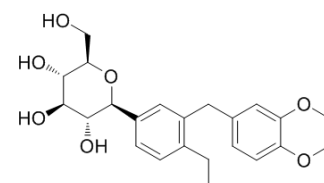


## Licogliflozin

<b>Cat. No.:</b>	HY-109092		
<b>CAS No.:</b>	1291094-73-9		
<b>Molecular Formula:</b>	C <sub>23</sub> H <sub>28</sub> O <sub>7</sub>		
<b>Molecular Weight:</b>	416.46		
<b>Target:</b>	SGLT		
<b>Pathway:</b>	Membrane Transporter/Ion Channel		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 100 mg/mL (240.12 mM; Need ultrasonic)  
 H<sub>2</sub>O : 2 mg/mL (4.80 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
	1 mM	2.4012 mL	12.0060 mL	24.0119 mL
	5 mM	0.4802 mL	2.4012 mL	4.8024 mL
	10 mM	0.2401 mL	1.2006 mL	2.4012 mL

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

#### In Vivo

- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)  
 Solubility: ≥ 2.5 mg/mL (6.00 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil  
 Solubility: ≥ 2.5 mg/mL (6.00 mM); Clear solution

### BIOLOGICAL ACTIVITY

#### Description

Licogliflozin is a sodium glucose cotransporter (SGLT1 and SGLT2) inhibitor.

#### IC<sub>50</sub> & Target

SGLT1, SGLT2<sup>[1]</sup>

#### In Vitro

Licogliflozin (LIK066) is a non-anti-fibrotic treatment agent for non-alcoholic steatohepatitis (NASH).  
 MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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## REFERENCES

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[1]. Monica A. Konerman, et al. Overview of Clinical Treatment Trials for NASH. *Curr Hepatol Rep.* 2017 Dec;16(4): 366-373.

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

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