## Epristeride

Cat. No.:	HY-107385				
CAS No.:	119169-78-7				
Molecular Formula:	C <sub>25</sub> H <sub>37</sub> NO <sub>3</sub>				
Molecular Weight:	399.57				
Target:	5 alpha Reductase				
Pathway:	Metabolic Enzyme/Protease				
Storage:	Powder	-20°C	3 years		
		4°C	2 years		
	In solvent	-80°C	6 months		
		-20°C	1 month		

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## SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (250.27 mM; Need ultrasonic)							
		Solvent Mass Concentration	1 mg	5 mg	10 mg			
	Preparing Stock Solutions	1 mM	2.5027 mL	12.5135 mL	25.0269 mL			
		5 mM	0.5005 mL	2.5027 mL	5.0054 mL			
		10 mM	0.2503 mL	1.2513 mL	2.5027 mL			
	Please refer to the so	lubility information to select the app	propriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 2.5 mg/mL (6.26 mM); Suspended solution; Need ultrasonic							
		2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.26 mM); Clear solution						
	ent one by one: 10% DMSO >> 90% corn oil 5 mg/mL (6.26 mM); Clear solution							

BIOLOGICAL ACTIVITY				
Description	Epristeride (ONO-9302) is a selective, specific and orally active uncompetitive inhibitor of human steroid 5 alpha-reductase isoform 2. Epristeride has inhibitory effects for SR isoenzymes types 2 with K <sub>i</sub> value of 0.7-2 nM. Epristeride can be used for the research of prostatic hyperplasia and acne <sup>[1]</sup> .			
IC <sub>50</sub> & Target	Ki: 0.7-2 nM(SR2) <sup>[1]</sup>			
In Vitro	Epristeride has inhibitory effects for SR isoenzymes types 2 (SR2) with $K_i$ values of 0.7-2 nM <sup>[1]</sup> .			

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

## REFERENCES

[1]. M A Levy, et al. Epristeride is a selective and specific uncompetitive inhibitor of human steroid 5 alpha-reductase isoform 2. J Steroid Biochem Mol Biol. 1994 Feb;48(2-3):197-206.

## 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