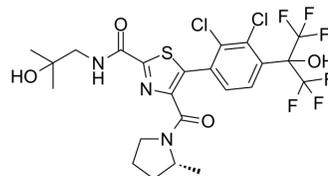


## JNJ-61803534

<b>Cat. No.:</b>	HY-139780		
<b>CAS No.:</b>	1917306-14-9		
<b>Molecular Formula:</b>	C <sub>23</sub> H <sub>23</sub> Cl <sub>2</sub> F <sub>6</sub> N <sub>3</sub> O <sub>4</sub> S		
<b>Molecular Weight:</b>	622.41		
<b>Target:</b>	ROR		
<b>Pathway:</b>	Metabolic Enzyme/Protease; Vitamin D Related/Nuclear Receptor		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 300 mg/mL (482.00 mM; Need ultrasonic)

Solvent	Mass	Concentration		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	1.6067 mL	8.0333 mL	16.0666 mL
	5 mM	0.3213 mL	1.6067 mL	3.2133 mL
	10 mM	0.1607 mL	0.8033 mL	1.6067 mL

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

### BIOLOGICAL ACTIVITY

#### Description

JNJ-61803534 is a potent and orally active ROR $\gamma$ t inverse agonist with an IC<sub>50</sub> of 9.6 nM. JNJ-61803534 has anti-inflammatory activity. JNJ-61803534 inhibits IL-17A production in human CD4+ T cells under Th17 differentiation conditions [1].

#### IC<sub>50</sub> & Target

ROR $\gamma$ t

#### In Vitro

JNJ-61803534 inhibits ROR $\gamma$ t transcription in HEK-293 T cells transfected with vectors encoding ROR $\gamma$ t, with an IC<sub>50</sub> of 9.6 nM<sup>[1]</sup>.  
 JNJ-61803534 (1 nM-1  $\mu$ M) inhibits IL-17A, IL-17F IFN $\gamma$  and IL-22 production in CD4+ T cells isolated from human blood<sup>[1]</sup>.  
 JNJ-61803534 does not impact in vitro Treg differentiation in CD4+ T cells<sup>[1]</sup>.  
 MCE has not independently confirmed the accuracy of these methods. They are for reference only.

#### In Vivo

JNJ-61803534 (100 mg/kg, p.o.) inhibits ex vivo stimulated IL-17A production in the blood of mice<sup>[1]</sup>.  
 JNJ-61803534 (3-100 mg/kg BID or 60 mg/kg QD, p.o.) alleviates inflammation, cartilage damage, bone destruction in mouse collagen-induced arthritis (CIA) model<sup>[1]</sup>.

JNJ-61803534 (30 and 100 mg/kg, p.o.) alleviates Imiquimod (HY-B0180)-induced dermal psoriatic-like inflammation in mice [1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Mouse collagen-induced arthritis (CIA) model <sup>[1]</sup>
Dosage:	3-100 mg/kg BID or 60 mg/kg QD
Administration:	Oral administration (p.o.)
Result:	Decreased clinical arthritis scores and hind paw histopathology scores.

## REFERENCES

[1]. Xue X, et al. Preclinical and clinical characterization of the ROR $\gamma$ t inhibitor JNJ-61803534. Sci Rep. 2021 May 26;11(1):11066.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

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