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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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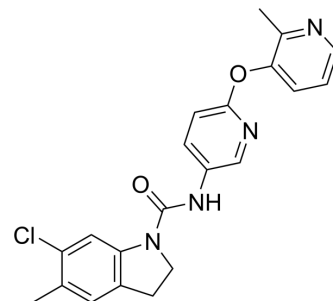
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SB 242084

Cat. No.:	HY-13409
CAS No.:	181632-25-7
Molecular Formula:	C ₂₁ H ₁₉ ClN ₄ O ₂
Molecular Weight:	394.85
Target:	5-HT Receptor
Pathway:	GPCR/G Protein; Neuronal Signaling
Storage:	Powder -20°C 3 years 4°C 2 years In solvent -80°C 2 years -20°C 1 year



SOLVENT & SOLUBILITY

In Vitro

DMSO : ≥ 44 mg/mL (111.43 mM)
 * "≥" means soluble, but saturation unknown.

	Solvent Concentration	Mass	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM		2.5326 mL	12.6630 mL	25.3261 mL
	5 mM		0.5065 mL	2.5326 mL	5.0652 mL
	10 mM		0.2533 mL	1.2663 mL	2.5326 mL

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

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
 Solubility: ≥ 2.5 mg/mL (6.33 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
 Solubility: ≥ 2.5 mg/mL (6.33 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

SB 242084 is a selective, competitive and high-affinity (pK_i=9.0) 5-HT_{2C} receptor antagonist (crosses the blood-brain barrier). SB 242084 increases basal activity of dopaminergic neurons in the ventral tegmental area (VTA) of the midbrain and dopamine release in the vomeronasal nucleus. SB 242084 also increases mitochondrial gene expression and oxidative metabolism via 5-HT_{2A} receptor. SB 242084 has good research potential in the negative symptoms of anxiety, depression and schizophrenia, as well as in acute organ damage^{[1][2][3]}.

IC₅₀ & Target

5-HT_{2C} Receptor
 9.0 (pKi)

In Vitro

SB 242084 (100 nM; 45 min) exhibits antagonism of the 5-HT stimulated increase in phosphatidylinositol hydrolysis at the human 5-HT_{2C} receptor in SH-SY5Y cells^[1].

SB 242084 (1-100 nM; 24 h) increases RPTC respiration and PGC-1 α mRNA expression in RPTC^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Viability Assay^[1]

Cell Line:	SH-SY5Y cells
Concentration:	100 nM
Incubation Time:	45 min
Result:	Antagonized 5-HT induced concentration-related increase in PI hydrolysis.

RT-PCR^[2]

Cell Line:	RPTC cells
Concentration:	1-100 nM
Incubation Time:	24 h
Result:	Increased FCCP-uncoupled respiration and PGC-1 α mRNA expression.

In Vivo

SB 242084 (0.1-1 mg/kg; i.p.; single; 20 min pre-test) improves the behavior of rats in social interaction tests^[1].

SB 242084 (5 mg/kg; i.p.; single; 20 min pre-test) improves mCPP-induced hypophagia in rats^[1].

SB 242084 (5, 10 mg/kg; i.p.; single) increases the levels of basal dialysate dopamine (DA) and dihydroxyphenylacetic acid (DOPAC) in the nucleus accumbens of rats^[3].

SB 242084 (160-640 μ g/kg; i.v.; single) dose-dependently and significantly increases the basal firing rate of VTA (ventral tegmental area) dopaminergic neurons, and the bursting activity is also enhanced in the same area, in vivo^[3].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male Sprague-Dawley (CD) rats ^[1] .
Dosage:	0.1-1 mg/kg
Administration:	Intraperitoneal injection; single; 20 min pre-test
Result:	Significantly increased the amount of time rats spent in social interaction over 15 min under brightly lit conditions and in an unfamiliar test box.

Animal Model:	Male Sprague-Dawley (CD) rats (mCPP-induced hypophagia model) ^[1] .
Dosage:	5 mg/kg
Administration:	Intraperitoneal injection; single; 20 min pre-test
Result:	Significantly reduced the amount of food consumed by 23 h food-deprived rats over a 1 hr test period from the time of food presentation.

Animal Model:	Rats ^[2] .
Dosage:	5, 10 mg/kg
Administration:	Intraperitoneal injection; single

Result:

Significantly increased basal dialysate dopamine (DA) and dihydroxyphenylacetic acid (DOPAC) in the nucleus accumbens.

CUSTOMER VALIDATION

- Authorea. September 19, 2022.

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REFERENCES

- [1]. Kennett GA, et al. SB 242084, a selective and brain penetrant 5-HT_{2C} receptor antagonist. *Neuropharmacology*. 1997 Apr-May;36(4-5):609-20.
- [2]. Harmon JL, et al. 5-HT₂ Receptor Regulation of Mitochondrial Genes: Unexpected Pharmacological Effects of Agonists and Antagonists. *J Pharmacol Exp Ther*. 2016 Apr;357(1):1-9.
- [3]. Kennett GA, et al. SB 242084, a selective and brain penetrant 5-HT_{2C} receptor antagonist. *Neuropharmacology*. 1997 Apr-May;36(4-5):609-20.

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

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