MCE ®

BI-9321

 $\begin{array}{lll} \textbf{Cat. No.:} & HY\text{-}114208 \\ \\ \textbf{CAS No.:} & 2387510\text{-}86\text{-}1 \\ \\ \textbf{Molecular Formula:} & C_{22}H_{21}FN_4 \\ \end{array}$

Molecular Weight: 360.43

Target: Histone Methyltransferase

Pathway: Epigenetics

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

Analysis.

BIOLOGICAL ACTIVITY

Description	BI-9321 is a potent, selective and cellular active nuclear receptor-binding SET domain 3 (NSD3)-PWWP1 domain antagonist with a K_d value of 166 nM. BI-9321 is inactive against NSD2-PWWP1 and NSD3-PWWP2. BI-9321 specifically disrupts histone interactions of the NSD3-PWWP1 domain with an IC $_{50}$ of 1.2 μ M in U2OS cells $^{[1]}$.
IC ₅₀ & Target	Kd: 166 nM (NSD3-PWWP1) ^[1]
In Vitro	BI-9321 is targeting the methyl-lysine binding site of the PWWP1 domain with sub-micromolar in vitro activity and cellular target engagement at 1 μ M. As a single agent, BI-9321 downregulates Myc messenger RNA expression and reduces proliferation in MOLM-13 cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Böttcher J, et al. Fragment-based discovery of a chemical probe for the PWWP1 domain of NSD3. Nat Chem Biol. 2019 Aug;15(8):822-829.

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

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