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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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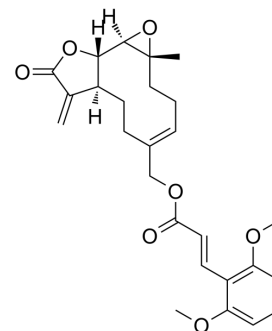
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anti-TNBC agent-1

Cat. No.:	HY-145143
CAS No.:	2289585-58-4
Molecular Formula:	C ₂₆ H ₃₀ O ₇
Molecular Weight:	454.51
Target:	Apoptosis
Pathway:	Apoptosis
Storage:	-20°C, protect from light, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light, stored under nitrogen)



SOLVENT & SOLUBILITY

In Vitro

DMSO : 50 mg/mL (110.01 mM; Need ultrasonic)

Concentration	Solvent	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.2002 mL	11.0009 mL	22.0017 mL
	5 mM	0.4400 mL	2.2002 mL	4.4003 mL
	10 mM	0.2200 mL	1.1001 mL	2.2002 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description

anti-TNBC agent-1 is a potent anti-triple-negative breast cancer (TNBC) agent. anti-TNBC agent-1 exhibits potent activity against different breast cancer cells with IC₅₀ values ranging from 0.20 μM to 0.27 μM. anti-TNBC agent-1 induces apoptosis of SUM-159 cells through mitochondria pathway and causes G1 phase arrest of SUM-159 cells^[1].

In Vitro

anti-TNBC agent-1 (compound 7) exhibits potent activity against MDA-MB231, SUM-159, MCF-7, Bcap-37, 4T1 cells with IC₅₀ values ranging from 0.20 μM to 0.27 μM. anti-TNBC agent-1 shows 11.6- to 18.6-fold improvement comparing to that of the parent compound parthenolide with IC₅₀ values of 2.68-4.63 μM. anti-TNBC agent-1 is more active than the positive control drug Adriamycin (ADR)^[1].

anti-TNBC agent-1 (2 μM and 5 μM; 48 hours; SUM-159 cells) exhibits significant stronger effect on induction of cell apoptosis compared with that of Parthenolide^[1].

anti-TNBC agent-1 has selective cytotoxicity against breast cancer cells (IC₅₀=0.22 μM) being compared with 3T3 cells (IC₅₀=8.13 μM)^[1].

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

REFERENCES

[1]. Ge W, et al. Synthesis and structure-activity relationship studies of parthenolide derivatives as potential anti-triple negative breast cancer agents. Eur J Med Chem. 2019;166:445-469.

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

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