## Axinelline A

MedChemExpress

Cat. No.:	HY-N11624	
CAS No.:	1593741-99-1	ОН
Molecular Formula:	C <sub>12</sub> H <sub>15</sub> NO <sub>6</sub>	OH
Molecular Weight:	269.25	
Target:	COX	
Pathway:	Immunology/Inflammation	Ö <sup>€</sup> _OH
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	On

BIOLOGICAL ACTIVITY			
Description	Axinelline A is a potent COX inhibitor with IC <sub>50</sub> s of 2.22 μM and 8.89 μM against COX-2 and COX-1, respectively. Axinelline A shows anti-inflammatory activity <sup>[1]</sup> .		
IC <sub>50</sub> & Target	COX-2 2.22 μM (IC <sub>50</sub> )	COX-1 8.89 μΜ (IC <sub>50</sub> )	
In Vitro	Axinelline A (2-30 μM; 24 h) inhibits Lipopolysaccharide (LPS; HY-D1056)-induced expression of pro-inflammatory factors (NO, TNF-α, IL-6, IL-1β, and PGE <sub>2</sub> ) in RAW264.7 cells <sup>[1]</sup> . Axinelline A (2-30 μM; 24 h) inhibits LPS-induced NF-κB signaling pathway in RAW264.7 cells <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis <sup>[1]</sup>		
	Cell Line:	RAW264.7 cells	
	Concentration:	2, 10 and 30 μM	
	Incubation Time:	24 h	
	Result:	Diminished LPS-induced expression of nitric oxide synthase (iNOS) and COX-2 protein levels. The phosphorylation level of NF-κB increased after 30 min of LPS treatment, but pretreatment with test compound reduced the level of phosphorylation in a dose- dependent manner. The phosphorylation of IKK and IκBα was inhibited in a dose- dependent manner.	

## REFERENCES

[1]. Ju Z, et al. Synthesis and Anti-Inflammatory Activity of the Natural Cyclooxygenase-2 Inhibitor Axinelline A and Its Analogues. J Nat Prod. 2023 Apr 28;86(4):958-965.

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

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA