

Product Data Sheet

PROTAC EGFR degrader 6

 Cat. No.:
 HY-146423

 CAS No.:
 2409793-28-6

 Molecular Formula:
 $C_{49}H_{57}FN_{12}O_5$

Molecular Weight: 913.05

Target: PROTACs; EGFR; Apoptosis

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

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

Analysis.

BIOLOGICAL ACTIVITY

Description	PROTAC EGFR degrader 6, a PROTAC EGFR degrader, potently degrades EGFR ^{Del19} in HCC827 cells with the DC ₅₀ of 45.2 nM. PROTAC EGFR degrader 6 significantly induces the apoptosis of HCC827 cells and arrest the cells in G1 phase ^[1] .
IC ₅₀ & Target	IC ₅₀ : 180 nM (EGFR ^{Del19} in HCC827 cells) ^[1] DC ₅₀ : 45.2 nM (EGFR ^{Del19} in HCC827 cells) ^[1]
In Vitro	PROTAC EGFR degrader 6 (compound 2) (0.001-10 μ M; 48 hours) exhibits potent EGFR degradation activity with certain concentration-dependent manner; and shows DC ₅₀ of 45.2 nM in HCC827 cells ^[1] . PROTAC EGFR degrader 6 (100 nM; 4-96 hours) degradates EGFR protein in a time-dependent manner, and reaches the maximum degradation rate (D _{max} = 87%) at 96 h ^[1] . PROTAC EGFR degrader 6 (0.1, 1 and 10 μ M; 48 hours) shows weak degradation activity on EGFR in H1975 and A549 cells ^[1] . PROTAC EGFR degrader 6 (0.1 and 1 μ M; 32 hours) induce HCC827 apoptosis, and the percentages of apoptosis cells are 53.72% and 32.31% at 0.1 and 1 μ M, respectively ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Hao Zhang, et al. Discovery of potent epidermal growth factor receptor (EGFR) degraders by proteolysis targeting chimera (PROTAC). Eur J Med Chem. 2020 Mar 1;189:112061.

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

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