

Data Sheet

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 Product Name
 : TAK-418

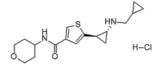
 Cat. No.
 : PC-20519

 CAS No.
 : 1818252-53-7

 Molecular Formula
 : C₁₇H₂₅CIN₂O₂S

Molecular Weight: 356.91

Target : Histone Demethylase
Solubility : 10 mM in DMSO



CAS: 1818252-53-7

Biological Activity

TAK-418 (TAK418) is a potent, specific inhibitor of LSD1 (KDM1A) enzyme activity with IC50 of 2.9 nM.

TAK-418 does not inhibit recombinant human MAO-A and MAO-B with IC50 of >100 uM.

TAK-418 (0.1-10 uM) increases H3K4me1/2/3 and H3K9me2 levels at the Ucp2 gene and induces Ucp2 mRNA expression in primary cultured rat neurons, also increases H3K4me1/2/3 at the Bdnf gene.

TAK-418 has minimal impact on the interaction between LSD1 and its cofactor, GFI1B.

Single administration of TAK-418 at 1 or 3 mg/kg increases H3K4me2 levels at Ucp2 gene in the mouse brain.

TAK-418 normalizes mRNA expression dysregulated in neurodevelopmental disorder model rodents.

TAK-418 produces inverse correlation against the histone modification and DNA methylation levels altered in VPA rats and poly I:C mice.

TAK-418 ameliorates some ASD-like behaviors in neurodevelopmental disorder model rodents.

References

Baba R, et al. *Sci Adv.* 2021 Mar 12;7(11):eaba1187.

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

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