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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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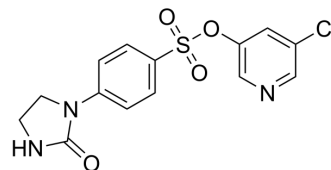
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PYRIB-SO 2

Cat. No.:	HY-163523
Molecular Formula:	C ₁₄ H ₁₂ ClN ₃ O ₄ S
Molecular Weight:	353.78
Target:	Microtubule/Tubulin
Pathway:	Cell Cycle/DNA Damage; Cytoskeleton
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	PYRIB-SO 2 is a potent antimetabolic agent. PYRIB-SO 2 shows antiproliferative activity and induces cell cycle arrest at G2/M phase. PYRIB-SO 2 reduces and disrupts microtubule structures. PYRIB-SO 2 binds to the colchicine-binding site (C-BS) of α , β -tubulin ^[1] .																
In Vitro	<p>PYRIB-SO 2 (150 nM; 24 h) induces cell cycle arrest at G2/M phase ^[1].</p> <p>PYRIB-SO 2 (138 nM; 24 h) reduces and disrupts microtubule structures in MCF7 cells ^[1].</p> <p>PYRIB-SO 2 (5, 50 μM) reduces the formation of the EB1:β-tubulin adduct ^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MCF7, MDA-MB-468, MDA-MB-231 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-500 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Showed antiproliferative activity with IC₅₀s of 50, 9, 220 nM for MCF7, MDA-MB-468 and MDA-MB-231 cells, respectively.</td> </tr> </table> <p>Cell Cycle Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MCF7 cells</td> </tr> <tr> <td>Concentration:</td> <td>150 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Induced cell cycle arrest at G2/M phase with the percentage of 77%.</td> </tr> </table>	Cell Line:	MCF7, MDA-MB-468, MDA-MB-231 cells	Concentration:	0-500 nM	Incubation Time:	48 h	Result:	Showed antiproliferative activity with IC ₅₀ s of 50, 9, 220 nM for MCF7, MDA-MB-468 and MDA-MB-231 cells, respectively.	Cell Line:	MCF7 cells	Concentration:	150 nM	Incubation Time:	24 h	Result:	Induced cell cycle arrest at G2/M phase with the percentage of 77%.
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REFERENCES

[1]. Ouellette V, et al. Modification of the phenyl ring B of phenyl 4-(2-oxoimidazolidin-1-yl)benzenesulfonates by pyridinyl moiety leads to novel antimetotics targeting the colchicine-binding site. *Bioorg Med Chem Lett*. 2024 Jun 1;105:129745.

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

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