# **Product** Data Sheet

# Resveratrol

Cat. No.: HY-16561 CAS No.: 501-36-0 Molecular Formula:  $C_{14}H_{12}O_3$ 

Molecular Weight: 228.24

Target: IKK; Autophagy; Mitophagy; Sirtuin; Apoptosis; Bacterial; Fungal; Antibiotic; Keap1-

NF-κΒ; Autophagy; Cell Cycle/DNA Damage; Epigenetics; Apoptosis; Anti-infection Pathway:

Storage: Powder -20°C 3 years

4°C 2 years

-80°C 2 years In solvent -20°C 1 year

#### **SOLVENT & SOLUBILITY**

In Vitro DMSO: 100 mg/mL (438.14 mM; Need ultrasonic)

Ethanol: 50 mg/mL (219.07 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	4.3814 mL	21.9068 mL	43.8135 mL
	5 mM	0.8763 mL	4.3814 mL	8.7627 mL
	10 mM	0.4381 mL	2.1907 mL	4.3814 mL

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

In Vivo

- 1. Add each solvent one by one: 0.5% CMC-Na/saline water Solubility: 16.67 mg/mL (73.04 mM); Suspended solution; Need ultrasonic
- 2. Add each solvent one by one: 50% PEG300 >> 50% saline Solubility: 12.5 mg/mL (54.77 mM); Suspended solution; Need ultrasonic
- 3. Add each solvent one by one: 10% EtOH >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 5 mg/mL (21.91 mM); Clear solution; Need ultrasonic
- 4. Add each solvent one by one: 10% EtOH >> 90% (20% SBE-β-CD in saline) Solubility: 5 mg/mL (21.91 mM); Clear solution; Need ultrasonic
- 5. Add each solvent one by one: 10% EtOH >> 90% corn oil Solubility: ≥ 5 mg/mL (21.91 mM); Clear solution
- 6. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (10.95 mM); Clear solution
- 7. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (10.95 mM); Clear solution

8. Add each solvent one by one: 10% DMSO >> 90% corn oil

Solubility: ≥ 2.5 mg/mL (10.95 mM); Clear solution

- 9. Add each solvent one by one: 5% DMSO >> 40% PEG300 >> 5% Tween-80 >> 50% saline Solubility: ≥ 2.5 mg/mL (10.95 mM); Clear solution
- 10. Add each solvent one by one: 5% DMSO >> 95% (20% SBE- $\beta$ -CD in saline) Solubility:  $\geq$  2.5 mg/mL (10.95 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

#### Description

Resveratrol (trans-Resveratrol; SRT501), a natural polyphenolic phytoalexin that possesses anti-oxidant, anti-inflammatory, cardioprotective, and anti-cancer properties. Resveratrol (SRT 501) has a wide spectrum of targets including mTOR, JAK,  $\beta$ -amyloid, Adenylyl cyclase, IKK $\beta$ , DNA polymerase. Resveratrol also is a specific SIRT1 activator<sup>[1][2][3][4]</sup>. Resveratrol is a potent pregnane X receptor (PXR) inhibitor<sup>[5]</sup>. Resveratrol is an Nrf2 activator, ameliorates aging-related progressive renal injury in mice model<sup>[6]</sup>. Resveratrol increases production of NO in endothelial cells<sup>[7]</sup>.

#### IC<sub>50</sub> & Target

Adenylyl cyclase 0.8 nM (IC<sub>50</sub>) IKKβ 1 μM (IC<sub>50</sub>) DNA polymerase  $\alpha$  3.3  $\mu$ M (IC<sub>50</sub>)

DNA polymerase  $\delta$ 

5 μM (IC<sub>50</sub>)

Autophagy

Mitophagy

Sirtuin

#### In Vitro

Resveratrol (trans-Resveratrol; SRT501) is one of the numerous polyphenolic compounds found in several vegetal sources In the vast majority of cases, Resveratrol displays inhibitory/activatory effects in the micromolar range, which is potentially attainable pharmacologically, although targets with affinities in the nanomolar range have also been reported  $^{[1]}$ . MCF-7 cells are plated in DME-F12 medium supplemented with 5% FBS in the presence of increasing concentrations of Resveratrol. Control cells are treated with the same volume of vehicle only (0.1% ethanol). Resveratrol inhibits the growth of MCF-7 cells in a dose-dependent fashion. Addition of 10  $\mu$ M Resveratrol results in an 82% inhibition of MCF-7 cell growth after 6 days while at 1  $\mu$ M, only a 10% inhibition is observed. The cells treated with 10  $\mu$ M Resveratrol have a doubling time of 60 hr whereas control cells doubled every 30 hr. Trypan blue exclusion assay shows that at concentrations of 10  $\mu$ M or lower, Resveratrol does not affect cell viability (90% viable cells) whereas at 100  $\mu$ M, only 50% of the cells are viable after 6 days of Resveratrol treatment. Moreover, MCF-7 cells do not undergo apoptosis after incubation with Resveratrol at concentration of 10  $\mu$ M as determined by ApoAlert Annexin V Apoptosis kit<sup>[2]</sup>.

Resveratrol increases the production of nitric oxide (NO) in endothelial cells by upregulating the expression of endothelial NO synthase (eNOS), stimulating eNOS enzymatic activity, and preventing eNOS uncoupling<sup>[7]</sup>.

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

#### In Vivo

The average tumor volume is reduced by treatment with Resveratrol (trans-Resveratrol; SRT501) at a dose of 50 mg/kg body weight (195.5 $\pm$ 124.8 mm<sup>3</sup>; P<0.05) or 100 mg/kg body weight (81.7 $\pm$ 70.5 mm<sup>3</sup>; P<0.001) compare with the vehicle-treated animals (315 $\pm$ 94 mm<sup>3</sup>). There is a good correlation between the tumor volume and the tumor mass<sup>[3]</sup>.

 ${\tt MCE}\ has\ not\ independently\ confirmed\ the\ accuracy\ of\ these\ methods.\ They\ are\ for\ reference\ only.$ 

## **PROTOCOL**

#### Cell Assay [2]

To determine the effect of Resveratrol on cell growth, MCF-7 cells are plated in 6-well plates at 10<sup>5</sup> cells per well in 2 mL of DME-F12 medium supplemented with 5% FBS in the presence or absence of increasing concentrations of Resveratrol. The cell number is measured every 2 days till day 6 with a hemocytometer after detaching the cells with trypsin-EDTA<sup>[2]</sup>.

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

# Animal Administration [3]

### Mice<sup>[3]</sup>

Female BALB/c (nu/nu) mice, 6 weeks old, are used. PA-1 cells ( $1 \times 10^7$  in 200  $\mu$ L PBS) are injected s.c. on the right hind flank. Tumor volume (length×width×depth×0.52) is measured three times a week. After 10 days of implantation, two groups (n=10)

are given Resveratrol (dissolved in 5% ethanol and 25% polyethyleneglycol 400 in distilled water) i.p. at a daily dose of 50 or 100 mg/kg body weight for consecutive 4 weeks, whereas the other group receive the vehicle only. Body weights are recorded everyday. Animals are given bromodeoxyuridine (BrdUrd; 10 mg/kg body weight, i.p.) 2 h before sacrifice. Xenograft tumors are weighed and frozen in liquid nitrogen or fixed in 10% formalin and embedded in paraffin. The BrdUrdlabeled cells in paraffin-embedded tissues are detected employing a monoclonal anti-BrdUrd antibody. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

#### **CUSTOMER VALIDATION**

- ACS Nano. 2023 Oct 2.
- Environ Int. 2023 Nov 25, 108354.
- Redox Biol. 2022 Jun;52:102310.
- Sci Total Environ. 2023 Sep 16;166954.
- PLoS Biol. 2022 Jun 30;20(6):e3001682.

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#### **REFERENCES**

- [1]. Pirola L, et al. Resveratrol: one molecule, many targets. IUBMB Life. 2008 May;60(5):323-32.
- [2]. Lu R, et al. Resveratrol, a natural product derived from grape, exhibits antiestrogenic activity and inhibits the growth of human breast cancer cells. J Cell Physiol. 1999 Jun;179(3):297-304.
- [3]. Lee MH, et al. Resveratrol suppresses growth of human ovarian cancer cells in culture and in a murine xenograft model: eukaryotic elongation factor 1A2 as a potential target. Cancer Res. 2009 Sep 15;69(18):7449-58.
- [4]. Du LL, et al. Activation of sirtuin 1 attenuates cerebral ventricular streptozotocin-induced tau hyperphosphorylation and cognitive injuries in rat hippocampi. Age (Dordr). 2014 Apr;36(2):613-23.
- [5]. Smutny T, et al. Resveratrol as an inhibitor of pregnane X receptor (PXR): another lesson in PXR antagonism. J Pharmacol Sci. 2014;126(2):177-8.
- [6]. Eun Nim Kim, et al. Resveratrol, an Nrf2 activator, ameliorates aging-related progressive renal injury. Aging (Albany NY). 2018 Jan; 10(1): 83–99.
- [7]. Huige Li, et al. Resveratrol and Vascular Function. Int J Mol Sci. 2019 Apr 30;20(9):2155.

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

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