## CB1 inverse agonist 2

Cat. No.:	HY-12095	
CAS No.:	1019839-52-1	
Molecular Formula:	C <sub>24</sub> H <sub>20</sub> CIFN <sub>2</sub> OS	/=
Molecular Weight:	438.94	$\searrow$
Target:	Cannabinoid Receptor	
Pathway:	GPCR/G Protein; Neuronal Signaling	M H S
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

Description	CB1 inverse agonist 2 is an orally active inverse agonist of Cannabinoid Receptor CB1. CB1 inverse agonist 2 effectively inhibits CP55940-induced hypothermia and anorexia in mice model <sup>[1]</sup> .			
IC <sub>50</sub> & Target	СВ1 9.7 (рКі)	CB2 6.0 (pKi)		
In Vitro	CB1 inverse agonist 2 (compound 12 h) shows selectivity on CB1 over CB2 with pK <sub>i</sub> values of 9.7 and 6.0, respectively <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			
In Vivo	CB1 inverse agonist 2 (compound 12 h) (1 mg/kg; i.p.; single dose) results 70% inhibition against hypothermia, as well as (1 mg/kg; p.o.; single dose) resulting 59% inhibition against food intake <sup>[1]</sup> . CB1 inverse agonist 2 (10 mg/kg; p.o.; once daily for 14 d) produces robust weight loss of approximately 13% of initial body weight in mouse <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

## REFERENCES

[1]. Pettersson H, et al. Synthesis and evaluation of dibenzothiazepines: a novel class of selective cannabinoid-1 receptor inverse agonists. J Med Chem. 2009 Apr 9;52(7):1975-82.

[2]. Synthesis and Evaluation of Dibenzothiazepines: A Novel Class of Selective Cannabinoid-1 Receptor Inverse Agonists J. Med. Chem., 2009, 52 (7), pp 1975-1982

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

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**Product** Data Sheet



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