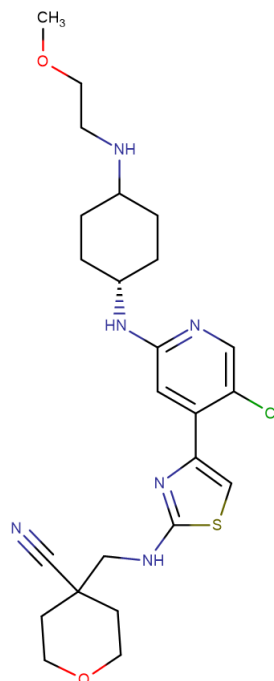


**Product Name** : JSH-150  
**Synonyms** : JSH150;JSH 150  
**Cat No.** : M13607  
**CAS Number** : 2247481-21-4  
**Molecular Formula** : C<sub>24</sub>H<sub>33</sub>ClN<sub>6</sub>O<sub>2</sub>S

**Formula Weight** : 505.08



**Chemical Name** : 4-(((4-(5-Chloro-2-(((1r,4r)-4-((2-methoxyethyl)amino)cyclohexyl)amino)pyridin-4-yl)thiazol-2-yl)amino)methyl)tetrahydro-2H-pyran-4-carbonitrile

**Description** : JSH-150 (JSH150) is a potent, highly selective inhibitor of CDK9 kinase with IC<sub>50</sub> of 1 nM in the biochemical assays, displays 300-10000-fold selectivity over other CDK kinase family members (CDK7 IC<sub>50</sub>=1.72 μM); also exhibits high selectivity over other 468 kinases/mutants (KINOMEscan S score(1)=0.01); displays potent antiproliferative effects against melanoma, neuroblastoma, hepatoma, colon cancer, lung cancer as well as leukemia cell lines, dose-dependently inhibits the phosphorylation of RNA Pol II, suppresses the expression of MCL-1 and c-Myc, arrests the cell cycle and induces the apoptosis in the leukemia cells; completely suppresses the tumor progression in MV4-11 cell-inoculated xenograft mouse model (10 mg/kg).

**Pathway** : Angiogenesis

**Target** : CDK

**Receptor** : CDK

**Solubility** : —

**SMILES** : COCCNC1CC[C@H](NC2=NC=C(Cl)C(C3=CSC(NCC4(C#N)CCOCC4)=N3)=C2)CC1

**Storage** : (-20°C)

**Stability** : ≥ 2 years

**Reference** :

1. Wang B, et al. Eur J Med Chem. 2018 Oct 5;158:896-916.