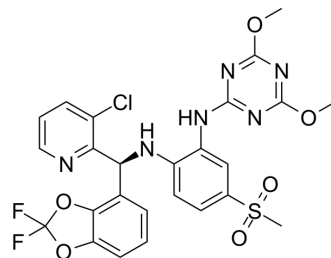


## Dot1L-IN-6

Cat. No.:	HY-135129
CAS No.:	2565705-01-1
Molecular Formula:	C <sub>25</sub> H <sub>21</sub> ClF <sub>2</sub> N <sub>6</sub> O <sub>6</sub> S
Molecular Weight:	606.99
Target:	Histone Methyltransferase
Pathway:	Epigenetics
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Dot1L-IN-6 is a potent disruptor of telomeric silencing 1-like protein (DOT1L) inhibitor with an IC <sub>50</sub> SPA DOT1L of 0.19 nM <sup>[1]</sup> .
<b>In Vitro</b>	Dot1L-IN-6 (Compound 9) is tested in cellular assays to assess the ability to inhibit the dimethylation of H3K79 in HeLa cells (ED <sub>50</sub> H3K79me2 Elisa=12 nM) and HOXA9 gene expression in Molm-13 cells (ED <sub>50</sub> HOXA9 RGA=170 nM) <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
<b>In Vivo</b>	Dot1L-IN-6 (Compound 9) shows an excellent blood exposure after a single dose (100 mg/kg) <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Frédéric Stauffer, et al. New Potent DOT1L Inhibitors for in Vivo Evaluation in Mouse. ACS Med. Chem. Lett. 2019, 10, 12, 1655-1660.

**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA