# BAY885

®

MedChemExpress

Cat. No.:	HY-112082	
CAS No.:	2307249-33-6	
Molecular Formula:	$C_{25}H_{28}F_{3}N_{7}O_{2}$	
Molecular Weight:	515.53	
Target:	ERK	
Pathway:	MAPK/ERK Pathway; Stem Cell/Wnt	Í
Storage:	-20°C, stored under nitrogen	Ņ,
	* In solvent : -80°C, 6 months; -20°C, 1 month (stored under nitrogen)	Ň

# SOLVENT & SOLUBILITY

In Vitro	DMSO : 16.67 mg/mL (32.34 mM; Need ultrasonic)						
		Mass Solvent Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	1.9398 mL	9.6988 mL	19.3975 mL		
		5 mM	0.3880 mL	1.9398 mL	3.8795 mL		
		10 mM	0.1940 mL	0.9699 mL	1.9398 mL		
	Please refer to the so	lubility information to select the app	propriate solvent.				
In Vivo	1. Add each solvent one by one: 5% DMSO >> 40% PEG300 >> 5% Tween-80 >> 50% saline Solubility: ≥ 2.5 mg/mL (4.85 mM); Clear solution						
	2. Add each solvent one by one: 5% DMSO >> 95% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (4.85 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 1.67 mg/mL (3.24 mM); Clear solution						
	4. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 1.67 mg/mL (3.24 mM); Clear solution						
	5. Add each solvent o Solubility: 0.5 mg/	one by one: 1% DMSO >> 99% salin /mL (0.97 mM); Suspended solution;	e Need ultrasonic				

BIOLOGICAL ACTIVITY			
Description	BAY885 is a highly potent and selective ERK5 inhibitor with an IC <sub>50</sub> of 35 nM. BAY885 shows weak inhibition on others kinases <sup>[1]</sup> .		
IC <sub>50</sub> & Target	ERK5		

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 $H_2N$ 

F↓F

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	35 nM (IC <sub>50</sub> )
In Vitro	The ERK5 probe 41 (BAY-885) showed potent ERK5 kinase and transcriptional inhibition in the SN12C-MEF2 reporter cell line (IC <sub>50</sub> = 115 nM/IC <sub>90</sub> = 691 nM) and had no effects on a reporter control cell line with constitutive luciferase expression (SN12C-CMV-luc, IC <sub>50</sub> > 30 μM) <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## REFERENCES

[1]. Nguyen D, et al. Discovery and Characterization of the Potent and Highly Selective (Piperidin-4-yl)pyrido[3,2-d]pyrimidine Based in Vitro Probe BAY-885 for the Kinase ERK5. J Med Chem. 2019 Jan 24;62(2):928-940.

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

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