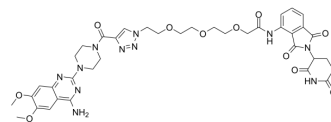


## α1A-AR Degradar 9c

Cat. No.:	HY-147100
CAS No.:	2863635-02-1
Molecular Formula:	C <sub>38</sub> H <sub>43</sub> N <sub>11</sub> O <sub>11</sub>
Molecular Weight:	829.82
Target:	Adrenergic Receptor; PROTACs
Pathway:	GPCR/G Protein; Neuronal Signaling; PROTAC
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	α1A-AR Degradar 9c (compound 9c) is a potent, selective and reversible α1A-AR (Adrenergic receptor) PROTAC degrader, with a DC <sub>50</sub> of 2.86 μM. α1A-AR Degradar 9c induces α1A-AR degradation can be attributed to proteasomal degradation. α1A-AR Degradar 9c inhibits the proliferation of PC-3 cells, with an IC <sub>50</sub> of 6.12 μM. α1A-AR Degradar 9c shows antitumor activity, and can be used for prostate cancer research <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	α1A-adrenergic receptor 2.86 μM (DC50)
<b>In Vivo</b>	α1A-AR Degradar 9c (compound 9c) (50 mg/kg, IP, once daily for 15 days) causes a significant suppression of tumor growth <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Li Z, et al. First small-molecule PROTACs for G protein-coupled receptors: inducing α1A-adrenergic receptor degradation. Acta Pharm Sin B. 2020 Sep;10(9):1669-1679.

**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