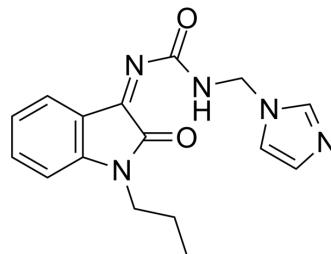


COX-2/PI3K-IN-2

Cat. No.:	HY-147912
CAS No.:	2459938-28-2
Molecular Formula:	C ₁₆ H ₁₇ N ₅ O ₂
Molecular Weight:	311.34
Target:	PI3K; COX
Pathway:	PI3K/Akt/mTOR; Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	COX-2/PI3K-IN-2 (compound 5f) is a potent PI3K inhibitor with IC ₅₀ value of 2.78 nM. COX-2/PI3K-IN-2 is a selective COX-2 inhibitor with K _i value of 3.02 nM. COX-2/PI3K-IN-2 shows anti-inflammatory and anti-cancer properties ^[1] .									
IC₅₀ & Target	PI3K 2.78 nM (IC ₅₀)	COX-2 3.02 nM (K _i)								
In Vitro	<p>COX-2/PI3K-IN-2 (compound 5f) (50-500 µg/mL, 72 hours) exerts potent antitumor activities against Breast cancer (MCF-7) cells^[1].</p> <p>COX-2/PI3K-IN-2 (compound 5f) (10-150 µg/mL) has antioxidant properties with H₂O₂ scavenging activity equivalent to ascorbic acid and higher lysozyme inhibition (IC₅₀ 2.25 nM)^[1].</p> <p><small>IC₅₀ 1.65 µM independently confirmed the accuracy of these methods. They are for reference only.</small></p> <p><small>Cell Cytotoxicity Assay^[1]</small></p> <table border="1"> <tr> <td>Cell Line:</td> <td>Breast cancer (MCF-7) cells and normal breast epithelial (MCF-10A) cells</td> </tr> <tr> <td>Concentration:</td> <td>50-500 µg/mL</td> </tr> <tr> <td>Incubation Time:</td> <td>72 hours</td> </tr> <tr> <td>Result:</td> <td>Caused 40 % cell death at 1.65 µM in MCF-7 cells.</td> </tr> </table>		Cell Line:	Breast cancer (MCF-7) cells and normal breast epithelial (MCF-10A) cells	Concentration:	50-500 µg/mL	Incubation Time:	72 hours	Result:	Caused 40 % cell death at 1.65 µM in MCF-7 cells.
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REFERENCES

[1]. Rajesh Kumar M, et al. p-TSA.H₂O mediated one-pot, multi-component synthesis of isatin derived imidazoles as dual-purpose drugs against inflammation and cancer. Bioorg Chem. 2020 Sep;102:104046.

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

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