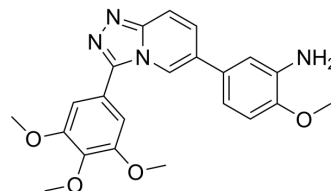


## Tubulin polymerization-IN-11

Cat. No.:	HY-146817
CAS No.:	2470063-59-1
Molecular Formula:	C <sub>22</sub> H <sub>22</sub> N <sub>4</sub> O <sub>4</sub>
Molecular Weight:	406.43
Target:	Microtubule/Tubulin; Apoptosis
Pathway:	Cell Cycle/DNA Damage; Cytoskeleton; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Tubulin polymerization-IN-11 is a potent tubulin polymerization inhibitor with an IC <sub>50</sub> value of 3.4 μM. Tubulin polymerization-IN-11 shows antiproliferative activity. Tubulin polymerization-IN-11 induces <a href="#">Apoptosis</a> and cell cycle arrest at G2/M phase. Tubulin polymerization-IN-11 decreases the expression of cyclin B1, p-cdc2, and Bcl-2 protein levels and increases the expression of cleaved PARP <sup>[1]</sup> .																
<b>In Vitro</b>	<p>Tubulin polymerization-IN-11 (compound 7i) (0-100 μM; 48 h) shows antiproliferative activity with IC<sub>50</sub>s of 0.012, &gt;100, 10.40, 40.40, 27.91 μM for HeLa, HEK-293, A549, MCF-7, T47D cells, respectively<sup>[1]</sup>.</p> <p>Tubulin polymerization-IN-11 (12, 24, 48 nM; 24 h) induces apoptosis and cell cycle arrest at G2/M phase in a dose-dependent manner<sup>[1]</sup>.</p> <p>Tubulin polymerization-IN-11 (12, 24, 48 nM; 24 h) decreases the expression of cyclin B1, p-cdc2, Bcl-2 protein levels and increases the expression of cleaved PARP in a dose-dependent manner<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>HeLa, HEK-293, A549, MCF-7, T47D cells</td> </tr> <tr> <td>Concentration:</td> <td>0-100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Showed antiproliferative activity with IC<sub>50</sub>s of 0.012, &gt;100, 10.40, 40.40, 27.91 μM for HeLa, HEK-293, A549, MCF-7, T47D cells, respectively.</td> </tr> </table> <p>Cell Cycle Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>HeLa cells</td> </tr> <tr> <td>Concentration:</td> <td>12, 24, 48 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Induced cell cycle arrest at G2/M phase with the percentage of cells was 13.90%, 26.00%, and 92.65% at 12, 24, 48 nM, respectively.</td> </tr> </table> <p>Western Blot Analysis<sup>[1]</sup></p>	Cell Line:	HeLa, HEK-293, A549, MCF-7, T47D cells	Concentration:	0-100 μM	Incubation Time:	48 h	Result:	Showed antiproliferative activity with IC <sub>50</sub> s of 0.012, >100, 10.40, 40.40, 27.91 μM for HeLa, HEK-293, A549, MCF-7, T47D cells, respectively.	Cell Line:	HeLa cells	Concentration:	12, 24, 48 nM	Incubation Time:	24 h	Result:	Induced cell cycle arrest at G2/M phase with the percentage of cells was 13.90%, 26.00%, and 92.65% at 12, 24, 48 nM, respectively.
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Cell Line:	HeLa cells
Concentration:	12, 24, 48 nM
Incubation Time:	24 h
Result:	Concentration-dependently decreased cyclin B1 and p-cdc2 protein levels.

#### Apoptosis Analysis<sup>[1]</sup>

Cell Line:	HeLa cells
Concentration:	12, 24, 48 nM
Incubation Time:	24 h
Result:	Induced apoptosis with the total numbers of early and late apoptotic cells were 8.44%, 26.87% and 53.3% at 12, 24, and 48 nM, respectively.

## REFERENCES

[1]. Yang F, et al. Synthesis, and biological evaluation of 3,6-diaryl-[1,2,4]triazolo[4,3-a]pyridine analogues as new potent tubulin polymerization inhibitors. Eur J Med Chem. 2020 Oct 15;204:112625.

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

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