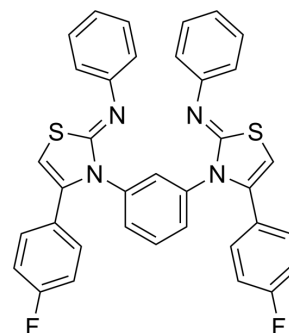


Pim-1 kinase inhibitor 9

Cat. No.:	HY-163119
Molecular Formula:	C ₃₆ H ₂₄ F ₂ N ₄ S ₂
Molecular Weight:	614.73
Target:	Pim
Pathway:	JAK/STAT Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Pim-1 kinase inhibitor 9 (compound 8b) is a selective inhibitor against Pim-1 kinase with IC ₅₀ value of 0.24 μM. Pim-1 kinase inhibitor 9 inhibits cell cycle of T47D at S phase. Pim-1 kinase inhibitor 9 reveals antitumor activity ^[1] .																	
IC₅₀ & Target	PIM1 0.24 μM (IC ₅₀)	PIM2 10.53 μM (IC ₅₀)																
In Vitro	<p>Pim-1 kinase inhibitor 9 inhibits the growth of breast cancer cells T47D with IC₅₀ values of 9.8 μM (8b, 48h) and 2.61 μM (8b, 96h)^[1].</p> <p>Pim-1 kinase inhibitor 9 inhibits the activities of Pim-1 kinase and Pim-2 kinase with IC₅₀ values of 0.24 μM (Pim-1 kinase) and 10.53 μM (Pim-2 kinase), respectively^[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>T47D</td> </tr> <tr> <td>Concentration:</td> <td>10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h and 96 h</td> </tr> <tr> <td>Result:</td> <td>Exhibited cytotoxicity activity to T47D with IC₅₀ value of 9.8 μM (48 h) and 2.61 μM (96h).</td> </tr> </table> <p>Cell Cycle Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>T47D</td> </tr> <tr> <td>Concentration:</td> <td>10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited cell cycle at S phase.</td> </tr> </table>		Cell Line:	T47D	Concentration:	10 μM	Incubation Time:	48 h and 96 h	Result:	Exhibited cytotoxicity activity to T47D with IC ₅₀ value of 9.8 μM (48 h) and 2.61 μM (96h).	Cell Line:	T47D	Concentration:	10 μM	Incubation Time:	48 h	Result:	Inhibited cell cycle at S phase.
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In Vivo	Pim-1 kinase inhibitor 9 (1 mg/kg/once weekly for 3 weeks; i.p.) inhibits the growth of Ehrlich solid tumours in mice by promoting caspase-3 expression and inhibiting the expression of VEGF ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.																	

Animal Model:	swiss albino mice/Ehrlich solid carcinoma model ^[1]
Dosage:	1 mg/kg
Administration:	i.p. injected once a week, for 3 weeks
Result:	Inhibited growth of Ehrlich solid tumours in mice.

REFERENCES

[1]. Al-Sanea MM, et al., Design, synthesis and cytotoxic evaluation of novel bis-thiazole derivatives as preferential Pim1 kinase inhibitors with in vivo and in silico study. J Enzyme Inhib Med Chem. 2023 Dec;38(1):2166936.

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

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