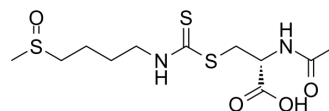


DL-Sulforaphane N-acetyl-L-cysteine

Cat. No.:	HY-132242
CAS No.:	334829-66-2
Molecular Formula:	C ₁₁ H ₂₀ N ₂ O ₄ S ₃
Molecular Weight:	340.48
Target:	HDAC; Apoptosis
Pathway:	Cell Cycle/DNA Damage; Epigenetics; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	DL-Sulforaphane N-acetyl-L-cysteine (SFN-NAC) is an orally active HDAC inhibitor and metabolite of sulforaphane (HY-13755) with longer half-life and better blood-brain barrier permeability. DL-Sulforaphane N-acetyl-L-cysteine activates autophagy-mediated downregulation of α -tubulin expression through the ERK pathway and can be used in cancer research ^{[1][2]} .																						
In Vitro	<p>SFN-NAC (24 h) decreases cell viability, with IC₅₀ values of 60.08 μM for HA, 35.20 μM for U87MG, 39.11 μM for U373MG, and 36.20 μM for the U87/TR cells^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cycle Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>U87MG and U373MG cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 30 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Induced cell-cycle arrest in the G2/M phase and triggered apoptosis at the same time.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>U87MG and U373MG cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 10, 20, 30, 40, 50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Activated ERK1/2 (Thr202/Tyr204), downregulated α-tubulin, and induced autophagy in a dose-dependent manner.</td> </tr> </table> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HA, U87MG, U373MG and U87/TR cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 10, 20, 30, 40, 50, 60, 70, 80, 90 μM for HA and U87MG cells or 0, 10, 20, 30, 40, 50, 60 μM for U373MG and U87/TR cells</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> </table>	Cell Line:	U87MG and U373MG cells	Concentration:	0, 30 μ M	Incubation Time:	24 h	Result:	Induced cell-cycle arrest in the G2/M phase and triggered apoptosis at the same time.	Cell Line:	U87MG and U373MG cells	Concentration:	0, 10, 20, 30, 40, 50 μ M	Incubation Time:	24 h	Result:	Activated ERK1/2 (Thr202/Tyr204), downregulated α -tubulin, and induced autophagy in a dose-dependent manner.	Cell Line:	HA, U87MG, U373MG and U87/TR cells	Concentration:	0, 10, 20, 30, 40, 50, 60, 70, 80, 90 μ M for HA and U87MG cells or 0, 10, 20, 30, 40, 50, 60 μ M for U373MG and U87/TR cells	Incubation Time:	24 h
Cell Line:	U87MG and U373MG cells																						
Concentration:	0, 30 μ M																						
Incubation Time:	24 h																						
Result:	Induced cell-cycle arrest in the G2/M phase and triggered apoptosis at the same time.																						
Cell Line:	U87MG and U373MG cells																						
Concentration:	0, 10, 20, 30, 40, 50 μ M																						
Incubation Time:	24 h																						
Result:	Activated ERK1/2 (Thr202/Tyr204), downregulated α -tubulin, and induced autophagy in a dose-dependent manner.																						
Cell Line:	HA, U87MG, U373MG and U87/TR cells																						
Concentration:	0, 10, 20, 30, 40, 50, 60, 70, 80, 90 μ M for HA and U87MG cells or 0, 10, 20, 30, 40, 50, 60 μ M for U373MG and U87/TR cells																						
Incubation Time:	24 h																						

	Result:	Decreased the cell viability of these cell lines in a dose-dependent manner.
In Vivo	SFN-NAC (10 µmol; 6 h; Oral gavage; single dose) significantly inhibits HDAC activity in mouse colon mucosa ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	mice ^[2]
	Dosage:	10 µmol
	Administration:	Oral gavage; 6 h; single dose
	Result:	Significantly inhibited HDAC activity in mouse colon mucosa.

REFERENCES

- [1]. Liu HJ, et al. Sulforaphane-N-Acetyl-Cysteine Induces Autophagy Through Activation of ERK1/2 in U87MG and U373MG Cells. *Cell Physiol Biochem*. 2018;51(2):528-542.
- [2]. Dashwood RH, et al. Dietary histone deacetylase inhibitors: from cells to mice to man[C]//Seminars in cancer biology. Academic Press, 2007, 17(5): 363-369.

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