

GSK-J2 (Sodium Salt)

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3-((6-(4,5-dihydro-1H-benzo[d]azepin-3(2H)-yl)-2-(pyridin-3-yl)pyrimidin-4-**ALTERNATE NAMES:**

yl)amino)propanoicacid,monosodiumsalt, 3-[[2-pyridin-3-yl-6-(1,2,4,5-tetrahydro-3-benzazepin-3-

yl)pyrimidin-4-yl]amino]propanoic acid;sodium

B2813-1 CATALOG #: B2813-5 5 mg

STRUCTURE:

MOLECULAR FORMULA: $C_{22}H_{23}N_5NaO_2$

MOLECULAR WEIGHT: 411.43

CAS NUMBER: 2108665-15-0

APPEARANCE: Crystalline solid

PURITY: ≥90%

Sparingly soluble in aqueous and organic solvents such as DMSO, ethanol and DMF SOLUBILITY:

DESCRIPTION: GSK-J2 (sodium salt) is an isomer of GSK-J1. GSK-1 is a selective inhibitor of the H3K27 histone

> demethylases JMJD3 and UTX, with an IC $_{50}$ of 60 nM for human JMJD3 in vitro. GSK-J2 poorly inhibits JMJD3 with an IC₅₀ > 100 μM in vitro. GSK-J2 serves as a negative control for in vitro studies with

GSK-J1.

STORAGE TEMPERATURE: -20°C. Store under desiccating conditions.

HANDLING: Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of

exposure.

RELATED PRODUCTS:

GSK-J4 hydrochloride (Cat. No. 2259) GSK-J1 sodium salt (Cat. No. 2260) GSK-J1 (Free acid) (Cat. No. 2761) DiscoveryPak™ Epigenetic Modulators Set I (Cat. No. K879)

GSK-J4 (Free base) (Cat. No. 2762)

DISCLAIMER: FOR RESEARCH USE ONLY! Not to be used on humans.