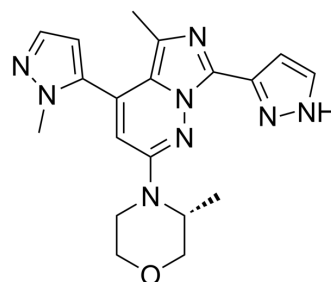


## ATR-IN-29

<b>Cat. No.:</b>	HY-153729
<b>CAS No.:</b>	2761193-67-1
<b>Molecular Formula:</b>	C <sub>19</sub> H <sub>22</sub> N <sub>8</sub> O
<b>Molecular Weight:</b>	378.43
<b>Target:</b>	ATM/ATR
<b>Pathway:</b>	Cell Cycle/DNA Damage; PI3K/Akt/mTOR
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



## BIOLOGICAL ACTIVITY

<b>Description</b>	ATR-IN-29 is a potent and orally active ATR kinase inhibitor with an IC <sub>50</sub> value of 1 nM. ATR-IN-29 shows antiproliferative activity <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	ATR 1 nM (IC <sub>50</sub> )
<b>In Vitro</b>	ATR-IN-29 (compound 1) (4 days) shows antiproliferative activity with IC <sub>50</sub> s of 156.70, 38.81, 22.48, 181.60, 19.02 nM for A549, HCC1806, HCT116, OVCAR-3, NCI-H460 cells, respectively <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
<b>In Vivo</b>	ATR-IN-29 (10 mg/kg; p.o.; once) shows good pharmacokinetic parameters with t <sub>1/2</sub> of 1.64 h, C <sub>max</sub> of 9343 ng/mL, AUC <sub>0-t</sub> of 98507 ng·h/mL, AUC <sub>0-inf</sub> of 98517 ng·h/mL in CD-1 (ICR) mice <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## REFERENCES

[1]. Sui Xiong Cai, et al. Substituted imidazo[1,5-b]pyridazine compounds as kinase inhibitors and use thereof. WO2022135560A1.

**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