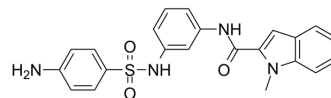


PHGDH-IN-2

Cat. No.:	HY-145860
Molecular Formula:	C ₂₂ H ₂₀ N ₄ O ₃ S
Molecular Weight:	420.48
Target:	Others
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	PHGDH-IN-2 is a potent and NAD ⁺ competitive PHGDH inhibitor with an IC ₅₀ of 5.2 μM. PHGDH-IN-2 inhibits the serine synthetic pathway in MDA-MB-468 cells. PHGDH-IN-2 inhibits the growth of PHGDH-dependent cancer cells ^[1] .								
IC₅₀ & Target	PHGDH ^[1]								
In Vitro	<p>PHGDH-IN-2 (compound C25) (10, 30 μM) inhibits the serine synthetic pathway in MDA-MB-468 cells^[1]. PHGDH-IN-2 (72 h) inhibits the growth of PHGDH-dependent cancer cells^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MDA-MB-468, NCI-H1975, HCC827, PC-9, ZR-75-1, MDA-MB-231 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>Inhibited the growth of PHGDH-dependent cancer cells.</td> </tr> </table>	Cell Line:	MDA-MB-468, NCI-H1975, HCC827, PC-9, ZR-75-1, MDA-MB-231 cells	Concentration:		Incubation Time:	72 h	Result:	Inhibited the growth of PHGDH-dependent cancer cells.
Cell Line:	MDA-MB-468, NCI-H1975, HCC827, PC-9, ZR-75-1, MDA-MB-231 cells								
Concentration:									
Incubation Time:	72 h								
Result:	Inhibited the growth of PHGDH-dependent cancer cells.								

REFERENCES

[1]. Zhang FM, et al. Discovery of PHGDH inhibitors by virtual screening and preliminary structure-activity relationship study. Bioorg Chem. 2022, 121:105705.

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

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