# **PI3K-IN-37**

Cat. No.: HY-147284 CAS No.: 1257547-40-2 Molecular Formula:  $C_{25}H_{26}N_6O_2$ Molecular Weight: 442.51 Target: PI3K; mTOR

Pathway: PI3K/Akt/mTOR

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

Analysis.

**Product** Data Sheet

## **BIOLOGICAL ACTIVITY**

Description PI3K-IN-37 (Example 84.1) is a PI3K  $\alpha/\beta/\delta$  inhibitor with IC<sub>50</sub>s of 6, 8, 4 nM, respectively. PI3K-IN-37 can also inhibit mTOR (IC <sub>50</sub>=4 nM)<sup>[1]</sup>.

IC<sub>50</sub> & Target ΡΙ3Κδ mTOR ΡΙ3Κα ΡΙ3Κβ 4 nM (IC<sub>50</sub>) 4 nM (IC<sub>50</sub>) 6 nM (IC<sub>50</sub>) 8 nM (IC<sub>50</sub>)

PI3K-IN-37 (0.005-10  $\mu$ M; 30 min) treatment inhibits PKB Ser473 phosphorylation in Rat1 cells<sup>[1]</sup>. In Vitro

 $P13K-IN-37\ (0.005-1\ \mu\text{M}; 60\ min)\ treatment\ inhibits\ S6\ Ribosomal\ Protein\ phosphorylation\ Ser 235/236\ in\ TSC1\ nult\ MEF\ cells$ 

[1].

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

Cell Viability Assay<sup>[1]</sup>

Cell Line:	Rat1 cells
Concentration:	0.005-10 μΜ
Incubation Time:	30 min
Result:	Inhibited Pl3-kinase isoforms alpha, beta or delta phosphorylation with the IC <sub>50</sub> s of <5 nM, 8 nM, and 18 nM, respectively.

### Cell Viability Assay[1]

Cell Line:	TSC1-/- MEF cells
Concentration:	0.005-1 μM
Incubation Time:	60 min
Result:	Inhibited S6 Ribosomal Protein phosphorylation (IC <sub>50</sub> =2 nM).

### **REFERENCES**

[1]. FURET Pascal, et al. 1H-IMIDAZO[4,5-c]QUINOLINONE DERIVATIVES. WO2010139731A1.						
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