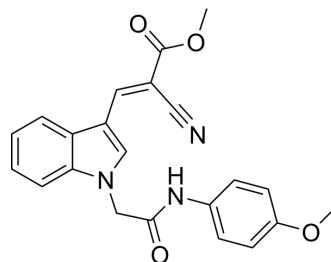


MCT1-IN-3

Cat. No.:	HY-148574
CAS No.:	2878360-80-4
Molecular Formula:	C ₂₂ H ₁₉ N ₃ O ₄
Molecular Weight:	389.4
Target:	Monocarboxylate Transporter; P-glycoprotein
Pathway:	Membrane Transporter/Ion Channel
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	MCT1-IN-3 is a monocarboxylate transporter 1 (MCT1) inhibitor with an IC ₅₀ value of 81.0 nM. MCT1-IN-3 has also significant inhibitivity against the multidrug transporter ABCB1. MCT1-IN-3 can be used for the research of cancer ^[1] .																
IC₅₀ & Target	IC ₅₀ : 81.0 nM (MCT1) ^[1] GI ₅₀ : 20 μM (A-549); 15.1 μM (MCF-7) ^[1]																
In Vitro	<p>MCT1-IN-3 (compound 24) has inhibitory activity for MCT1 with an IC₅₀ value of 81.0 nM^[1].</p> <p>MCT1-IN-3 has great antiproliferative activities against the MCT1-expressing cancer cell lines A-549 and MCF-7 with GI₅₀ values of 20 μM and 15.1 μM, respectively^[1].</p> <p>MCT1-IN-3 (5 μM; 24 h) leads to cancer cell cycle arrest as well as apoptosis^[1].</p> <p>MCT1-IN-3 (1.0, 2.0, 3.5, and 5.0 μM) shows to sensitize these cancer cells toward an antineoplastic agent^[1].</p> <p>MCT1-IN-3 has also significant inhibitory power against the multidrug transporter ABCB1 and shows to reverse ABCB1-mediated multidrug resistance (MDR)^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cycle Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>A-549 cells</td> </tr> <tr> <td>Concentration:</td> <td>5 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Caused a disruption of the cell cycle of A-549 cancer cells, indicated by a shift from the predominant G₀/G₁ phase.</td> </tr> </table> <p>Apoptosis Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>A-549 cells</td> </tr> <tr> <td>Concentration:</td> <td>5 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Increased the percentage of total apoptotic cells by a factor of 13 from 0.51 to 6.68%.</td> </tr> </table>	Cell Line:	A-549 cells	Concentration:	5 μM	Incubation Time:	24 h	Result:	Caused a disruption of the cell cycle of A-549 cancer cells, indicated by a shift from the predominant G ₀ /G ₁ phase.	Cell Line:	A-549 cells	Concentration:	5 μM	Incubation Time:	24 h	Result:	Increased the percentage of total apoptotic cells by a factor of 13 from 0.51 to 6.68%.
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REFERENCES

[1]. Sachin Puri, et al. Indole Derivatives as New Structural Class of Potent and Antiproliferative Inhibitors of Monocarboxylate Transporter 1 (MCT1; SLC16A1). J Med Chem. 2023 Jan 12;66(1):657-676.

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