GSK 690

Cat. No.: HY-117226 CAS No.: 2101305-84-2 Molecular Formula: $C_{24}H_{23}N_3O$ Molecular Weight: 369.46

Target: Histone Demethylase

Pathway: **Epigenetics**

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

Product Data Sheet

BIOLOGICAL ACTIVITY

Description	GSK 690 is a reversible inhibitor of lysine specific demethylase 1 (LSD1), with a K _d value of 9 nM and a biochemical IC ₅₀ of 37 nM.
In Vitro	GSK690 (1-10 μ M) acts together with JNJ-26481585 to induce cell death in all four tested RMS cells lines (RD, RH30, RMS13, and TE381.T cells) ^[2] . GSK690/JNJ-26481585 cotreatment alters the balance between pro- and antiapoptotic proteins with 1 μ M GSK690 for RD cells) and 10 μ M GSK690 for RH30 cells ^[2] . GSK690/JNJ-26481585 cotreatment induces caspase-dependent cell death with 1 μ M GSK690 for RD cells and 10 μ M GSK690 for RH30 cells ^[2] . The addition of GSK690 further enhances the JNJ-26481585-stimulated G2/M arrest ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

• Oncogene. 2021 Mar 12.

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REFERENCES

[1]. Mould DP, et al. Development of (4-Cyanophenyl)glycine Derivatives as Reversible Inhibitors of Lysine Specific Demethylase 1. J Med Chem. 2017 Oct 12;60(19):7984-7999.

[2]. Haydn T, et al. Concomitant epigenetic targeting of LSD1 and HDAC synergistically induces mitochondrial apoptosis in rhabdomyosarcoma cells. Cell Death Dis. 2017 Jun 15;8(6):e2879.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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