# **Product** Data Sheet

#### CK-869

 $\begin{array}{lll} \textbf{Cat. No.:} & \textbf{HY-16927} \\ \textbf{CAS No.:} & 388592\text{-}44\text{-}7 \\ \textbf{Molecular Formula:} & \textbf{C}_{17}\textbf{H}_{16}\textbf{BrNO}_{3}\textbf{S} \\ \end{array}$ 

Molecular Weight: 394.28

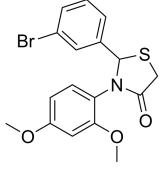
Target: Arp2/3 Complex
Pathway: Cytoskeleton

Storage: Powder -20°C

-20°C 3 years 4°C 2 years

In solvent -80°C 2 years

-20°C 1 year



#### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 25 mg/mL (63.41 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.5363 mL	12.6813 mL	25.3627 mL
	5 mM	0.5073 mL	2.5363 mL	5.0725 mL
	10 mM	0.2536 mL	1.2681 mL	2.5363 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:  $\geq$  2.5 mg/mL (6.34 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- $\beta$ -CD in saline) Solubility:  $\geq$  2.5 mg/mL (6.34 mM); Suspended solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.34 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	CK-869 is an Actin-Related Protein 2/3 (ARP2/3) complex inhibitor, with an IC $_{50}$ of 7 $\mu$ M.	
IC <sub>50</sub> & Target	IC50: 7 μM (ARP2/3) <sup>[1]</sup> .	
In Vitro	CK-869 is an Actin-Related Protein 2/3 (ARP2/3) complex inhibitor, with an IC $_{50}$ of 7 $\mu$ M $^{[1]}$ . CK-869 significantly inhibits MT polymerization even at a concentration of 25 $\mu$ M $^{[2]}$ . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

## **CUSTOMER VALIDATION**

• Cell Rep. 2021 Jul 6;36(1):109318.

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#### **REFERENCES**

[1]. Hetrick B, et al. Small molecules CK-666 and CK-869 inhibit actin-related protein 2/3 complex by blocking an activating conformational change. Chem Biol. 2013 May 23;20(5):701-12.

[2]. Yamagishi Y, et al. Use of CK-548 and CK-869 as Arp2/3 complex inhibitors directly suppresses microtubule assembly both in vitro and in vivo. Biochem Biophys Res Commun. 2018 Feb 12;496(3):834-839.

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

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