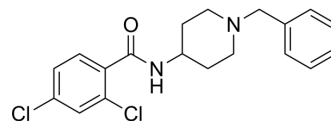


BPDBA

Cat. No.:	HY-119591		
CAS No.:	312281-74-6		
Molecular Formula:	C ₁₉ H ₂₀ Cl ₂ N ₂ O		
Molecular Weight:	363.28		
Target:	GABA Receptor		
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling		
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 : 100 mg/mL (275.27 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	2.7527 mL	13.7635 mL	27.5270 mL
5 mM	0.5505 mL	2.7527 mL	5.5054 mL
10 mM	0.2753 mL	1.3763 mL	2.7527 mL

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

BIOLOGICAL ACTIVITY

Description

BPDBA is a selective and noncompetitive betaine/GABA transporter (BGT-1) inhibitor with IC₅₀s of 20 μM and 35 μM against human BGT-1 and mouse GAT2, respectively^[1].

In Vivo

BPDBA is predicted to have excellent oral absorption and blood brain barrier penetration properties^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Kragholm B, et al. Discovery of a subtype selective inhibitor of the human betaine/GABA transporter 1 (BGT-1) with a non-competitive pharmacological profile. *Biochem Pharmacol.* 2013 Aug 15;86(4):521-8.

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