

Product Information

Product ID U524081

CAS No. 1431612-23-5

Chemical Name

Synonym

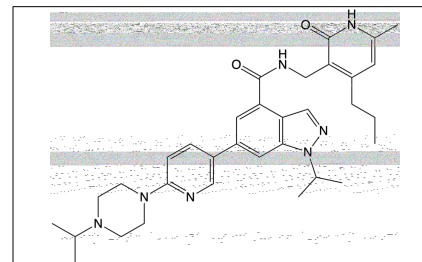
Formula C₃₃H₄₃N₇O₂

Formula Wt. 569.7

Melting Point

Purity ≥98%

Solubility



Pricing and Availability

Bulk quantities available upon request

Product ID	Size	List Price
U524081	5 mg	\$125.00
U524081	25 mg	\$350.00

Store Temp -20° C

Ship Temp Ambient

Description UNC1999 is a potent, orally bioavailable inhibitor that selectively inhibits both EZH2 (enhancer of zeste homolog 2) and EZH1, the two catalytic subunits of the Polycomb Repressive Complex 2 (PRC2) responsible for tri-methylation of histone H3 at lysine 27 (H3K27me3). Unlike EZH2-selective inhibitors, UNC1999 dual inhibition provides more complete suppression of H3K27 methylation because EZH1 can compensate for EZH2 loss. UNC1999 is an important chemical probe for studying PRC2-mediated gene silencing, differentiation, and epigenetic dependencies in cancer.

- References**
1. Konze KD, Ma A, Li F, et al. An orally bioavailable chemical probe of the Lysine Methyltransferases EZH2 and EZH1. *ACS Chem Biol*. 2013;8(6):1324-34. PMID: 23614352.
 2. Morin RD, Johnson NA, Severson TM, et al. Somatic mutations altering EZH2 (Tyr641) in follicular and diffuse large B-cell lymphomas of germinal-center origin. *Nat Genet*. 2010;42(2):181-5. PMID: 20081860.
 3. Margueron R, Reinberg D. The Polycomb complex PRC2 and its mark in life. *Nature*. 2011;469(7330):343-9. PMID: 21248841.
 4. McCabe MT, Ott HM, Ganji G, et al. EZH2 inhibition as a therapeutic strategy for lymphoma with EZH2-activating mutations. *Nature*. 2012;492(7427):108-12. PMID: 23051747.
 5. Xu B, On DM, Ma A, et al. Selective inhibition of EZH2 and EZH1 enzymatic activity by a small molecule suppresses MLL-rearranged leukemia. *Blood*. 2015;125(2):346-57. PMID: 25395426.

Caution: This product is intended for laboratory and research use only. It is not for human or drug use.