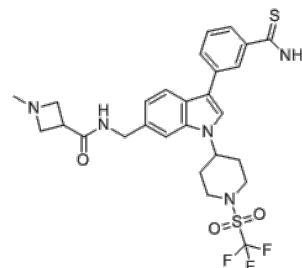


**Product Name** : ASH1L inhibitor AS-99  
**Cat. No.** : PC-72494  
**CAS No.** : 2323623-93-2  
**Molecular Formula** : C<sub>27</sub>H<sub>30</sub>F<sub>3</sub>N<sub>5</sub>O<sub>3</sub>S<sub>2</sub>  
**Molecular Weight** : 593.684  
**Target** : Histone Methyltransferase (HMTase)  
**Solubility** : 10 mM in DMSO



## Biological Activity

ASH1L inhibitor AS-99 (AS99) is a first-in-class, potent, selective inhibitor of **ASH1L** histone methyltransferase with IC<sub>50</sub> of 0.79 μM.

AS-99 strongly bind to the ASH1L SET domain with K<sub>d</sub> values of 0.89 μM.

AS-99 displayed no significant inhibition (>100-fold selectivity) at 50 μM against a panel of 20 histone methyltransferases, including NSD1, NSD2, NSD3, and SETD2.

AS-99 inhibits the growth of leukemia cells (MV4;11, MOLM13, and KOPN8) harboring different MLL1 translocations with the GI<sub>50</sub> values of 1.8-3.6 μM, showed a several fold weaker effect on the proliferation of leukemia cells without MLL1 translocations, such as SET2 and K562, without toxicity in normal cells.

AS-99 impairs transcriptional program of MLL fusion proteins and reduces leukemia burden.

AS-99 reduced the leukemia burden in the xenotransplantation mouse model of MLL leukemia without affecting blood counts in normal mice.

## References

David S Rogawski, et al. **Nat Commun.** 2021 May 14;12(1):2792.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

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