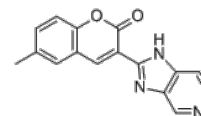


Product Name : iJMJD6
Cat. No. : PC-49244
CAS No. : 899548-78-8
Molecular Formula : C₁₆H₁₁N₃O₂
Molecular Weight : 277.283
Target : Histone Demethylase
Solubility : 10 mM in DMSO



Biological Activity

iJMJD6 is a potent, specific small-molecule JMJD6 inhibitor that specifically binds to **JMJD6** (ITC K_d=3.83 μM) and competes with α-KG and the substrate for binding, inhibits the demethylase activity of JMJD6 with IC₅₀ of 149.6 nM. iJMJD6 suppresses the malignant phenotypes of HeLa cervical cancer cells, inhibits cell proliferation with EC₅₀ of 1.49, 9.37, and 9.53 μM for HeLa, SMCC7721, and MCF7 cells, respectively. iJMJD6 inhibits JMJD6-mediated demethylation in HeLa cells, inhibits the expression of oncogenes, including c-Myc, N-myc, and CCND1, inhibit the transcription of these oncogenes through inhibiting JMJD6-mediated demethylation of H4R3me2(s). iJMJD6 (25 mg/kg, 50 mg/kg, i.p.) significantly suppressed the tumor growth of HeLa cell-derived xenografts and patient-derived xenograft mouse models. iJMJD6 exhibits synergistic effects on tumor growth when combined with BRD4 inhibitor JQ1 and ERα degrader fulvestrant.

References

Rong-Quan Xiao, et al. *Proc Natl Acad Sci U S A*. 2022 Aug 23;119(34):e2200753119.

Caution: Product has not been fully validated for medical applications. Lab Use Only!

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