Inhibitors

Screening Libraries

Product Data Sheet

5-NIdR

Pathway:

Cat. No.: HY-115567 CAS No.: 191421-10-0 Molecular Formula: $C_{13}H_{14}N_{2}O_{5}$ Molecular Weight: 278.26 Target: **Apoptosis**

Storage: Powder

-20°C 3 years 4°C 2 years

In solvent -80°C 6 months

Apoptosis

-20°C 1 month

BIOLOGICAL ACTIVITY

Description	5-NIdR (1-(β-D-2-Deoxyribofuranosyl)-5-nitroindole), an artificial nucleoside, exhibits the ability to inhibit the replication of DNA lesions generated by Temozolomide (HY-17364). 5-NIdR induces cancer cells apoptosis and arrests cell cycle at G0 phase. 5-NIdR enhances Temozolomide anti-tumor efficacy in murine glioblastoma model ^[1] .
In Vitro	5-NIdR (12.5-100 μM; 24-72 h) inhibits the growth of human glioblastoma cell lines (U87, A172, and SW1088) in a dose-dependent manner ^[1] . 5-NIdR (1-100 μg/mL; 72 h) induces cell apoptosis in U87 cells, and arrests cell cycle at G0 phase with 100 μg/mL overnight incubation ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	5-NIdR (100 mg/kg; ip; for 5 consecutive days), together with Temozolomide (40 mg/kg), results complete tumor regression in a murine xenograft model of glioblastoma. Temozolomide alone only delayed tumor growth ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Choi JS, et al. Inhibition of Translesion DNA Synthesis as a Novel Therapeutic Strategy to Treat Brain Cancer. Cancer Res. 2018 Feb 15;78(4):1083-1096.

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

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