## **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		
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 : 25 mg/mL (65	DMSO : 25 mg/mL (65.10 mM; Need ultrasonic)				
F	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	<ol> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 40% PEG300 &gt;&gt; 5% Tween-80 &gt;&gt; 45% saline Solubility: 2.5 mg/mL (6.51 mM); Suspended solution; Need ultrasonic</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2.5 mg/mL (6.51 mM); Clear solution</li> </ol>					

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> .		
IC <sub>50</sub> & Target	Kd: 76 nM (LARG) <sup>[1]</sup>		
In Vitro	Y16 (10-30 μM; 24 hours; NIH 3T3 cells) could inhibit RhoA-GTP formation induced by serum dose dependently and is specific for RhoA <sup>[1]</sup> . 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> .		

# Product Data Sheet

MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay<sup>[1]</sup>

10 μΜ, 30 μΜ
24 hours
Inhibited RhoA-GTP formation induced by serum dose dependently and was specific for RhoA.

Cell Line:	NIH 3T3 cells
Concentration:	10 μΜ, 30 μΜ
Incubation Time:	24 hours
Result:	Inhibited serum or SDF-1 $\alpha$ -induced phospho-MLC and phospho-FAK formation, which were downstream of RhoA.

### **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.

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

Tel: 609-228-6898Fax: 609-228-5909E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA