Product Data Sheet

Syk Inhibitor II dihydrochloride

Cat. No.: HY-112390 CAS No.: 227449-73-2

Molecular Weight: 413.23

Molecular Formula:

Target: 5-HT Receptor; Syk

Pathway: GPCR/G Protein; Neuronal Signaling; Protein Tyrosine Kinase/RTK

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

Analysis.

 $C_{14}H_{17}Cl_{2}F_{3}N_{6}O$

BIOLOGICAL ACTIVITY

Description	Syk Inhibitor II dihydrochloride is a potent, high selective and ATP-competitive Syk inhibitor with an IC ₅₀ of 41 nM. Syk Inhibitor II dihydrochloride inhibits 5-HT release from RBL-cells with an IC ₅₀ of 460 nM. Syk Inhibitor II dihydrochloride shows less potent against other kinases and has anti-allergic effect ^[1] .	
IC ₅₀ & Target	5-HT Receptor 460 nM (IC ₅₀)	
In Vitro	Syk Inhibitor II (compound 9a) dihydrochloride shows less potent against PKC ϵ , PKC β 2, ZAP-70, Btk, and Itk with IC $_{50}$ values of 5.1 μ M, 11 μ M, 11.2 μ M, 15.5 μ M, and 22.6 μ M, respectively ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
In Vivo	Syk Inhibitor II (Compound 9a; 10-100 mg/kg) dihydrochloride is subcutaneously administered to mice 30 min before antigen challenge. Syk Inhibitor II inhibits the anaphylaxis reaction dose-dependently with an ID_{50} value of 13.2 mg/kg ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	$ICRmice^{[1]}$
	Dosage:	10, 30, 100 mg/kg
	Administration:	S.c.; 30 min before antigen challenge
	Result:	Inhibited the anaphylaxis reaction dose-dependently with an ID ₅₀ value of 13.2 mg/kg.

REFERENCES

[1]. Hiroyuki Hisamichi, et al. Synthetic studies on novel Syk inhibitors. Part 1: Synthesis and structure-activity relationships of pyrimidine-5-carboxamide derivatives. Bioorg Med Chem. 2005 Aug 15;13(16):4936-51.

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

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