

Product Data Sheet

$PI3K\alpha/HDAC6-IN-1$

 $\begin{tabular}{lll} \textbf{Cat. No.:} & HY-156091 \\ \textbf{CAS No.:} & 3007565-26-3 \\ \begin{tabular}{lll} \textbf{Molecular Formula:} & $C_{27}H_{30}F_3N_7O_6S_2$ \\ \end{tabular}$

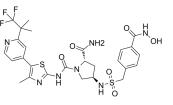
Molecular Weight: 669.7

Target: PI3K; HDAC

Pathway: PI3K/Akt/mTOR; Cell Cycle/DNA Damage; Epigenetics

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

Analysis.



BIOLOGICAL ACTIVITY

Description	PI3Kα/HDAC6-IN-1 (compound 21j) is a dual PI3Kα/HDAC6 inhibitor with IC $_{50}$ of 2.9 and 26 nM, respectively. PI3Kα/HDAC6-IN-1 also inhibits AKT(Ser473) phosphorylation and induces the accumulation of acetylated α-tubulin without affecting acetylated histones H3 and H4. PI3Kα/HDAC6-IN-1 efficiently inhibits L-363 cell line (IC $_{50}$ =0.17 μM) and has good anti-cancer activity ^[1] .
IC ₅₀ & Target	IC50: 2.9 nM (PI3Kα), 26 nM (HDAC6) ^[1]

REFERENCES

[1]. Zhang Y, et al. Discovery of (S)-N1-(thiazol-2-yl) pyrrolidine-1,2-dicarboxamide derivatives targeting PI3Ka/HDAC6 for the treatment of cancer. Bioorg Med Chem Lett. 2023 Oct 1;94:129462...

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

Inhibitors