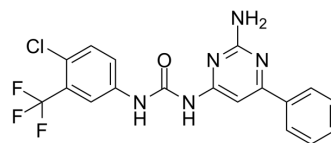


## Anticancer agent 71

Cat. No.:	HY-147906
CAS No.:	2453228-45-8
Molecular Formula:	C <sub>18</sub> H <sub>13</sub> ClF <sub>3</sub> N <sub>5</sub> O
Molecular Weight:	407.78
Target:	Apoptosis; PARP
Pathway:	Apoptosis; Cell Cycle/DNA Damage; Epigenetics
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Anticancer agent 71 (Compound 4b) is a potent anticancer agent and induces apoptosis. Anticancer agent 71 arrests cell cycle at G2/M phase and induces apoptosis through upregulating Bax, Ikb- $\alpha$ and cleaved PARP and downregulating Bcl-2 expression levels. Anticancer agent 71 shows antiproliferative activity <sup>[1]</sup> .																				
<b>In Vitro</b>	<p>Anticancer agent 71 (Compound 4b) (0.1-50 <math>\mu</math>M, 24 hours; SW480 and PC3 cells) increases the cytotoxicity and exerts potent antitumor activities against SW480 and PC3 cells<sup>[1]</sup>.</p> <p>Anticancer agent 71 (Compound 4b) (11.08 <math>\mu</math>M, 24 hours; SW480 cells) induces a significant G2/M phase arrest<sup>[1]</sup>.</p> <p>Anticancer agent 71 (Compound 4b) (11.08 <math>\mu</math>M, 24 hours) induces apoptosis in SW480 cells by upregulating Bax, Ikb-<math>\alpha</math> and cleaved PARP and downregulating Bcl-2 expression<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>SW480 and PC3 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.1, 1, 12.5, 25 and 50 <math>\mu</math>M</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibited with IC<sub>50</sub> values of 11.08 and 17.39 <math>\mu</math>M for SW480 and PC3 cells, respectively.</td> </tr> </table> <p>Cell Cycle Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>SW480 cells</td> </tr> <tr> <td>Concentration:</td> <td>11.08 <math>\mu</math>M</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>The cell population percent at G2/M phase increased to 45.3%.</td> </tr> </table> <p>Apoptosis Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>SW480 cells</td> </tr> <tr> <td>Concentration:</td> <td>11.08 <math>\mu</math>M</td> </tr> </table>	Cell Line:	SW480 and PC3 cells	Concentration:	0.1, 1, 12.5, 25 and 50 $\mu$ M	Incubation Time:	24 hours	Result:	Inhibited with IC <sub>50</sub> values of 11.08 and 17.39 $\mu$ M for SW480 and PC3 cells, respectively.	Cell Line:	SW480 cells	Concentration:	11.08 $\mu$ M	Incubation Time:	24 hours	Result:	The cell population percent at G2/M phase increased to 45.3%.	Cell Line:	SW480 cells	Concentration:	11.08 $\mu$ M
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Incubation Time:	24 hours
Result:	The percentage of cells in early and late apoptosis was recorded as 5.69% and 3.52%, respectively.
Western Blot Analysis <sup>[1]</sup>	
Cell Line:	SW480 cells
Concentration:	11.08 $\mu$ M
Incubation Time:	24 hours
Result:	The expression of antiapoptotic protein Bcl-2 decreased and the expression of Pro apoptotic Bax increased.

## REFERENCES

[1]. Kilic-Kurt Z, et al. Synthesis and anticancer activity of some pyrimidine derivatives with aryl urea moieties as apoptosis-inducing agents. Bioorg Chem. 2020 Aug;101:104028.

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

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