# Z57346765

Cat. No.: HY-W195984 CAS No.: 1016340-64-9 Molecular Formula:  $C_{17}H_{18}N_4O$ Molecular Weight: 294.35 Target: Others Pathway: Others

Storage: Powder

-20°C 3 years 2 years

In solvent -80°C 6 months

> -20°C 1 month

**Product** Data Sheet

### **SOLVENT & SOLUBILITY**

In Vitro

DMSO : ≥ 100 mg/mL (339.73 mM)

\* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.3973 mL	16.9866 mL	33.9732 mL
	5 mM	0.6795 mL	3.3973 mL	6.7946 mL
	10 mM	0.3397 mL	1.6987 mL	3.3973 mL

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (8.49 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- $\beta$ -CD in saline) Solubility: ≥ 2.5 mg/mL (8.49 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	Z57346765 is a PGK1-specific inhibitor that reduces the activity of metabolic enzymes in PGK1 glycolysis and inhibits PGK1-dependent cell proliferation. Z57346765 exerts inhibitory effects against clear cell renal cell carcinoma (KIRC) <sup>[1]</sup> .
IC <sub>50</sub> & Target	$PGK1^{[1]}$

## **REFERENCES**

1]. He Y, et al. Novel inhibitors targeting the PGK1 metabolic enzyme in glycolysis exhibit effective antitumor activity against kidney renal clear cell carcinoma in vitro and n vivo. Eur J Med Chem. 2024 Mar 5;267:116209.						
	Caution: Product has no	ot been fully validated for me	dical applications. For research use onl	y.		
	Tel: 609-228-6898	Fax: 609-228-5909 Deer Park Dr, Suite Q, Monmo	E-mail: tech@MedChemExpress.com	m		
	Address. I	beer rank bi, baite Q, monnie	attrounction, 143 00032, 03/1			

Page 2 of 2 www.MedChemExpress.com