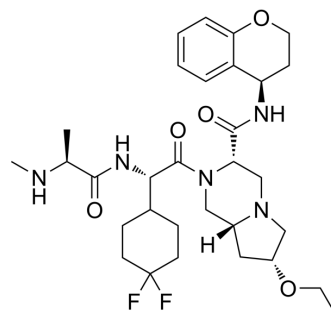


T-3256336

Cat. No.:	HY-100682
CAS No.:	1266227-69-3
Molecular Formula:	C ₃₁ H ₄₅ F ₂ N ₅ O ₅
Molecular Weight:	605.72
Target:	IAP
Pathway:	Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	T-3256336 is a potent and orally active cIAP1 and XIAP inhibitor with IC ₅₀ s of 1.3, 200 nM, respectively. T-3256336 shows anti-tumor activity ^[1] .									
IC₅₀ & Target	cIAP1 1.3 nM (IC ₅₀)	XIAP 200 nM (IC ₅₀)								
In Vivo	<p>T-3256336 (compound 45) (30, 100 mg/kg; p.o.) increases caspase-3/7 activity and shows anti-tumor activity in mouse^[1]. 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/cAJcl mice (MDA-MB-231-Luc xenograft)^[1]</td> </tr> <tr> <td>Dosage:</td> <td>30, 100 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>P.o.</td> </tr> <tr> <td>Result:</td> <td>Stimulated caspase-3/7 activity in a dose-dependent manner, induced tumor regression in a dose-dependent manner with a percent tumor growth inhibition (T/C) of 53% and 62% at 30 mg/kg and 100 mg/kg, respectively.</td> </tr> </table>		Animal Model:	Female BALB/cAJcl mice (MDA-MB-231-Luc xenograft) ^[1]	Dosage:	30, 100 mg/kg	Administration:	P.o.	Result:	Stimulated caspase-3/7 activity in a dose-dependent manner, induced tumor regression in a dose-dependent manner with a percent tumor growth inhibition (T/C) of 53% and 62% at 30 mg/kg and 100 mg/kg, respectively.
Animal Model:	Female BALB/cAJcl mice (MDA-MB-231-Luc xenograft) ^[1]									
Dosage:	30, 100 mg/kg									
Administration:	P.o.									
Result:	Stimulated caspase-3/7 activity in a dose-dependent manner, induced tumor regression in a dose-dependent manner with a percent tumor growth inhibition (T/C) of 53% and 62% at 30 mg/kg and 100 mg/kg, respectively.									

REFERENCES

[1]. Hashimoto K, et al. Design and synthesis of potent inhibitor of apoptosis (IAP) proteins antagonists bearing an octahydropyrrolo[1,2-a]pyrazine scaffold as a novel proline mimetic. J Med Chem. 2013 Feb 14;56(3):1228-46.

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