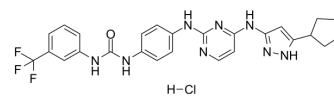


CD532 hydrochloride

Cat. No.:	HY-112273A
CAS No.:	2926498-81-7
Molecular Formula:	C ₂₆ H ₂₆ ClF ₃ N ₈ O
Molecular Weight:	558.99
Target:	Aurora Kinase
Pathway:	Cell Cycle/DNA Damage; Epigenetics
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (178.89 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg
		Concentration				
		1 mM		1.7889 mL	8.9447 mL	17.8894 mL
		5 mM		0.3578 mL	1.7889 mL	3.5779 mL
10 mM		0.1789 mL	0.8945 mL	1.7889 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 (4.47 mM); Suspended solution; Need ultrasonic					

BIOLOGICAL ACTIVITY

Description	CD532 hydrochloride is a potent Aurora A kinase inhibitor with an IC ₅₀ of 45 nM. CD532 hydrochloride has the dual effect of blocking Aurora A kinase activity and driving degradation of MYCN. CD532 hydrochloride also can directly interact with AURKA and induces a global conformational shift. CD532 hydrochloride can be used for the research of cancer ^{[1][2]} .
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REFERENCES

- [1]. Gustafson WC, et, al. Drugging MYCN through an allosteric transition in Aurora kinase A. *Cancer Cell*. 2014 Sep 8;26(3):414-427.
- [2]. Lee JK, et, al. N-Myc Drives Neuroendocrine Prostate Cancer Initiated from Human Prostate Epithelial Cells. *Cancer Cell*. 2016 Apr 11;29(4):536-547.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA