

Product Data Sheet

PI3K-IN-35

 Cat. No.:
 HY-147900

 CAS No.:
 2458163-99-8

 Molecular Formula:
 $C_{25}H_{23}N_7O_2$

Molecular Weight: 453.5

Target: PI3K; Apoptosis

Pathway: PI3K/Akt/mTOR; Apoptosis

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

BIOLOGICAL ACTIVITY

IC₅₀ & Target

PI3K-IN-35 (Compound 6l) is a highly selective PI3K inhibitor with IC₅₀ values of 13.98, 7.22 and 10.94 μM for PI3K-α⊠PI3K-β and PI3K-δ, respectively. PI3K-IN-35 arrests cell cycle at G2/M phase and induces apoptosis. PI3K-IN-35 can be used in leukemia research^[1].

ΡΙ3Κβ ΡΙ3Κδ ΡΙ3Κα

7.22 μ M (IC₅₀) 10.94 μ M (IC₅₀) 13.98 μ M (IC₅₀)

In Vitro PI3K-IN-35 (Compound 6l) (10 μM, 48 hours) possess potent antiproliferative activity can inhibit proliferation of Leukemia SR

cell lines in vitro^[1].

PI3K-IN-35 (Compound 6l) induces cell cycle arrest toward the G2-M phase (41.61%) and leads to DNA synthesis and apoptosis^[1].

PI3K-IN-35 (Compound 6l) (10 μ M, 48 hours) results in apoptotic with 16.98% of cells in the early and latest age [1].

 $\label{eq:mce} \mbox{MCE has not independently confirmed the accuracy of these methods. They are for reference only.}$

Cell Cytotoxicity Assay^[1]

Cell Line:	Leukemia SR cell lines.
Concentration:	10 μΜ
Incubation Time:	48 hours
Result:	Antiproliferative activity with values of 13.59 μM .

REFERENCES

[1]. Helwa AA, et al. Novel antiproliferative agents bearing morpholinopyrimidine scaffold as PI3K inhibitors and apoptosis inducers; design, synthesis and molecular docking. Bioorg Chem. 2020 Sep;102:104051.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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