Me-Indoxam

BIOLOGICAL ACTIV

Cat. No.:	HY-125157	
CAS No.:	172732-62-6	
Molecular Formula:	C ₂₆ H ₂₂ N ₂ O ₅	
Molecular Weight:	442.46	
Target:	Phospholipase	
Pathway:	Metabolic Enzyme/Protease	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	НС

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VITY		
Me-Indoxam is a potent and cell-impermeable secreted phospho	olipase A_2 (sPLA ₂) inhibitor ^[1] .	
sPLA2		

Description	Me-Indoxam is a potent and cell-impermeable secreted phospholipase A_2 (sPLA ₂) inhibitor ^[1] .
IC ₅₀ & Target	sPLA2
In Vitro	 Me-Indoxam shows different potencies on the various sPLA₂ isoforms in in vitro assays with an IC₅₀ of less than 100 nM for hGIIA, hGIIE, and hGV; an IC₅₀ of between 200 and 600 nM for hGIB and hGX; and an IC₅₀ of greater than 2 mM for hGIID, hGIIF, hGIII, and hGXIIA^[1]. Me-Indoxam is able to inhibit the activity of sPLA₂s only when they are secreted in the extracellular space^[1]. Me-Indoxam (0-10 µM; 30 min) inhibits Leukotriene C₄ (LTC₄) production from anti-IgE-stimulated primary human lung mast cells (HLMCs)^[1]. Me-indoxam (0.01–10 M) did not affect the basal secretion of TNF-α and IL-6 from human lung macrophages (HLM). Preincubation (15 min, 37°C) of hGX (1 µg/mL) with various concentrations of Me-indoxam before the addition to HLM dose-dependently inhibited TNF-α and IL-6 release. The IC₅₀ values of Me-indoxam were 253±72 and 320±87 nM on TNF-α and IL-6 release, respectively^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Triggiani M, et al. Lung mast cells are a source of secreted phospholipases A2. J Allergy Clin Immunol. 2009 Sep;124(3):558-65, 565.e1-3.

[2]. Granata F, et al. Activation of cytokine production by secreted phospholipase A2 in human lung macrophages expressing the M-type receptor. J Immunol. 2005 Jan 1;174(1):464-74.

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

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