Proteins

Inhibitors

MYF-01-37

Cat. No.: HY-139603 CAS No.: 2416417-65-5 Molecular Formula: $C_{15}H_{17}F_{3}N_{2}O$ Molecular Weight: 298.3

YAP Target:

Pathway: Stem Cell/Wnt

Storage: Powder -20°C 3 years

2 years

-80°C In solvent 6 months

> -20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 250 mg/mL (838.08 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.3523 mL	16.7617 mL	33.5233 mL
	5 mM	0.6705 mL	3.3523 mL	6.7047 mL
	10 mM	0.3352 mL	1.6762 mL	3.3523 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: ≥ 2.08 mg/mL (6.97 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (6.97 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

MYF-01-37 is a covalent TEAD inhibitor targeting Cys380. MYF-01-37 has a reversible inhibition on YAP/TEAD interaction^[1].

In Vitro

MYF-01-37 (10 μ M; 24h) results in inhibition of direct YAP/TEAD interaction in HEK 293T cells, and in the reduction in canonical YAP target gene CTGF expression in PC-9 cells^[1].

MYF-01-37 (0.1, 1, 10, 100 μ M) has minimal impact on cell viability of several EGFR-mutant NCSLC cell lines [1].

MYF-01-37 (10 μ M; 10 days) combinated with OT (combination of osimertinib and trametinib) leads to a dramatic decrease in dormant cells compared to OT alone^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES	
l]. Kari J Kurppa, et al. Treatme 3;37(1):104-122.e12.	nt-Induced Tumor Dormancy through YAP-Mediated Transcriptional Reprogramming of the Apoptotic Pathway. Cancer Cell. 2020 Jan
	Caution: Product has not been fully validated for medical applications. For research use only.
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