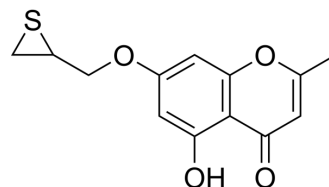


## HSP27 inhibitor J2

<b>Cat. No.:</b>	HY-124653		
<b>CAS No.:</b>	2133499-85-9		
<b>Molecular Formula:</b>	C <sub>13</sub> H <sub>12</sub> O <sub>4</sub> S		
<b>Molecular Weight:</b>	264.3		
<b>Target:</b>	HSP		
<b>Pathway:</b>	Cell Cycle/DNA Damage; Metabolic Enzyme/Protease		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 20 mg/mL (75.67 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	<b>Preparing Stock Solutions</b>	1 mM	3.7836 mL	18.9179 mL	37.8358 mL
		5 mM	0.7567 mL	3.7836 mL	7.5672 mL
10 mM		0.3784 mL	1.8918 mL	3.7836 mL	
Please refer to the solubility information to select the appropriate solvent.					
<b>In Vivo</b>	<ol style="list-style-type: none"> <li>Add each solvent one by one: 50% PEG300 &gt;&gt; 50% saline Solubility: 5 mg/mL (18.92 mM); Suspended solution; Need ultrasonic and warming and heat to 48°C</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2 mg/mL (7.57 mM); Clear solution</li> </ol>				

### BIOLOGICAL ACTIVITY

<b>Description</b>	HSP27 inhibitor J2 (J2) is a HSP27 inhibitor, which significantly induces abnormal HSP27 dimer formation and inhibits a production of HSP27 giant polymers, thereby having an effect of inhibiting a chaperone function of the HSP27 and reducing a cell protection function thereof. HSP27 inhibitor J2 (J2) remarkably enhances the antiproliferative activity of 17-AAG and sensitizes cisplatin-induced lung cancer cell growth inhibition <sup>[1][2]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	HSP27 <sup>[2]</sup>
<b>In Vitro</b>	HSP27 inhibitor J2 (10 μM; 12 h) induces significant abnormal HSP27 dimer formation in NCI-H460 <sup>[1]</sup> . HSP27 inhibitor J2 (10 μM; 24 h) increases 17-AAG-induced apoptosis in shCTRL cells <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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## CUSTOMER VALIDATION

- J Exp Clin Cancer Res. 2023 Apr 4;42(1):80.
- Int J Mol Sci. 2023 Jul 18, 24(14), 11598.
- Biomedicines. 2022, 10(10), 2489.
- University of Zagreb. Department of Biology. 2021 Sep.

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## REFERENCES

- [1]. Hwang SY, et al. Synthesis and biological effect of chrom-4-one derivatives as functional inhibitors of heat shockprotein 27. Eur J Med Chem. 2017 Oct 20;139:892-900.
- [2]. Younghwa Na, et al. Methods for treating pulmonary fibrosis using chromenone derivatives.
- 

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

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