

CFI-402257

## Chemical Properties

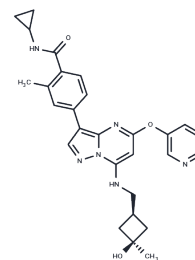
CAS No. : 1610759-22-2

Formula: C<sub>28</sub>H<sub>30</sub>N<sub>6</sub>O<sub>3</sub>

Molecular Weight: 498.58

Storage: Store at low temperature, Keep away from moisture  
Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Actual storage temperature shall be subject to the COA.



## Biological Description

Description	CFI-402257 (Luvixasertib) is a highly selective oral TTK (Mps1, Monopolar Spindle 1) kinase inhibitor that potently inhibits TTK kinase activity (Mps1 Ki = 0.09 nM; EC <sub>50</sub> = 6.5 nM). It is intended for use in research on hepatocellular carcinoma and breast cancer.
Targets(IC50)	Kinesin, Tyrosine Kinases
In vitro	<b>Methods:</b> MHCC97L cells were treated with 100 nM CFI-402257 for 72 hours, followed by flow cytometric analysis of DNA content. <b>Results:</b> CFI-402257 induced a significant increase in the proportion of aneuploid cells. [1]
In vivo	<b>Methods:</b> MHCC97L-luc cells were implanted in BALB/cAnN-nu mice. Treatment commenced 2 weeks post-implantation when tumor formation initiated. CFI-402257 was administered orally via gavage at 30 mg/kg once daily for 4 consecutive weeks. <b>Results:</b> Tumor volume significantly decreased, and the number of pulmonary metastatic nodules markedly reduced. [1] <b>Methods:</b> A carotid artery wire injury model was established in C57BL/6 mice via carotid artery wire injury. Treatment commenced postoperatively with oral gavage of CFI-402257 at 30 mg/kg once daily for 28 consecutive days. <b>Results:</b> The area of newly formed endothelium and the intima-media ratio were significantly reduced, while re-endothelialization remained unaffected (confirmed by CD31 staining), demonstrating superiority over conventional drug-eluting stents. [2]

## Solubility Information

Solubility	DMSO: 37.7 mg/mL (75.61 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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### Preparing Stock Solutions

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	1mg	5mg	10mg
1 mM	2.0057 mL	10.0285 mL	20.057 mL
5 mM	0.4011 mL	2.0057 mL	4.0114 mL
10 mM	0.2006 mL	1.0028 mL	2.0057 mL
50 mM	0.0401 mL	0.2006 mL	0.4011 mL

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Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

### Reference

Chan CY, et al. CFI-402257, a TTK inhibitor, effectively suppresses hepatocellular carcinoma. Proc Natl Acad Sci U S A. 2022 Aug 9;119(32):e2119514119.

Wu JH, et al. TTK Inhibition Alleviates Postinjury Neointimal Formation and Atherosclerosis. Adv Sci (Weinh). 2025 Feb;12(6):e2409250.

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