**Proteins** 



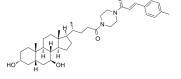
## Akt/NF-kB/MAPK-IN-1

Cat. No.: HY-149496 Molecular Formula:  $C_{38}H_{56}N_2O_4$ Molecular Weight: 604.86

Target: p38 MAPK; ERK; JNK; Interleukin Related; TNF Receptor; NO Synthase; COX Pathway: MAPK/ERK Pathway; Stem Cell/Wnt; Immunology/Inflammation; Apoptosis

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.



**Product** Data Sheet

## **BIOLOGICAL ACTIVITY**

Description	Akt/NF-κB/MAPK-IN-1 (compound 2m) is a potent and orally active inhibitor against NO (IC <sub>50</sub> =7.70 μM) with no significant toxicity. Akt/NF-κB/MAPK-IN-1 shows anti-inflammatory activity by inhibiting Akt/NF-κB and MAPK signaling pathways <sup>[1]</sup> .			
IC <sub>50</sub> & Target	IL-1β	IL-6	iNOS	COX-2
In Vitro	Akt/NF- $\kappa$ B/MAPK-IN-1 (compound 2m) (0-20 $\mu$ M, 24 h) significantly decreases the levels of TNF- $\alpha$ , IL-1 $\beta$ , IL-6 and PGE2, down-regulates the expression of iNOS and COX-2 in LPS (10 ng/mL)-induced RAW264.7 cells <sup>[1]</sup> . Akt/NF- $\kappa$ B/MAPK-IN-1 (0-20 $\mu$ M, 24 h) can block LPS-induced phosphorylation of p38, ERK and JNK in RAW264.7 cells <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			
In Vivo	Akt/NF-κB/MAPK-IN-1 (compound 2m) (10-20 mg/kg, IG, for 7 consecutive days) reduces LPS-induced inflammatory disease in mice <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.			
	Animal Model:	Male ICR mice (weighing 20-25 g) <sup>[1]</sup>		
	Dosage:	10 and 20 mg/kg		
	Administration:	Intragastric administration, for 7 consecutive days, and then LPS was intraperitoneally injected		
	Result:	The levels of IL-6, IL-1 $\beta$ and TNF- $\alpha$ in the LPS group were significantly increased compared to the control group, and the administration of 2m effectively prevented the increase.		

## **REFERENCES**

[1]. Li X, et al. Design, synthesis and evaluation of ursodeoxycholic acid-cinnamic acid hybrids as potential anti-inflammatory agents by inhibiting Akt/NF-кВ and MAPK signaling pathways. Eur J Med Chem. 2023 Aug 31;260:115785.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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