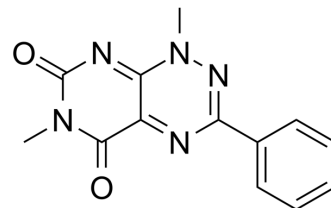


## 3-Phenyltoxoflavin

<b>Cat. No.:</b>	HY-125759		
<b>CAS No.:</b>	32502-63-9		
<b>Molecular Formula:</b>	C <sub>13</sub> H <sub>11</sub> N <sub>5</sub> O <sub>2</sub>		
<b>Molecular Weight:</b>	269.26		
<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	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 7.14 mg/mL (26.52 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
<b>1 mM</b>	3.7139 mL	18.5694 mL	37.1388 mL
<b>5 mM</b>	0.7428 mL	3.7139 mL	7.4278 mL
<b>10 mM</b>	0.3714 mL	1.8569 mL	3.7139 mL

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

### BIOLOGICAL ACTIVITY

**Description** 3-Phenyltoxoflavin, a derivative of Toxoflavin, is an Hsp90 inhibitor, with a K<sub>d</sub> of 585 nM for the interaction of Hsp90-TPR2A. 3-Phenyltoxoflavin has anti-cancer activity<sup>[1][2]</sup>.

**IC<sub>50</sub> & Target** HSP90  
585 nM (Kd)

**In Vitro** 3-Phenyltoxoflavin (1 nM-100 μM; 4 d) inhibits BT474 cells proliferation in a concentration-dependent manner, with an IC<sub>50</sub> of 690 nM<sup>[1]</sup>.  
3-Phenyltoxoflavin (0.56 nM-100 μM; 2 h) competes with biotinylated Hsp90 peptide for its binding to TPR2A in a dose-dependent manner<sup>[1]</sup>.  
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

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[1]. Yi F, et, al. A novel class of small molecule inhibitors of Hsp90. ACS Chem Biol. 2008 Oct 17;3(10):645-54.

[2]. Koh S, et, al. A novel light-dependent selection marker system in plants. Plant Biotechnol J. 2011 Apr;9(3):348-58.

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

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