



SZABO SCANDIC

Part of Europa Biosite

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

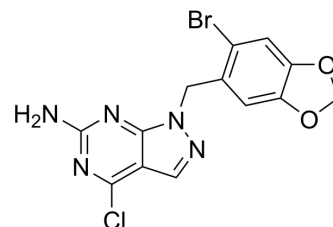
mail@szabo-scandic.com

www.szabo-scandic.com

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

DN401

Cat. No.:	HY-115781
CAS No.:	2135749-60-7
Molecular Formula:	C ₁₃ H ₉ BrClN ₅ O ₂
Molecular Weight:	382.6
Target:	HSP
Pathway:	Cell Cycle/DNA Damage; Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	DN401 is a potent TRAP1 inhibitor with an IC ₅₀ of 79 nM. DN401 shows a weak inhibition of Hsp90 (IC ₅₀ of 698 nM). DN401 inactivates mitochondrial TRAP1 and has potent anticancer activities ^[1] .																	
IC₅₀ & Target	TRAP1 79 nM (IC ₅₀)	HSP90 698 nM (IC ₅₀)																
In Vitro	<p>DN401 (5-20 μM; 24 hours) has superior cancer-specific cytotoxicity but reduced cytotoxic effects in normal cells^[1]. DN401 (20 μM; 6 hours) inhibits the expression of Chk1 and Akt in HeLa cells^[1]. 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"> <tr> <td>Cell Line:</td> <td>HeLa, SK-HEP-1, T98G, H460, and PC3 cells</td> </tr> <tr> <td>Concentration:</td> <td>5 μM, 10 μM, 15 μM, 20 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Exhibited cytotoxic activity.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HeLa cells</td> </tr> <tr> <td>Concentration:</td> <td>20 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>6 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibited the expression of Chk1 and Akt.</td> </tr> </table>		Cell Line:	HeLa, SK-HEP-1, T98G, H460, and PC3 cells	Concentration:	5 μM, 10 μM, 15 μM, 20 μM	Incubation Time:	24 hours	Result:	Exhibited cytotoxic activity.	Cell Line:	HeLa cells	Concentration:	20 μM	Incubation Time:	6 hours	Result:	Inhibited the expression of Chk1 and Akt.
Cell Line:	HeLa, SK-HEP-1, T98G, H460, and PC3 cells																	
Concentration:	5 μM, 10 μM, 15 μM, 20 μM																	
Incubation Time:	24 hours																	
Result:	Exhibited cytotoxic activity.																	
Cell Line:	HeLa cells																	
Concentration:	20 μM																	
Incubation Time:	6 hours																	
Result:	Inhibited the expression of Chk1 and Akt.																	
In Vivo	<p>DN401 (30 mg/kg; ip; once daily; for 14 days) reduces tumor growth and increases apoptotic cell death in tumor tissues, and does not cause weight loss or histologic abnormalities^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>																	

Animal Model:	6-week-old BALB/c nu/nu male mice injected with PC3 cells ^[1]
Dosage:	30 mg/kg
Administration:	ip; once daily; for 14 days
Result:	Reduced tumor growth.

REFERENCES

[1]. Hye-Kyung Park, et al. Paralog Specificity Determines Subcellular Distribution, Action Mechanism, and Anticancer Activity of TRAP1 Inhibitors. J Med Chem. 2017 Sep 14;60(17):7569-7578.

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