

Product Data Sheet

EGFR-IN-97

Cat. No.: HY-157432 CAS No.: 3020681-05-1 Molecular Formula: $C_{37}H_{36}N_8O_2$ Molecular Weight: 624.73

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

Description	EGFR-IN-97 (compound 6q) is a EGFR inhibitor. EGFR-IN-97 shows inhibitory activity against Ba/F3-EGFR ^{L858R/T790M/C797S} and Ba/F3-EGFR ^{Del19/T790M/C797S} cells, with IC $_{50}$ values of 0.42 μ M and 0.41 μ M, respectively. EGFR-IN-97 can promote apoptosis of NCI-H1975-EGFR ^{L858R/T790M/C797S} cells at the concentration of 0.8 μ M ^[1] .
In Vitro	EGFR-IN-97 (compound 6q) shows excellent activity against mutant NSCLC cell line NCI-H1975-EGFR ^{L858R} /T790M/C797S cells, with IC ₅₀ value of 0.82 μ M, which was superior to that of osimertinib (HY-15772) (IC ₅₀ = 2.94 μ M), JBJ-04-125-02 (HY-135805) (IC ₅₀ = 3.66 μ M), and coadministration of JBJ-04-125-02 and osimertinib (IC ₅₀ = 1.25 μ M) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Hu L, et al. Identification of novel aminopyrimidine derivatives for the treatment of mutant NSCLC. Eur J Med Chem. 2023 Dec 20;265:116074.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA