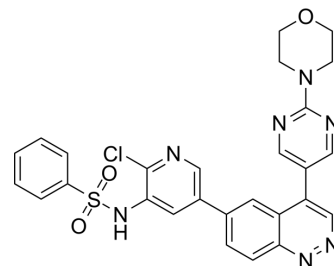


PI3K-IN-29

Cat. No.:	HY-144450
CAS No.:	2768005-77-0
Molecular Formula:	C ₂₇ H ₂₂ ClN ₇ O ₃ S
Molecular Weight:	560.03
Target:	PI3K; Akt
Pathway:	PI3K/Akt/mTOR
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	PI3K-IN-29 is a potent PI3K inhibitor. PI3K-IN-29 displays good inhibition potencies against U87MG, HeLa and HL60 cells with IC ₅₀ values of 0.264, 2.04 and 1.14 μM, respectively. PI3K-IN-29 inhibits PI3K/Akt pathway by inhibiting phosphorylation of Akt that is catalyzed by PI3K ^[1] .																
In Vitro	<p>PI3K-IN-29 (compound 25) (72 h) displays good inhibition potencies against U87MG, HeLa and HL60 cells with IC₅₀ values of 0.264, 2.04 and 1.14 μM, respectively^[1].</p> <p>PI3K-IN-29 (U87MG cells; 1, 5 μM; 1 h) inhibits PI3K/Akt pathway by inhibiting phosphorylation of Akt that is catalyzed by PI3K^[1].</p> <p>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>U87MG, HeLa, HepG2, A549, HL60, MCF7 cells</td> </tr> <tr> <td>Concentration:</td> <td></td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Displayed good inhibition potencies against U87MG, HeLa and HL60 cells with IC₅₀ values of 0.264, 2.04 and 1.14 μM, respectively.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>U87MG cells</td> </tr> <tr> <td>Concentration:</td> <td>1, 5 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>1 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited PI3K/Akt pathway by inhibiting phosphorylation of Akt that was catalyzed by PI3K.</td> </tr> </table>	Cell Line:	U87MG, HeLa, HepG2, A549, HL60, MCF7 cells	Concentration:		Incubation Time:	72 h	Result:	Displayed good inhibition potencies against U87MG, HeLa and HL60 cells with IC ₅₀ values of 0.264, 2.04 and 1.14 μM, respectively.	Cell Line:	U87MG cells	Concentration:	1, 5 μM	Incubation Time:	1 h	Result:	Inhibited PI3K/Akt pathway by inhibiting phosphorylation of Akt that was catalyzed by PI3K.
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REFERENCES

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

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