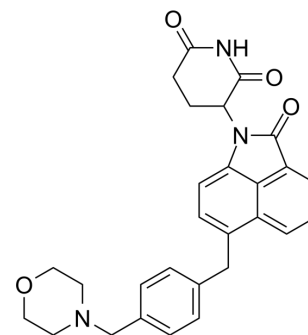


(Rac)-Cemsidomide

Cat. No.:	HY-144841A		
CAS No.:	2504233-68-3		
Molecular Formula:	C ₂₈ H ₂₇ N ₃ O ₄		
Molecular Weight:	469.53		
Target:	Others		
Pathway:	Others		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 62.5 mg/mL (133.11 mM; ultrasonic and warming and heat to 60°C)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.1298 mL	10.6489 mL	21.2979 mL
		5 mM	0.4260 mL	2.1298 mL	4.2596 mL
10 mM		0.2130 mL	1.0649 mL	2.1298 mL	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (4.43 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	(Rac)-Cemsidomide ((Rac)-CFT7455) is a zinc finger transcription factors Ikaros (IKZF1) and Aiolos (IKZF3) degrader, acting via the ubiquitin proteasome pathway, with a GI ₅₀ of 0.05 nM for NCIH929.1 cells. (Rac)-Cemsidomide is the racemic isomer of Cemsidomide (HY-144841A) which is a IKZF1/IKZF3 degrader with anticancer activity ^{[1][2]} .
IC₅₀ & Target	Ikaros, Aiolos ^[1]

REFERENCES

[1]. David Proia et al. Advantageous therapies for disorders mediated by ikaros or aiolos. WO2022032132A1 (compound 5)

[2]. James A. Henderson, et al. Abstract LB007: CFT7455: A novel, IKZF1/3 degrader that demonstrates potent tumor regression in IMiD-resistant multiple myeloma (MM) xenograft models. Cancer Res (2021) 81 (13_Supplement): LB007.

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

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