

Product Data Sheet

T-1-PMPA

Cat. No.: HY-158149

CAS No.: 1323883-62-0

Molecular Formula: C₁₆H₁₇N₅O₃

Molecular Weight: 327.34

Target: EGFR; Apoptosis

Pathway: JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Apoptosis

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

Analysis.

BIOLOGICAL ACTIVITY

DescriptionT-1-PMPA is a potent EGFR inhibitor with apoptotic properties. T-1-PMPA effectively inhibits EGFR^{WT} and EGFR^{790m}, with IC

 $_{\rm 50}$ values of 86 nM and 561.73 nM, respectively $^{[1]}.$

IC₅₀ & Target EGFR^{WT} EGFR^{T790M}

86 nM (IC₅₀) 561.73 nM (IC₅₀)

In Vitro T-1-PMPA (0.312-10 μM; 24 h) shows significant suppression of the proliferation of HepG2 and MCF7 malignant cell lines, with

IC $_{50}$ values of 3.51 μM and 4.13 μM, respectively $^{[1]}$. In HepG2 cells, T-1-PMPA increases the proportion of cells in the early stage of apoptosis from 0.77 to 29.17%, the late stage of apoptosis from 0.17 to 8.81%, and the overall stage from 3.05 to 42.03%. Additionally, the percentage of necrotic cells increased to 4.05% compared to 2.21% in the control cells. The qRT-PCR analysis further supported the apoptotic effects by revealing significant increases in the levels of caspase-3 and caspase-9. T-1-PMPA controlls the levels of TNFα and IL2 by 74 and $50\%^{[1]}$.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Viability Assay^[1]

Cell Line:	HepG2 and MCF7 malignant cell lines
Concentration:	0.312 μΜ, 0.625 μΜ, 1.25 μΜ, 2.5 μΜ, 5 μΜ, 10 μΜ
Incubation Time:	24 h
Result:	Showed significant suppression of the proliferation of HepG2 and MCF7 malignant cell lines.

REFERENCES

[1]. Ibrahim H Eissa, et al. New Theobromine Apoptotic Analogue with Anticancer Potential Targeting the EGFR Protein: Computational and In Vitro Studies. ACS Omega. 2024 Mar 27;9(14):15861-15881.

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

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