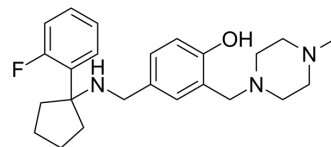


ARN5187

Cat. No.:	HY-103691
CAS No.:	1287451-26-6
Molecular Formula:	C ₂₄ H ₃₂ FN ₃ O
Molecular Weight:	397.53
Target:	Autophagy; Apoptosis
Pathway:	Autophagy; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	ARN5187 is a lysosomotropic REV-ERBβ ligand with a dual inhibitory activity toward REV-ERB-mediated transcriptional regulation and autophagy. ARN5187 shows lysosomotropic potency and cytotoxicity. ARN5187 induces apoptosis ^{[1][2]} .																
IC₅₀ & Target	REV-ERBβ ^[1]																
In Vitro	<p>ARN5187 (compound 1) (0-100 μM; 48 h) shows cytotoxicity with EC₅₀ of 23.5 μM in BT-474 cells and IC₅₀ of 30.14 μM, >100 μM for BT-474 and HMEC cells, respectively^{[1][2]}.</p> <p>ARN5187 (0-100 μM) activates the RevRE reporter in a concentration-dependent manner in HEK-293 cells^[1].</p> <p>ARN5187 (25, 50 μM) is a lysosomotropic-independent REV-ERB antagonistic activity^[1].</p> <p>ARN5187 (50 μM; 24 h) shows autophagy inhibition^[1].</p> <p>ARN5187 (50 μM; 2, 8, 24 h) effects autophagy formation and maturation^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>BT-474 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Showed cytotoxicity with EC₅₀ of 23.5 μM.</td> </tr> </table> <p>RT-PCR^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>BT-474 cells</td> </tr> <tr> <td>Concentration:</td> <td>25, 50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td></td> </tr> <tr> <td>Result:</td> <td>Significantly enhanced the expression of BMAL1, PER1 and PEPCK in a dose-dependent manner.</td> </tr> </table> <p>Western Blot Analysis^[1]</p>	Cell Line:	BT-474 cells	Concentration:	0-100 μM	Incubation Time:	48 h	Result:	Showed cytotoxicity with EC ₅₀ of 23.5 μM.	Cell Line:	BT-474 cells	Concentration:	25, 50 μM	Incubation Time:		Result:	Significantly enhanced the expression of BMAL1, PER1 and PEPCK in a dose-dependent manner.
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Cell Line:	BT-474 cells
Concentration:	50 μ M
Incubation Time:	24 h
Result:	Significantly increased the expression of α -LC3-II, α -p62, α -Cleaved PARP.

REFERENCES

- [1]. De Mei C, et al. Dual inhibition of REV-ERB β and autophagy as a novel pharmacological approach to induce cytotoxicity in cancer cells. *Oncogene*. 2015 May 14;34(20):2597-608.
- [2]. Torrente E, et al. Synthesis and in Vitro Anticancer Activity of the First Class of Dual Inhibitors of REV-ERB β and Autophagy. *J Med Chem*. 2015 Aug 13;58(15):5900-15.
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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

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