# **Screening Libraries**

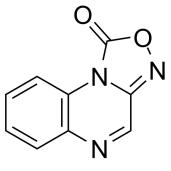
# **ODQ**

Cat. No.: HY-101255 CAS No.: 41443-28-1 Molecular Formula:  $C_9H_5N_3O_2$ Molecular Weight: 187.15

Target: Guanylate Cyclase; Apoptosis Pathway: GPCR/G Protein; Apoptosis Storage: Powder -20°C 3 years

> 4°C 2 years In solvent -80°C 2 years

> > -20°C 1 year



**Product** Data Sheet

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (534.33 mM; Need ultrasonic)

| Preparing<br>Stock Solutions | Solvent Mass<br>Concentration | 1 mg      | 5 mg       | 10 mg      |
|------------------------------|-------------------------------|-----------|------------|------------|
|                              | 1 mM                          | 5.3433 mL | 26.7165 mL | 53.4331 mL |
|                              | 5 mM                          | 1.0687 mL | 5.3433 mL  | 10.6866 mL |
|                              | 10 mM                         | 0.5343 mL | 2.6717 mL  | 5.3433 mL  |

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

In Vivo

1. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 5 mg/mL (26.72 mM); Clear solution

# **BIOLOGICAL ACTIVITY**

| Description | ODQ is a potent and selective soluble guanylyl cyclase (sGC, nitric oxide-activated enzyme) inhibitor. ODQ enhances the proapoptotic effects of Cisplatin in human mesothelioma cells <sup>[1]</sup> .  |
|-------------|---|
| In Vitro    | At 30 and 50 $\mu$ M, ODQ causes significant induction of apoptosis in the NCI-H2452 cells, elevating apoptotic levels by 12 fold and 15 fold, respectively. At 10 $\mu$ M, a concentration below the threshold for induction of apoptosis by ODQ, ODQ in combination with Cisplatin enhanced (in fact, doubled) the pro-apoptotic effects of Cisplatin at 1 $\mu$ M <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |
| In Vivo     | ODQ (2 mg/kg; i.p.) reduces the multiple organ injury and dysfunction caused by wall fragments of Gram-positive or Gram-negative bacteria in the anesthetized rat <sup>[2]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.   |

| Animal Model:   | Anesthetized, male Wistar rats <sup>[2]</sup>   |  |
|-----------------|---|--|
| Dosage:         | 2 mg/kg   |  |
| Administration: | l.p.  |  |
| Result:         | Attenuated the renal dysfunction, lung injury, and hepatocellular injury caused by lipoteichoic acid/peptidoglycan or lipopolysaccharide. |  |

### **REFERENCES**

[1]. Ronald Fiscus, et al. ODQ, an inhibitor of soluble guanylyl cyclase (nitric oxide-activated enzyme), enhances the pro-apoptotic effects of cisplatin in human mesothelioma cells. Cancer Res May 1 2007 (67) (9 Supplement) LB-43.

[2]. Zacharowski K, et al. The selective guanylate cyclase inhibitor ODQ reduces multiple organ injury in rodent models of Gram-positive and Gram-negative shock. Crit Care Med. 2001;29(8):1599-1608.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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