Cat No. 74-089

BIX01294

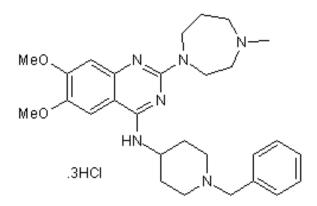
REAGENTS DIRECT

5 mg

For research purposes only

BIX01294 is a histone lysine methyltransferase (HMTase) inhibitor that modulates the epigenetic status of chromatin. It selectively inhibits the G9a HMTase and the generation of histone H3 lysine 9 methylation. BIX01294, when used in combination with HDAC inhibitor Valproic Acid has been used as a replacement for ectopic OCT4 (POU5F1) and cMyc respectively in pluripotent stem cell induction (iPS) recipes. BIX01294 is a useful for the study of histone lysine methylation, differentiation and reprogramming.

TECHNICAL INFORMATION



Other Names: 2-(Hexahydro-4-methyl-1*H*-1,4-diazep in-1-yl)-6,7-dimethoxy-*N*-[1-(phenylmethyl)-4-piper idinyl]-4-quinazolinamine trihydrochloride

Chemical Formula: C₂₈H₃₈N₆O₂.3HCl

CAS Number: 935693-62-2 Molecular Weight: 600.02

Purity: >99%

Appearance: white solid

Solubility: DMSO

STORAGE AND HANDLING

Storage: Store at 4°C and protected from light. Following reconstitution, store aliquots at -20°C.

Stability: Stock solutions stable at -20°C for up to 2 years.

Shipping Conditions: Shipped at room temperature.

PRODUCT USE

Soluble in DMSO. For a 10mM concentrated stock solution, reconstitute the compound in 1.02ml of DMSO to the entire contents of the vial. If precipitate is observed, vortex for 5 minutes. For most cells, the maximum tolerance to DMSO is less than 0.5%.

REFERENCES

- Chang et al. (2009) Structural basis for G9a-like protein lysine methyltransferase inhibition by BIX-01294. Nat Struct Mol Biol. 16(3):312-17.
- 2. Kubick et al. (2007) Reversal of H3K9me2 by a small-molecule inhibitor for the G9a histone methyltransferase. Mol Cell. 25(3): 473-81.
- 3. Shi et al. (2008) Induction of pluripotent stem cells from mouse embryonic fibroblasts by Oct4 and Klf4 with small-molecule compounds. Cell Stem Cell. 3 (5):568-74.



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