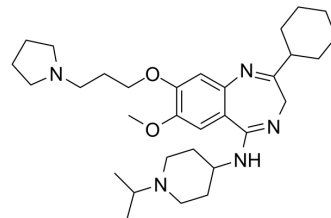


## EML741

<b>Cat. No.:</b>	HY-111544
<b>CAS No.:</b>	2328074-38-8
<b>Molecular Formula:</b>	C <sub>31</sub> H <sub>49</sub> N <sub>5</sub> O <sub>2</sub>
<b>Molecular Weight:</b>	523.75
<b>Target:</b>	Histone Methyltransferase
<b>Pathway:</b>	Epigenetics
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	EML741 is a histone lysine methyltransferase G9a/GLP inhibitor, with an IC <sub>50</sub> of 23 nM, K <sub>d</sub> of 1.13 μM for G9a. EML741 also inhibits DNMT1 (IC <sub>50</sub> , 3.1 μM), with no effect on DNMT3a or DNMT3b. EML741 exhibits low cell toxicity, and is membrane permeable and blood-brain barrier penetrated <sup>[1]</sup> .			
<b>IC<sub>50</sub> &amp; Target</b>	G9a 23 nM (IC <sub>50</sub> )	G9a 1.13 μM (K <sub>d</sub> )	GLP	DNMT1 3.1 μM (IC <sub>50</sub> )
<b>In Vitro</b>	EML741 (Compound 12a) shows a similar high inhibition potency against G9a (97%, 98% inhibition at 10 μM and 25 μM, respectively) and GLP (95%, 98% inhibition at 10 μM and 25 μM, respectively) <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

### REFERENCES

[1]. Milite, et al. Discovery of a Novel Chemotype of Histone Lysine Methyltransferase EHMT1/2 (GLP/G9a) Inhibitors: Rational Design, Synthesis, Biological Evaluation, and Co-crystal Structure. *J Med Chem.* 2019 Mar 14;62(5):2666-2689.

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

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