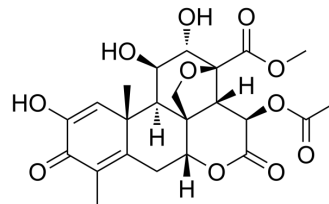


Dehydrobruceine B

Cat. No.:	HY-N10313
CAS No.:	53730-90-8
Molecular Formula:	C ₂₃ H ₂₆ O ₁₁
Molecular Weight:	478.45
Target:	Bcl-2 Family; Keap1-Nrf2; Apoptosis
Pathway:	Apoptosis; NF-κB
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Dehydrobruceine B, a quassinoid, can be isolated from Brucea javanica. Dehydrobruceine B shows a synergistic effect with Cisplatin (HY-17394) to induce apoptosis via mitochondrial method. Dehydrobruceine B increases apoptosis-inducing factor (AIF) and Bax expression and suppresses Keap1-Nrf2 ^[1] .																		
IC₅₀ & Target	Bax	Bcl-xL	Bcl-2																
In Vitro	<p>Dehydrobruceine B (1 μM; 48 h) and Cisplatin (3-18 μM; 48 h) combination synergizes the cytotoxic and apoptotic effects in A549 cells^[1].</p> <p>Dehydrobruceine B (1 μM; 24 h) induces excessive generation of intracellular ROS in Cisplatin (3 μM, 6 μM; 24 h)-treated A549 cells^[1].</p> <p>Dehydrobruceine B (1 μM; 24 h) increases depolarization of mitochondrial membrane potential (MMP) and translocation of cytochrome c in Cisplatin (3 μM, 6 μM; 24 h)-treated A549 cells^[1].</p> <p>Dehydrobruceine B (1 μM; 24 h) enhances the change of anti-apoptotic and pro-apoptotic protein levels in A549 cells^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>A549 cells</td> </tr> <tr> <td>Concentration:</td> <td>1 μM; with or without Cisplatin</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> <tr> <td>Result:</td> <td>Upregulated the protein level of Bax, while downregulated the levels of Bcl-2 and Bcl-xL. Enhanced caspase activation and PARP cleavage.</td> </tr> </table> <p>Apoptosis Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>A549 cells</td> </tr> <tr> <td>Concentration:</td> <td>1 μM; with or without Cisplatin</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibited cell viability with 9 μM and 18 μM Cisplatin, respectively. Induced cell apoptosis with 3 μM and 6 μM Cisplatin, respectively.</td> </tr> </table>			Cell Line:	A549 cells	Concentration:	1 μM; with or without Cisplatin	Incubation Time:	48 hours	Result:	Upregulated the protein level of Bax, while downregulated the levels of Bcl-2 and Bcl-xL. Enhanced caspase activation and PARP cleavage.	Cell Line:	A549 cells	Concentration:	1 μM; with or without Cisplatin	Incubation Time:	48 hours	Result:	Inhibited cell viability with 9 μM and 18 μM Cisplatin, respectively. Induced cell apoptosis with 3 μM and 6 μM Cisplatin, respectively.
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Immunofluorescence^[1]

Cell Line:	A549 cells
Concentration:	1 μ M; with or without 3 μ M and 6 μ M Cisplatin, respectively
Incubation Time:	24 hours
Result:	Resulted apoptosis-inducing factor (AIF) translocated from cytosol into nucleus dramatically in the co-treatment condition.

REFERENCES

[1]. Huang Z, et al. Dehydrobruceine B enhances the cisplatin-induced cytotoxicity through regulation of the mitochondrial apoptotic pathway in lung cancer A549 cells. Biomed Pharmacother. 2017 May;89:623-631.

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

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