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## Produktinformation



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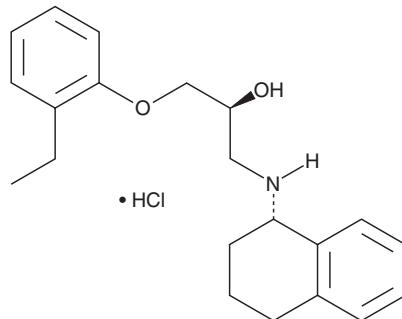
# PRODUCT INFORMATION



## SR 59230A (hydrochloride)

Item No. 21407

CAS Registry No.:	1135278-41-9
Formal Name:	(2S)-1-(2-ethylphenoxy)-3-[(1S)-1,2,3,4-tetrahydro-1-naphthalenyl]amino]-2-propanol, monohydrochloride
MF:	C <sub>21</sub> H <sub>27</sub> NO <sub>2</sub> • HCl
FW:	361.9
Purity:	≥99%
Supplied as:	A solid
Storage:	Room temperature
Stability:	≥1 year



Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

### Laboratory Procedures

SR 59230A (hydrochloride) is supplied as a solid. A stock solution may be made by dissolving the SR 59230A (hydrochloride) in the solvent of choice. SR 59230A (hydrochloride) is soluble in the organic solvent DMSO, which should be purged with an inert gas, at a concentration of up to approximately 100 mM. It is also soluble in water. The solubility of SR 59230A (hydrochloride) in water is up to approximately 10 mM. We do not recommend storing the aqueous solution for more than one day.

### Description

SR 59230A (hydrochloride) is a β<sub>3</sub>-adrenergic receptor (β<sub>3</sub>-AR) antagonist (pA<sub>2</sub>s = 8.76, 7.31, and 6.63 in rat proximal colon, guinea pig atrium, and guinea pig trachea, respectively).<sup>1</sup> It is less selective for β<sub>3</sub>-AR in cells transfected with the human β-AR subtypes (K<sub>i</sub>s = 16.4, 61.9, and 122 nM for β<sub>1</sub>-, β<sub>2</sub>-, and β<sub>3</sub>-AR, respectively).<sup>2</sup> At low concentrations, SR 59230A blocks MDMA-induced hyperthermia, while at high concentrations it blocks hyperthermia but also increases heat loss through an α<sub>1</sub>-AR antagonistic mechanism.<sup>3</sup> In adipocytes, it induces phosphorylation of p38 MAPK via the G<sub>s</sub> pathway.<sup>4</sup> It has also been used in studies of heart failure to elucidate the role of the β<sub>3</sub>-ARs.<sup>5</sup>

### References

1. Manara, L., Badone, D., Baroni, M., et al. Functional identification of rat atypical β-adrenoceptors by the first β<sub>3</sub>-selective antagonists, aryloxypropanolaminotetralins. *Br. J. Pharmacol.* **117**(3), 435-442 (1996).
2. Hoffman, C., Leitz, M.R., Oberdorf-Maass, S., et al. Comparative pharmacology of human β-adrenergic receptor subtypes—characterization of stably transfected receptors in CHO cells. *Naunyn-Schmiedeberg's Arch. Pharmacol.* **369**(2), 151-159 (2004).
3. Bexis, S. and Docherty, J.R. Role of α<sub>1</sub>- and β<sub>3</sub>-adrenoceptors in the modulation by SR59230A of the effects of MDMA on body temperature in the mouse. *Br. J. Pharmacol.* **158**(1), 259-266 (2009).
4. Mizuno, K., Kanda, Y., Kuroki, Y., et al. Stimulation of β<sub>3</sub>-adrenoceptors causes phosphorylation of p38 mitogen-activated protein kinase via a stimulatory G protein-dependent pathway in 3T3-L1 adipocytes. *Br. J. Pharmacol.* **135**(4), 951-960 (2002).
5. Gan, R.T., Li, W.M., Xiu, C.H., et al. Chronic blocking of β<sub>3</sub>-adrenoceptor ameliorates cardiac function in rat model of heart failure. *Chin. Med. J. (Engl.)* **120**(24), 2250-2255 (2007).

#### WARNING

THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

#### SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

#### WARRANTY AND LIMITATION OF REMEDY

Buyer agrees to purchase the material subject to Cayman's Terms and Conditions. Complete Terms and Conditions including Warranty and Limitation of Liability information can be found on our website.

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