LSD1-IN-17

Cat. No.: Molecular Formula: Molecular Weight: Target: Pathway: Storage:	HY-144758 C ₂₀ H ₁₈ N ₂ OS 334.43 Histone Demethylase; Monoamine Oxidase Epigenetics; Neuronal Signaling Please store the product under the recommended conditions in the Certificate of Analysis.	S O NH2
---	---	---------

BIOLOGICAL ACTIV				
Description	LSD1-IN-17 (compound 5b) is a potent LSD1 inhibitor. LSD1-IN-17 can inhibit LSD1-CoREST, MAO-A and MAO-B, with IC ₅₀ values of 0.005, 0.028, and 0.820 μM, respectively. LSD1-IN-17 displays cell growth arrest in prostate cancer LNCaP cells, with an IC ₅₀ of 17.2 μM ^[1] .			
IC₅₀ & Target	MAO-A 0.028 ± 0. μM (IC ₅₀)	MAO-B 0.820 ± 0. μM (IC ₅₀)		

REFERENCES

[1]. Fioravanti R, et al. Heterocycle-containing tranylcypromine derivatives endowed with high anti-LSD1 activity. J Enzyme Inhib Med Chem. 2022 Dec;37(1):973-985.

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

Fax: 609-228-5909 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA

Product Data Sheet

Tel: 609-228-6898

E-mail: tech@MedChemExpress.com