

## LSD1-IN-14

## Chemical Properties

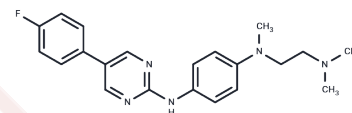
CAS No. : 2698340-11-1

Formula: C<sub>21</sub>H<sub>24</sub>FN<sub>5</sub>

Molecular Weight: 365.45

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Actual storage temperature shall be subject to the COA.



## Biological Description

|                            |  |
|----------------------------|--|
| Description                | LSD1-IN-14, a powerful and specific inhibitor of LSD1 (with an IC <sub>50</sub> value of 0.89 μM), effectively suppresses the growth of A549 and THP-1 cells and promotes the apoptosis of tumor cells [1].  |
| Targets(IC <sub>50</sub> ) | Apoptosis,Histone Demethylase,Others   |
| In vitro                   | LSD1-IN-14 (compound x43) inhibits the proliferation of A549 and THP-1 cells at concentrations of 0-20 μM over 72 hours, with IC <sub>50</sub> values of 1.62 μM and 1.21 μM, respectively [1]. At 0-3 μM, it increases H3K4me2 and H3K9me2 expression in A549 cells and induces apoptosis in 53.6% of these cells in a dose-dependent manner [1]. LSD1-IN-14 demonstrates metabolic stability in human liver microsomes with a half-life of 103.3 minutes, showing minimal inhibition of cytochrome P450 enzymes (Cl int(mic) value of 13.4 μL/min/mg at 1 mM for 60 minutes) [1]. This compound's antiproliferative effects, epigenetic modulation, and apoptosis induction make it a promising candidate for further research [1].  |
| In vivo                    | LSD1-IN-14, administered at dosages of 2 mg/kg intravenously (i.v.) and 10 mg/kg intragastrically (i.g.) to male Sprague-Dawley rats, exhibits an acceptable half-life and oral bioavailability. Pharmacokinetic analysis revealed: for i.v. administration, a peak concentration (C max) of 41.1 ng/mL, a half-life (T 1/2) of 1.0 hour, a steady-state volume of distribution (Vd ss) of 6.6 L/kg, a clearance rate (Cl) of 156 mL/min/kg, and an area under the curve (AUC 0-t) of 126 ng.h/mL extending to 152 ng.h/mL (AUC 0-∞). For i.g. administration, the peak concentration reached 575 ng/mL at a T max of 0.8 hours, a half-life of 2.8 hours, with an AUC 0-t of 211 ng.h/mL extending to 214 ng.h/mL (AUC 0-∞), reflecting an oral bioavailability of 11.9%. These findings suggest LSD1-IN-14 has promising pharmacokinetic properties for further development. |

### Preparing Stock Solutions

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|       | <b>1mg</b> | <b>5mg</b> | <b>10mg</b> |
|-------|------------|------------|-------------|
| 1 mM  | 2.7364 mL  | 13.6818 mL | 27.3635 mL  |
| 5 mM  | 0.5473 mL  | 2.7364 mL  | 5.4727 mL   |
| 10 mM | 0.2736 mL  | 1.3682 mL  | 2.7364 mL   |
| 50 mM | 0.0547 mL  | 0.2736 mL  | 0.5473 mL   |

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Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

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