

## **Product** Data Sheet

## sAJM589

Cat. No.:HY-122683CAS No.:2089-82-9Molecular Formula: $C_{16}H_{10}N_2O$ Molecular Weight:246.26Target:c-MycPathway:Apoptosis

Storage: Powder -20°C

20°C 3 years 4°C 2 years

In solvent -80°C 6 months

-20°C 1 month

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 5 mg/mL (20.30 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	4.0607 mL	20.3037 mL	40.6075 mL
	5 mM	0.8121 mL	4.0607 mL	8.1215 mL
	10 mM	0.4061 mL	2.0304 mL	4.0607 mL

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

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Description	sAJM589 is a Myc inhibitor which potently disrupts the Myc-Max heterodimer with an IC $_{50}$ of 1.8 $\mu$ M $^{[1]}$ .
IC <sub>50</sub> & Target	IC50: 1.8 μM (Myc) <sup>[1]</sup>
In Vitro	sAJM589 potently disrupts the Myc-Max heterodimer to reduce Myc protein levels in a dose dependent manner, with an IC <sub>50</sub> of 1.8 μM <sup>[1]</sup> .  ?sAJM589 suppresses cellular proliferation in diverse Myc-dependent cancer cell lines and anchorage independent growth of Raji cells <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **CUSTOMER VALIDATION**

• Biochim Biophys Acta Gen Subj. 2022 Jan 20;130093.

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REFERENCES	
[1]. Choi SH, et al. Targeted Disruption of Myc-Max Oncoprotein Complex by a Small Molecule. ACS Chem Biol. 2017 Nov 17;12(11):2715-2719.	
Caution: Product has not been fully validated for medical applications. For research use only.	
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