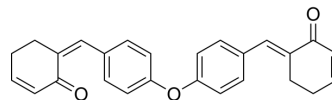


## TrxR-IN-5

Cat. No.:	HY-147803		
Molecular Formula:	C <sub>26</sub> H <sub>22</sub> O <sub>3</sub>		
Molecular Weight:	382.45		
Target:	Reactive Oxygen Species		
Pathway:	Immunology/Inflammation; Metabolic Enzyme/Protease; NF-κB		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 50 mg/mL (130.74 mM; ultrasonic and heat to 60°C)

Concentration	Mass			
	1 mg	5 mg	10 mg	
1 mM	2.6147 mL	13.0736 mL	26.1472 mL	
5 mM	0.5229 mL	2.6147 mL	5.2294 mL	
10 mM	0.2615 mL	1.3074 mL	2.6147 mL	

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

### BIOLOGICAL ACTIVITY

#### Description

TrxR-IN-5 (compound 4f) is a potent TrxR (thioredoxin reductase) inhibitor, with an IC<sub>50</sub> of 0.16 μM. TrxR-IN-5 increases the levels of ROS, thus leading to potent antiproliferative effects. TrxR-IN-5 exhibits prominent anticancer and anti-metastasis effects<sup>[1]</sup>.

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

IC<sub>50</sub>: 0.16 ± 0.02 μM (TrxR)<sup>[1]</sup>

#### In Vivo

TrxR-IN-5 (compound 4f) (MDA-MB-231 xenograft in BALB/c nude mice, 0-25 mg/kg, IP, once) shows strong inhibition potency toward solid tumors of breast cancer in vivo<sup>[1]</sup>.  
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Meng C, et al. Efficacy of novel methylenecyclohexenone derivatives as TrxR inhibitors in suppressing the proliferation and metastasis of human cancer cells. Bioorg Chem. 2020 Dec;105:104360.

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

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