Masofaniten

Cat. No.:	HY-136582		
CAS No.:	2416716-62-	-4	
Molecular Formula:	C ₂₄ H ₂₄ Cl ₂ N ₄ O ₄ S		
Molecular Weight:	535.44		
Target:	Androgen R	eceptor	
Pathway:	Vitamin D R	elated/Nu	Iclear Receptor
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month

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SOLVENT & SOLUBILITY

	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	1.8676 mL	9.3381 mL	18.6762 ml
	5 mM	0.3735 mL	1.8676 mL	3.7352 mL
	10 mM	0.1868 mL	0.9338 mL	1.8676 mL

BIOLOGICAL ACTIV	
Description	Masofaniten (Androgen receptor-IN-2) is a potent and orally active androgen receptor inhibitor. Masofaniten has antitumor activity against prostate cancer ^[1] .
IC ₅₀ & Target	Androgen receptor ^[1]
In Vitro	Masofaniten (compound A109, androgen-induced PSA-Luciferase assay) inhibits androgen binding to androgen receptor with an IC ₅₀ of 535 nM ^[1] . Masofaniten inhibits cell proliferation in LNCaP and LNCaP95 cells (IC ₅₀ : 0.44 μM, 3.78 μM respectively) ^[1] . Masofaniten shows a metabolic stability in liver microsome with a t _{1/2} of more than 120 min, and in hepatocytes with a t _{1/2} of more than 360 min ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	Masofaniten (compound A109) (60 mg/kg, p.o.) induces partial regressions of tumor growth in NCG mice bearing LNCaP tumors ^[1] . Masofaniten (5 mg/kg, p.o., single dose, in male CD-1 mice) shows a t _{1/2} of 8.1 h, C _{max} of 2673.3 ng/mL, F (%) of 33.6 ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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Product Data Sheet

Animal Model:	LNCaP Xenografts Model ^[1]
Dosage:	60 mg/kg
Administration:	Oral administration (p.o.)
Result:	Inhibited tumor growth no obvious drug related toxicity (bodyweight change).

REFERENCES

[1]. Han-Jie Zhou, et al. Androgen receptor modulators and methods for their use. Patent. US20200123117A1.

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

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