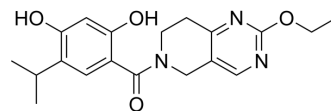


CPUY201112

Cat. No.:	HY-111524
CAS No.:	1860793-58-3
Molecular Formula:	C ₁₉ H ₂₃ N ₃ O ₄
Molecular Weight:	357.4
Target:	HSP; Apoptosis
Pathway:	Cell Cycle/DNA Damage; Metabolic Enzyme/Protease; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	CPUY201112 is a potent heat shock protein Hsp90 inhibitor with K _d of 27 nM. CPUY201112 induces p53-mediated apoptosis in MCF-7 cells, resulting in cell cycle arrest, which can be used in cancer research ^[1] .																
In Vitro	<p>CPUY201112 (0-9 μM, 7 days) reduces the viability of multiple cancer cell lines such as HCT116 colon cells, HepG2 hepatocellular carcinoma and other cancer cells in a dose-dependent manner^[1].</p> <p>CPUY201112 (0-2 μM, 24 h) can induce apoptosis in a dose-dependent manner^[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>HCT116, HepG2, MCF-7, A549</td> </tr> <tr> <td>Concentration:</td> <td>0-9 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>7 days</td> </tr> <tr> <td>Result:</td> <td>Inhibited MCF-7, A549, HCT116 and HepG2 cells with the IC₅₀ values of 0.624, 0.543, 0.763 and 0.342 μM, respectively.</td> </tr> </table> <p>Apoptosis Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MCF-7 and HCT116 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-2 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Induced cell cycle arrest in G2/M phase and induced apoptosis in more than 35% of MCF-7 cells. Induced p53-mediated apoptosis in HCT116 cells.</td> </tr> </table>	Cell Line:	HCT116, HepG2, MCF-7, A549	Concentration:	0-9 μM	Incubation Time:	7 days	Result:	Inhibited MCF-7, A549, HCT116 and HepG2 cells with the IC ₅₀ values of 0.624, 0.543, 0.763 and 0.342 μM, respectively.	Cell Line:	MCF-7 and HCT116 cells	Concentration:	0-2 μM	Incubation Time:	24 h	Result:	Induced cell cycle arrest in G2/M phase and induced apoptosis in more than 35% of MCF-7 cells. Induced p53-mediated apoptosis in HCT116 cells.
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In Vivo	<p>CPUY201112 (5-40 mg/kg, i.p., daily, 3 weeks) inhibits tumor growth in the MCF-7 tumor xenograft model^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Female BALB/c nude mice with MCF-7 cell^[1]</td> </tr> </table>	Animal Model:	Female BALB/c nude mice with MCF-7 cell ^[1]														
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Dosage:	5 mg/kg, 20 mg/kg and 40 mg/kg
Administration:	i.p., daily, 3 weeks
Result:	Reduced tumor volume by 11.92%, 26.58% and 39.63%, respectively, when using 5 mg/kg, 20 mg/kg and 40 mg/kg. Significantly induced the expression of Hsp70 and reduced the expression of Akt at 40 mg/kg.

REFERENCES

[1]. Xiao-Li X, et al. CPUY201112, a novel synthetic small-molecule compound and inhibitor of heat shock protein Hsp90, induces p53-mediated apoptosis in MCF-7 cells. Sci Rep. 2016 Jan 8;6:19004.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA