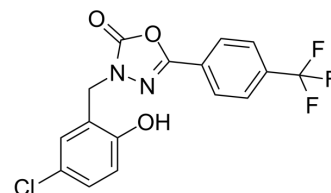


BMS-191011

Cat. No.:	HY-108593		
CAS No.:	202821-81-6		
Molecular Formula:	C ₁₆ H ₁₀ ClF ₃ N ₂ O ₃		
Molecular Weight:	370.71		
Target:	Potassium Channel		
Pathway:	Membrane Transporter/Ion Channel		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (269.75 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
	Preparing Stock Solutions	1 mM	2.6975 mL	13.4876 mL
	5 mM	0.5395 mL	2.6975 mL	
	10 mM	0.2698 mL	1.3488 mL	
	Please refer to the solubility information to select the appropriate solvent.			
In Vivo	<p>1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (5.61 mM); Clear solution</p> <p>2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (5.61 mM); Clear solution</p>			

BIOLOGICAL ACTIVITY

Description	BMS 191011 (BMS-A) is a potent BK _{Ca} channel opener (large-conductance Ca ²⁺ -activated potassium channel). BMS-191011 shows neuroprotective activities in rodent models of stroke ^[1] .	
In Vitro	BMS-191011 (20 or 40 μM) activates BK channels in IGR39 cells, and leads to potentiation in Panc-1 cells ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay ^[3]	
	Cell Line:	IGR39 and Panc-1 cells

	Concentration:	20 or 40 μ M
	Incubation Time:	
	Result:	Activated BK channels in less than 50% IGR39 cells and led to less than two-fold potentiation in Panc-1 cells. Promoted the activation of a voltage-independent K ⁺ conductance at negative voltages.
In Vivo	BMS-191011 (intravenous injection; 10-100 μ g/kg; once) dilates rat retinal arterioles through activation of iberiotoxin-sensitive BK(Ca) channels in vivo ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	Male Wistar rats (8- to 10-week-old) treated with Tetrodotoxin (50 μ g/kg, intravenously (i.v.)) ^[2]
	Dosage:	10-100 μ g/kg
	Administration:	Intravenous injection; 10-100 μ g/kg; once
	Result:	Increased the diameter of retinal arterioles, showed no significant effect on mean arterial pressure and heart rate.

CUSTOMER VALIDATION

- Br J Pharmacol. 2021 Jul 3.
- Eur J Pharmacol. 2023 Apr 19;175743.

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REFERENCES

- [1]. Asami Mori, et al. BMS-191011, an opener of large-conductance Ca²⁺-activated potassium channels, dilates rat retinal arterioles in vivo. Biol Pharm Bull. 2011;34(1):150-2.
- [2]. Remigante A, et al. NS-11021 Modulates Cancer-Associated Processes Independently of BK Channels in Melanoma and Pancreatic Duct Adenocarcinoma Cell Lines. Cancers (Basel). 2021 Dec 6;13(23):6144.
- [3]. Romine JL, et al. 3-[[5-Chloro-2-hydroxyphenyl)methyl]-5-[4-(trifluoromethyl)phenyl]-1,3,4-oxadiazol-2(3H)-one, BMS-191011: opener of large-conductance Ca(2+)-activated potassium (maxi-K) channels, identification, solubility, and SAR. J Med Chem. 2007 Fe

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

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