES

- [1]. Tomicic MT, et al. Apoptosis induced by temozolomide and nimustine in glioblastoma cells is supported by JNK/c-Jun-mediated induction of the BH3-only protein BIM. *Oncotarget*. 2015 Oct 20;6(32):33755-68.
- [2]. Shimizu F, et al. Effects of combined treatment with nimustine hydrochloride and radiation on solid FM3A tumor in mice. *Jpn J Cancer Res*. 1987 Jul;78(7):756-62.
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Caution: Product has not been fully validated for medical applications. For research use only.

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