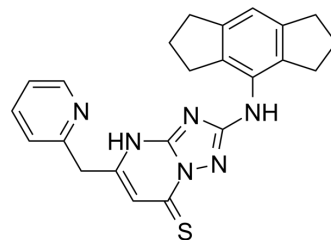


NDT-30805

Cat. No.:	HY-150965
Molecular Formula:	C ₂₃ H ₂₂ N ₆ S
Molecular Weight:	414.53
Target:	NOD-like Receptor (NLR)
Pathway:	Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	NDT-30805 is a selective NLRP3 inflammasome inhibitor. NDT-30805 is a triazolopyrimidinone derivative and inhibits IL-1 β release in PBMCs with an IC ₅₀ of 0.013 μ M. NDT-30805 can be used for the research of inflammation and innate immunity ^[1] .																	
IC₅₀ & Target	NLRP3	NLRP3 inflammasome																
In Vitro	<p>NDT-30805 (0-2 μM; 30 min) inhibits IL-1β release in PBMCs^[1].</p> <p>NDT-30805 (0-2 μM; 30 min) inhibits ASC speck formation in THP1-ASC-GFP cells^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Human PBMCs</td> </tr> <tr> <td>Concentration:</td> <td>0-2 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>30 min</td> </tr> <tr> <td>Result:</td> <td>Inhibited IL-1β release in PBMCs with an IC₅₀ of 0.013 μM.</td> </tr> </table> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>THP1-ASC-GFP cells</td> </tr> <tr> <td>Concentration:</td> <td>0-2 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>30 min</td> </tr> <tr> <td>Result:</td> <td>Inhibited ASC which is a component of NLRP3 inflammasome in PBMCs with an IC₅₀ of 0.034 μM.</td> </tr> </table>		Cell Line:	Human PBMCs	Concentration:	0-2 μ M	Incubation Time:	30 min	Result:	Inhibited IL-1 β release in PBMCs with an IC ₅₀ of 0.013 μ M.	Cell Line:	THP1-ASC-GFP cells	Concentration:	0-2 μ M	Incubation Time:	30 min	Result:	Inhibited ASC which is a component of NLRP3 inflammasome in PBMCs with an IC ₅₀ of 0.034 μ M.
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Concentration:	0-2 μ M																	
Incubation Time:	30 min																	
Result:	Inhibited ASC which is a component of NLRP3 inflammasome in PBMCs with an IC ₅₀ of 0.034 μ M.																	

REFERENCES

[1]. David H, et al. Discovery and Optimization of Triazolopyrimidinone Derivatives as Selective NLRP3 Inflammasome Inhibitors. ACS Medicinal Chemistry Letters. 2022.

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

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