Product Data Sheet

Antitumor agent-82

Cat. No.: HY-151914 $\label{eq:hybrid} \mbox{Molecular Formula:} \qquad \mbox{C_{32}H}_{42}\mbox{N_6}$

Molecular Weight: 510.72

Target: Autophagy; Atg7

Pathway: Autophagy

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

BIOLOGICAL ACTIVITY

Description

Antitumor agent-82 (compound 6g) is a potent anti-tumor agent. Antitumor agent-82 shows anti-proliferative activity. Antitumor agent-82 induces cell Autophagy by the ATG5/ATG7 signaling pathway^[1].

In Vitro

Antitumor agent-82 (compound 6g) (0-100 μ M; 48 h) shows anti-proliferative activity with IC₅₀s of 24.8, 13.5, 11.5, 2.71, 2.02, 4.53 μ M for BGC-823, MCF7, A375, 786-0, HT-29, Blu-87 cells, respectively^[1].

Antitumor agent-82 (0-4 μ M; 0-7 days) inhibits cell growth in a dose and time-dependent manner against HCT116 cells^[1]. Antitumor agent-82 (0-5 μ M; 0-60 h) activates autophagy by the ATG5/ATG7 signaling pathway in HCT116 cells^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Proliferation Assay^[1]

Cell Line:	BGC-823, MCF7, A375, 786-O, HT-29, Blu-87 cells
Concentration:	0-100 μΜ
Incubation Time:	48 h
Result:	Showed antiproliferative activity with IC $_{50}$ s of 24.8, 13.5, 11.5, 2.71, 2.02, 4.53 μ M for BGC-823, MCF7, A375, 786-O, HT-29, Blu-87 cells, respectively.

Western Blot Analysis^[1]

Cell Line:	HCT116 cells
Concentration:	0-5 μΜ
Incubation Time:	0-60 h
Result:	Induced autophagy with no influences on the expression of caspase-3, cleaved caspase-3 and p53 protein, significantly increased the expression of LC3-II and p62.

In Vivo

Antitumor agent-82 (45 mg/kg; i.p.; every two days for 16 days) shows anti-cancer activity for mouse^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	6-week-old male BALB/c mice $^{[1]}$

Dosage:	45 mg/kg
Administration:	I.p.; every two days for 16 days
Result:	Reduced tumor volume and resulted in a considerable reduction of the tumor weight of 69.69%.

REFERENCES

[1]. Ao J, et al. Design, synthesis and pharmacological evaluation of β -carboline derivatives as potential antitumor agent via targeting autophagy. Eur J Med Chem. 2022 Nov 26;246:114955.

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

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