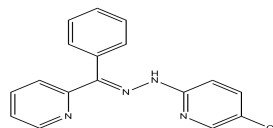


**Product: Jumonji HDM Inhibitor,
JIB-04****ALTERNATE NAME:** 5-Chloro-2-[(*E*)-2-[phenyl(pyridin-2-yl)methylidene]hydrazin-1-yl]pyridine; NSC693627**CATALOG #:** 2474-5, 25**AMOUNT:** 5 mg, 25 mg**STRUCTURE:****MOLECULAR FORMULA:** C₁₇H₁₃ClN₄**MOLECULAR WEIGHT:** 308.76**CAS No.** 199596-05-9**APPEARANCE:** White solid**SOLUBILITY:** DMSO (~20 mg/ml)**PURITY:** >98% by HPLC**STORAGE:** Store at -20°C. Protect from air and light**DESCRIPTION:** JIB-04 (*E*-isomer) is a potent, cell-permeable inhibitor of Jumonji histone demethylases in vitro and in vivo. (In vitro IC₅₀ values are 230, 340, 435, 445, 855 and 1100 nM for JARID1A, JMJD2E, JMJD2B, JMJD2A, JMJD3 and JMJD2C respectively). JIB-04 is not a competitive inhibitor of α-ketoglutarate ; it selectively inhibits viability of several human cancer cell lines with little toxicity towards normal cells, and also diminishes tumor growth in xenograft mouse models.**REFERENCES:** Wang, L., *et al.* (2013). *Nat. Commun.* **4**, 2045.**HANDLING:** Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of exposure.**RELATED PRODUCTS:**

Caffeic acid (Cat. No. 2303-50, 250)

Daminozide (Cat. No. 2438-100, 500)

Disulfiram (Cat. No. 2308-10, 50)

Ebselen (Cat. No. 2169-5, 25)

GSK-J4 hydrochloride (Cat. No. 2259-1, 5)

GSK-J1 sodium (Cat. No. 2260-1, 5)

IOX1 (Cat. No. 2266-5, 25)

β-Lapachone (Cat. No. 2262-5, 25)

2,4-PDCA (Cat. No. 2304-100, 500)

Tranylcypromine hemisulfate (Cat. No. 1816-25, 100)

FOR RESEARCH USE ONLY! Not to be used in humans.