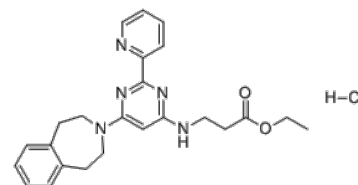


Product Name : GSK-J4
Cat. No. : PC-21211
CAS No. : 1373423-53-0
Molecular Formula : C₂₄H₂₈ClN₅O₂
Molecular Weight : 453.97
Target : Histone Demethylase
Solubility : 10 mM in DMSO



Biological Activity

GSK-J4 is a potent dual inhibitor of H3K27me₃/me₂-demethylases JMJD3/KDM6B and UTX/KDM6A with IC₅₀s of 8.6 and 6.6 μM, respectively, GSK-J4 is a cell permeable prodrug of GSK-J1.

GSK-J4 inhibits LPS-induced TNF-α production in human primary macrophages with IC₅₀ of 9 μM.

GSK-J4 prevents the JMJD3-induced loss of nuclear H3K27me₃ immunostaining in Flag-JMJD3-transfected HeLa cells.

GSK-J4 inhibits JMJD3 expression that is induced by TGF-β₁, inhibits H3K4 demethylation at Xist, Nodal, and HoxC13 in female embryonic stem cells.

GSK-J4 attenuates the development of kidney disease in diabetic mice.

References

Kruidenier L, et al. Nature. 2012 Aug 16;488(7411):404-8.

Majumder S, et al. J Clin Invest. 2018 Jan 2;128(1):483-499.

Donas C, et al. J Autoimmun. 2016 Dec;75:105-117.

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

E-mail: tech@probechem.com