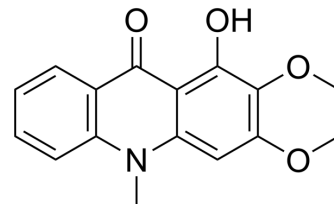


Arborinine

Cat. No.:	HY-128912
CAS No.:	5489-57-6
Molecular Formula:	C ₁₆ H ₁₅ NO ₄
Molecular Weight:	285.29
Target:	Histone Demethylase
Pathway:	Epigenetics
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Arborinine is a potent and orally active LSD1 inhibitor. Arborinine increases the expression of H3K4me1/2, H3K9me1/2, E-cad protein and decreases the expression of UBE2O protein level. Arborinine induces cell cycle arrest at S phase. Arborinine shows antitumor activity ^{[1][2]} .																
IC₅₀ & Target	LSD1																
In Vitro	<p>Arborinine (0, 5, 15, 30 μM; 48 h) increases the expression of H3K4me1/2, H3K9me1/2, E-cad protein^[1]. Arborinine (20 μM) decreases the expression of UBE2O protein level in 786O cells^[1]. Arborinine (20 μM) induces cell cycle arrest at the S phase and inhibits cell apoptosis^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>786O, A498, 769P, Caki1, OSRC2 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48, 72 h</td> </tr> <tr> <td>Result:</td> <td>Showed antiproliferative activity with IC₅₀s of 30.62, 39.09, 15.67, 31.42, 30.35 μM at 48 h, 20.92, 27.01, 14.94, 30.26, 17.37 μM at 72 h for 786O, A498, 769P, Caki1, OSRC2 cells, respectively.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>786O cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 5, 15, 30 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Increased the expression of H3K4me1/2 and H3K9me1/2 protein in a dose-dependent manner.</td> </tr> </table> <p>Cell Cycle Analysis^[1]</p>	Cell Line:	786O, A498, 769P, Caki1, OSRC2 cells	Concentration:	0-100 μM	Incubation Time:	48, 72 h	Result:	Showed antiproliferative activity with IC ₅₀ s of 30.62, 39.09, 15.67, 31.42, 30.35 μM at 48 h, 20.92, 27.01, 14.94, 30.26, 17.37 μM at 72 h for 786O, A498, 769P, Caki1, OSRC2 cells, respectively.	Cell Line:	786O cells	Concentration:	0, 5, 15, 30 μM	Incubation Time:	48 h	Result:	Increased the expression of H3K4me1/2 and H3K9me1/2 protein in a dose-dependent manner.
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	Cell Line:	786O and A498 cells
	Concentration:	20 μ M
	Incubation Time:	
	Result:	Induced increased population in S phase and decreased population in G1 phase, significantly inhibited both early and late apoptosis of ccRCC cells
In Vivo	Arborinine (40, 80 mg/kg; p.o.; for 21 days) shows antitumor activity in mouse ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	BALB/c nude mice (SGC-7901 cells and SGC-7901/ADR cells) ^[2]
	Dosage:	40, 80 mg/kg
	Administration:	P.o.; for 21 days
	Result:	Inhibited the growth of tumors while unchanged the body weights.

REFERENCES

- [1]. Feng C, et al. Arborinine from Glycosmis parva leaf extract inhibits clear-cell renal cell carcinoma by inhibiting KDM1A/UBE2O signaling. Food Nutr Res. 2022 Sep 16;66.
- [2]. Chu Y, et al. Arborinine, a potential LSD1 inhibitor, inhibits epithelial-mesenchymal transition of SGC-7901 cells and adriamycin-resistant gastric cancer SGC-7901/ADR cells. Invest New Drugs. 2021 Jun;39(3):627-635.

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