## **Product** Data Sheet

Inhibitors

**Screening Libraries** 

**Proteins** 

## **FY-56**

Molecular Weight: 390.41

Target: Histone Demethylase

Pathway: Epigenetics

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

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	FY-56 is a highly potent and selective LSD1/KDM1A inhibitor (IC $_{50}$ =42 nM) and exhibits high selectivity over MAO-A/B. FY-56 induces differentiation of MOLM-13 and MV4-11 cell and has the potential for AML research <sup>[1]</sup> .
IC <sub>50</sub> & Target	IC50: 42 nM (LSD1/KDM1A) <sup>[1]</sup>

## **REFERENCES**

[1]. Chao Yang, et al. Discovery of natural product-like spirooxindole derivatives as highly potent and selective LSD1/KDM1A inhibitors for AML treatment. Bioorg Chem. 2022 Mar; 120:105596

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA