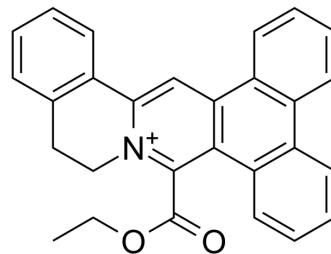


DC_C66

Cat. No.:	HY-100855
CAS No.:	108181-00-6
Molecular Formula:	C ₂₈ H ₂₂ NO ₂ ⁺
Molecular Weight:	404.48
Target:	Histone Methyltransferase
Pathway:	Epigenetics
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	DC_C66 is a cell-permeable, selective coactivator associated arginine methyltransferase 1 (CARM1) inhibitor with an IC ₅₀ of 1.8 μM. DC_C66 has a good selectivity for CARM1 against PRMT1 (IC ₅₀ =21 μM), PRMT6 (IC ₅₀ = 47μM), and PRMT5 ^[1] .								
IC₅₀ & Target	IC ₅₀ : 1.8 μM (CARM1) ^[1]								
In Vitro	<p>DC_C66 (6.25-100 μM; 24-72 hours) exhibits antiproliferation activity in several CARM1-associated cancer cell lines^[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>HELA, K562, MCF7 cells</td> </tr> <tr> <td>Concentration:</td> <td>6.25, 12.5, 25, 50, 100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24, 48, 72 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibits proliferation of cancer cells in a time-dependent and dose-dependent manner in HELA, K562, and MCF7 cells.</td> </tr> </table>	Cell Line:	HELA, K562, MCF7 cells	Concentration:	6.25, 12.5, 25, 50, 100 μM	Incubation Time:	24, 48, 72 hours	Result:	Inhibits proliferation of cancer cells in a time-dependent and dose-dependent manner in HELA, K562, and MCF7 cells.
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REFERENCES

[1]. Ye F, et al. Identification of Novel Inhibitors against Coactivator Associated Arginine Methyltransferase 1 Based on Virtual Screening and Biological Assays. Biomed Res Int. 2016;2016:7086390.

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

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