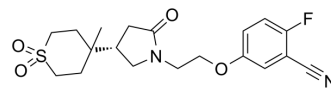


Lp-PLA2-IN-2

Cat. No.:	HY-133148
CAS No.:	2071636-15-0
Molecular Formula:	C ₁₉ H ₂₃ FN ₂ O ₄ S
Molecular Weight:	394.46
Target:	Phospholipase
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Lp-PLA2-IN-2 is a potent and selective lipoprotein-associated phospholipase A2 (Lp-PLA2) inhibitor, with an IC ₅₀ of 120 nM for recombinant human Lp-PLA2 ^[1] .								
IC₅₀ & Target	Lp-PLA2 120 nM (IC ₅₀)								
In Vitro	Lp-PLA2-IN-2 inhibits Human Lp-PLA2 with an IC ₅₀ of 62 nM in PED6 assay ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.								
In Vivo	Lp-PLA2-IN-2 (1 mg/kg; i.v.) treatment in rats shows that the Cl, Vss and t _{1/2} values are 67 mL/min/kg, 1.2 L/kg, and 0.34 hours, respectively ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.								
	<table border="1"> <tr> <td>Animal Model:</td> <td>Male Han Wistar rats (250-300 g)^[1]</td> </tr> <tr> <td>Dosage:</td> <td>1 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>i.v. (Pharmacokinetic Analysis)</td> </tr> <tr> <td>Result:</td> <td>The Cl, Vss and t_{1/2} values were 67 mL/min/kg, 1.2 L/kg, and 0.34 hours, respectively.</td> </tr> </table>	Animal Model:	Male Han Wistar rats (250-300 g) ^[1]	Dosage:	1 mg/kg	Administration:	i.v. (Pharmacokinetic Analysis)	Result:	The Cl, Vss and t _{1/2} values were 67 mL/min/kg, 1.2 L/kg, and 0.34 hours, respectively.
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REFERENCES

[1]. Woolford AJ, et al. Fragment-Based Approach to the Development of an Orally Bioavailable Lactam Inhibitor of Lipoprotein-Associated Phospholipase A2 (Lp-PLA2). *J Med Chem.* 2016 Dec 8;59(23):10738-10749.

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

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