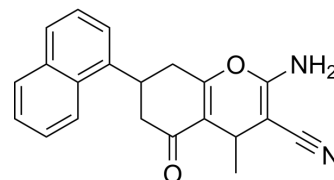


UCPH-102

Cat. No.:	HY-118858
CAS No.:	1229591-56-3
Molecular Formula:	C ₂₁ H ₁₈ N ₂ O ₂
Molecular Weight:	330.38
Target:	EAAT
Pathway:	Membrane Transporter/Ion Channel
Storage:	4°C, protect from light, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light, stored under nitrogen)



SOLVENT & SOLUBILITY

In Vitro

DMSO : 8 mg/mL (24.21 mM; Need ultrasonic and warming)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	3.0268 mL	15.1341 mL	30.2682 mL
5 mM	0.6054 mL	3.0268 mL	6.0536 mL
10 mM	0.3027 mL	1.5134 mL	3.0268 mL

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

BIOLOGICAL ACTIVITY

Description

UCPH-102 is a highly selective EAAT1 inhibitor with an IC₅₀ of 0.43 μM. UCPH-102 exhibits a specific anti-proliferative effect on T-ALL cells. UCPH-102 also shows good blood-brain permeability, which can be used in studies of amyotrophic lateral sclerosis, Alzheimer's disease, chronic pain and obsessive compulsive disorder^{[1][2][3][4]}.

IC₅₀ & Target

EAAT1

REFERENCES

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[2]. Haym I, et al. Bioavailability Studies and in vitro Profiling of the Selective Excitatory Amino Acid Transporter Subtype 1 (EAAT1) Inhibitor UCPH-102. *ChemMedChem*. 2016 Feb 17;11(4):403-19.

[3]. Stanulović V S, et al. Proliferation and Survival of T-cell Acute Lymphoblastic Leukaemia Depends on mTOR-regulated Glutamine Uptake and EAAT1-dependent Conversion of Glutamine to Aspartate and Nucleotides. *bioRxiv*. 2020.

[4]. Abrahamsen B, et al. Allosteric modulation of an excitatory amino acid transporter: the subtype-selective inhibitor UCPH-101 exerts sustained inhibition of EAAT1 through an intramonomeric site in the trimerization domain. J Neurosci. 2013 Jan 16;33(3):1068-87.

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

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