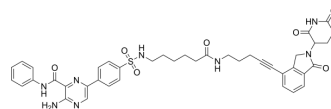


Abd110

Cat. No.:	HY-157767
CAS No.:	3021581-79-0
Molecular Formula:	C ₄₁ H ₄₂ N ₈ O ₇ S
Molecular Weight:	790.89
Target:	ATM/ATR; PROTACs
Pathway:	Cell Cycle/DNA Damage; PI3K/Akt/mTOR; PROTAC
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Abd110 (compound 42i) is a Lenalidomide-based PROTAC ATR kinase degrader. Abd110 selectively decreases ATR and phospho-ATR without affecting related kinases ATM and DNA-PKcs ^[1] .									
IC₅₀ & Target	ATR	Cereblon								
In Vitro	<p>Abd110 (compound 42i; 0.5-2 μM; 24 h) selectively degrades ATR through the proteasome, dependent on the E3 ubiquitin ligase component cereblon, and without affecting the associated kinases ATM and DNA-PKcs^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MIA PaCa-2 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.5 μM, 1 μM, and 2 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Decreased ATR and p-ATR levels without affecting related checkpoint kinases.</td> </tr> </table>		Cell Line:	MIA PaCa-2 cells	Concentration:	0.5 μM, 1 μM, and 2 μM	Incubation Time:	24 h	Result:	Decreased ATR and p-ATR levels without affecting related checkpoint kinases.
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Concentration:	0.5 μM, 1 μM, and 2 μM									
Incubation Time:	24 h									
Result:	Decreased ATR and p-ATR levels without affecting related checkpoint kinases.									

REFERENCES

[1]. Abdallah M Alfayomy, et al. Design, synthesis, and biological characterization of proteolysis targeting chimera (PROTACs) for the ataxia telangiectasia and RAD3-related (ATR) kinase. *Eur J Med Chem.* 2024 Mar 5:267:116167.

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

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