**Proteins** 

# **Y16**

Cat. No.: HY-12649 CAS No.: 429653-73-6 Molecular Formula:  $C_{24}H_{20}N_{2}O_{3}$ 

Molecular Weight: 384 Target: Ras

Pathway: GPCR/G Protein; MAPK/ERK Pathway

-20°C Storage: Powder

3 years 4°C 2 years

-80°C In solvent 2 years

> -20°C 1 year

**Product** Data Sheet

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 25 mg/mL (65.10 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.6042 mL	13.0208 mL	26.0417 mL
	5 mM	0.5208 mL	2.6042 mL	5.2083 mL
	10 mM	0.2604 mL	1.3021 mL	2.6042 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 (6.51 mM); Suspended solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.51 mM); Clear solution

### **BIOLOGICAL ACTIVITY**

Description Y16 is a specific inhibitor of Leukemia-associated Rho guanine nucleotide exchange factor (LARG) with a K<sub>d</sub> value of 76 nM. Y16 is active in blocking the interaction of LARG and related G-protein-coupled Rho GEFs with RhoA. Y16 shows no detectable effect on other diffuse B-cell lymphoma (Dbl) family Rho GEFs, Rho effectors, or a RhoGAP<sup>[1]</sup>.

Kd: 76 nM (LARG)[1] IC<sub>50</sub> & Target

Y16 (10-30 μM; 24 hours; NIH 3T3 cells) could inhibit RhoA-GTP formation induced by serum dose dependently and is specific In Vitro for  $RhoA^{[1]}$ .

> Y16 (10-30 μM; 24 hours; NIH 3T3 cells) efficiently inhibits serum or SDF-1α-induced phospho-MLC and phospho-FAK formation, which are downstream of RhoA<sup>[1]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability  $Assay^{[1]}$ Cell Line: NIH 3T3 cells Concentration:  $10~\mu\text{M}, 30~\mu\text{M}$ **Incubation Time:** 24 hours Result: Inhibited RhoA-GTP formation induced by serum dose dependently and was specific for RhoA. Western Blot Analysis<sup>[1]</sup> Cell Line: NIH 3T3 cells Concentration:  $10 \, \mu M$ ,  $30 \, \mu M$ **Incubation Time:** 24 hours Result: Inhibited serum or SDF- $1\alpha$ -induced phospho-MLC and phospho-FAK formation, which

### **CUSTOMER VALIDATION**

- J Cell Mol Med. 2020 Jul;24(14):8179-8193.
- Front Endocrinol. 2021 Feb 4;11:621944.
- Oncol Lett. 2022 Jun;23(6):173.
- J Breast Cancer. 2019 Apr 22;22(2):185-195.

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#### **REFERENCES**

[1]. Shang X, et al. Small-molecule inhibitors targeting G-protein-coupled Rho guanine nucleotide exchange factors. Proc Natl Acad Sci U S A. 2013 Feb 19;110(8):3155-60.

were downstream of RhoA.

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

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