ΡΙ3Κα-ΙΝ-8

| Cat. No.: | HY-147983 | |
|--------------------|---|------|
| CAS No.: | 2315320-24-0 | Br |
| Molecular Formula: | C ₂₆ H ₂₇ BrN ₄ O ₂ | ■ NH |
| Molecular Weight: | 507.42 | |
| Target: | PI3K; Reactive Oxygen Species; Apoptosis | |
| Pathway: | PI3K/Akt/mTOR; Immunology/Inflammation; Metabolic Enzyme/Protease; NF-кВ; Apoptosis | |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. | |

| BIOLOGICAL ACTIVITY | | | | | |
|---------------------|--|--------------------------------------|--------------------------------------|--------------------------------------|--|
| Description | PI3Kα-IN-8 (Compound 9g) is a selective PI3Kα inhibitor with an IC ₅₀ of 0.012 μM. PI3Kα-IN-8 increases intracellular reactive oxygen species level, decreases mitochondrial membrane potential and induces apoptosis ^[1] . | | | | |
| IC₅₀ & Target | ΡΙ3Κα 0.012 μΜ (IC ₅₀) | ΡΙ3Κδ 0.11 μΜ (IC ₅₀) | ΡΙ3Κγ 0.18 μΜ (IC ₅₀) | РІЗКβ 0.21 μM (IC ₅₀) | |
| In Vitro | PI3Kα-IN-8 (Compound 9g) exhibits anticancer activity with IC ₅₀ values of 0.18 ± 0.03, 0.43 ± 0.05, 0.71 ± 0.08 and 0.63 ± 0.09 μ M against HCT-116, MCF-7, HeLa and HepG2 cells, respectively ^[1] . PI3Kα-IN-8 upregulates Bax and cleaved caspase-3/9 levels, and downregulates Bcl-2 level ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | | | | |

REFERENCES

[1]. Yang YQ, et al. Synthesis and anticancer evaluation of novel 1H-benzo[d]imidazole derivatives of dehydroabietic acid as PI3Ka inhibitors. Bioorg Chem. 2020 Jul;100:103845.

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

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Product Data Sheet

