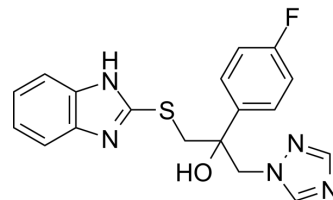


Antifungal agent 51

Cat. No.:	HY-149822
CAS No.:	2896209-47-3
Molecular Formula:	C ₁₈ H ₁₆ FN ₃ OS
Molecular Weight:	369.42
Target:	Fungal
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Antifungal agent 51 (Compound 5c) has potent antifungal activity, especially against <i>Candida albicans</i> FDC 151, <i>Candida parapsilosis</i> ATCC 22019 and <i>Candida tropicalis</i> FDC 138, with the MIC value is less than 0.063 µg/mL, and it has low toxicity to cells and no carcinogenicity ^[1] .								
In Vitro	<p>Antifungal agent 51 has potent antifungal activity, especially against <i>Candida albicans</i> FDC 151, <i>Candida parapsilosis</i> ATCC 22019 and <i>Candida tropicalis</i> FDC 138, with the MIC value is less than 0.063 µg/mL, and against <i>Candida albicans</i> CMRC 19 and <i>Candida glabrata</i> FDC 192 with MIC value of 0.125 µg/mL, against <i>Candida albicans</i> CMRC 192, with MIC value of 0.25 µg/mL, against <i>Candida glabrata</i> CMRC 89, with MIC value of 1 µg/mL^[1].</p> <p>Antifungal agent 51 (6.25, 12.5, 25 µg/mL; 48 h) partially affects the viability of cell lines even at relatively high concentration of 25 µg/mL in BEAS-2B and HepG2 cells^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 30%;">Cell Line:</td> <td>BEAS-2B and HepG2 cells</td> </tr> <tr> <td>Concentration:</td> <td>6.25, 12.5, and 25 µg/mL</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Partially affected the viability of cell lines even at relatively high concentration of 25 µg/mL in BEAS-2B and HepG2 cells</td> </tr> </table>	Cell Line:	BEAS-2B and HepG2 cells	Concentration:	6.25, 12.5, and 25 µg/mL	Incubation Time:	48 h	Result:	Partially affected the viability of cell lines even at relatively high concentration of 25 µg/mL in BEAS-2B and HepG2 cells
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Concentration:	6.25, 12.5, and 25 µg/mL								
Incubation Time:	48 h								
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REFERENCES

[1]. Ghobadi E, et al. Design, synthesis and biological activity of hybrid antifungals derived from fluconazole and mebendazole. *Eur J Med Chem.* 2023 Mar 5;249:115146.

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

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