YLF-466D

Cat. No.:	HY-15840		
CAS No.:	1273323-67	-3	
Molecular Formula:	C ²⁹ H ²⁰ CINO	3	
Molecular Weight:	465.93		
Target:	AMPK		
Pathway:	Epigenetics	; PI3K/Al	t/mTOR
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year

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SOLVENT & SOLUBILITY

In Vitro	DMSO : ≥ 100 mg/mL (214.62 mM) * "≥" means soluble, but saturation unknown.					
		Solvent Mass Concentration	1 mg	5 mg	10 mg	
	Preparing Stock Solutions	1 mM	2.1462 mL	10.7312 mL	21.4625 mL	
		5 mM	0.4292 mL	2.1462 mL	4.2925 mL	
		10 mM	0.2146 mL	1.0731 mL	2.1462 mL	
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent o Solubility: ≥ 2.5 mg	one by one: 10% DMSO >> 40% PE(g/mL (5.37 mM); Clear solution	G300 >> 5% Tween-8) >> 45% saline		
	2. Add each solvent o Solubility: ≥ 2.5 mg	one by one: 10% DMSO >> 90% cor g/mL (5.37 mM); Clear solution	n oil			

BIOLOGICALACTIVITY				
Description	YLF-466D is a newly developed AMPK activator, which inhibits platelet aggregation.			
IC ₅₀ & Target	АМРК			
In Vitro	The effect of YLF-466D on platelet AMPK and aggregation are examined to test whether YLF-466D can stimulate AMPK in platelets and thereby suppress aggregation. Platelet AMPK is activated by YLF-466D, which is confirmed with activation-dependent phosphorylation at Thr172. Consistent with this result, YLF-466D inhibits platelet aggregation induced by thrombin. Such inhibition is observed in the aggregation elicited by ADP and collagen as well as thrombin, indicating that the antiaggregatory effect of YLF-466D is not platelet-agonist specific but common, regardless of agonist type. All the effects			

Product Data Sheet

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on AMPK and aggregation are concentration-dependent with the highest efficacy at 150 μ M. IC₅₀ against thrombin-, ADPand collagen-induced aggreation are approximately 84, 55 and 87 μ M, respectively^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

PROTOCOL)
Kinase Assay ^[1]	Blood is collected from the abdominal aorta of ether anesthetized rats using 3.2% sodium citrate as an anticoagulant (sodium citrate:blood=1:9) and diluted with normal saline (1:1). Blood is incubated with YLF-466D (0, 50, 100 and 150 μM) for 3 min and aggregation is induced with 7.5 μg/mL Collagen. Aggregation is assessed by measuring the impedance change with a whole blood aggregometer ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Eur J Pharmacol. 2015 Aug 5;760:81-7.
- PLoS One. 2020 Oct 14;15(10):e0240517.

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REFERENCES

[1]. Liu Y, et al. Antiplatelet effect of a newly developed AMP-activated protein kinase activator YLF-466D. Eur J Pharmacol. 2015 Aug 5;760:81-7.

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

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