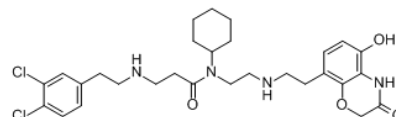


**Product Name** : AZ505  
**Cat. No.** : PC-44245  
**CAS No.** : 1035227-43-0  
**Molecular Formula** : C<sub>29</sub>H<sub>38</sub>Cl<sub>2</sub>N<sub>4</sub>O<sub>4</sub>  
**Molecular Weight** : 577.5424  
**Target** : Histone Methyltransferase (HMTase)  
**Solubility** : 10 mM in DMSO



### Biological Activity

AZ505 (AZ-505, AZ 505) is a potent and selective inhibitor of protein lysine methyltransferase **SMYD2** with IC<sub>50</sub> of 0.12 μM, displays >600-fold against other histone methyltransferases, such as SMYD3 and EZH2.

AZ505 suppresses BMP2-induced SMAD1/5 phosphorylation, delays cyst growth in both early- and later-stage Pkd1 conditional knockout mouse models.

AZ505 also significantly reduced tumor growth in vivo in triple-negative breast cancer (TNBC) models.

### References

Ferguson AD, et al. **Structure**. 2011 Sep 7;19(9):1262-73.

Gao S, et al. **J Biol Chem**. 2017 Jul 28;292(30):12702-12712.

Li LX, et al. **J Clin Invest**. 2017 Jun 30;127(7):2751-2764.

Li LX, et al. **Cell Death Dis**. 2018 Feb 27;9(3):326.

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

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