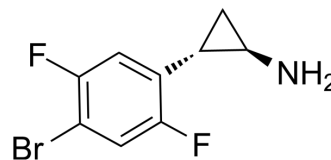


LSD1-IN-22

Cat. No.:	HY-151191
CAS No.:	2821068-05-5
Molecular Formula:	C ₉ H ₈ BrF ₂ N
Molecular Weight:	248.07
Target:	Histone Demethylase
Pathway:	Epigenetics
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	LSD1-IN-22 is a potent Lysine-specific demethylase 1 (LSD1) inhibitor with a K _i value of 98 nM. LSD1-IN-22 has anti-proliferative activity against certain cancer cells ^[1] .								
IC₅₀ & Target	KDM1/LSD1								
In Vitro	<p>LSD1-IN-22 (compound 7t) has anti-proliferative activity against CCRF-CEM, Jurkat and hERG cells^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>CCRF-CEM, Jurkat and hERG</td> </tr> <tr> <td>Concentration:</td> <td>0-50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Exhibited anti-proliferative activity against CCRF-CEM, Jurkat and hERG cells with IC₅₀s of 23±0.8 μM, 26±1.7 μM and 7.7±1.3 μM.</td> </tr> </table>	Cell Line:	CCRF-CEM, Jurkat and hERG	Concentration:	0-50 μM	Incubation Time:	72 h	Result:	Exhibited anti-proliferative activity against CCRF-CEM, Jurkat and hERG cells with IC ₅₀ s of 23±0.8 μM, 26±1.7 μM and 7.7±1.3 μM.
Cell Line:	CCRF-CEM, Jurkat and hERG								
Concentration:	0-50 μM								
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REFERENCES

[1]. Hideaki Niwa, et al. Structure-Activity Relationship and In Silico Evaluation of cis- and trans-PCPA-Derived Inhibitors of LSD1 and LSD2. ACS Medicinal Chemistry Letters Article ASAP.

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

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