

# Produktinformation



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# PRODUCT INFORMATION



## Lasofoxifene

Item No. 21754

CAS Registry No.: 180916-16-9

Formal Name: (5R,6S)-5,6,7,8-tetrahydro-6-

phenyl-5-[4-[2-(1-pyrrolidinyl)

ethoxy]phenyl]-2-naphthalenol

Synonym: CP 336,156 MF:  $C_{28}H_{31}NO_{2}$ FW: 413.6 **Purity:** ≥98%

Supplied as: A crystalline solid

Storage: -20°C Stability: ≥2 years

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



Lasofoxifene is supplied as a crystalline solid. A stock solution may be made by dissolving the lasofoxifene in the solvent of choice. Lasofoxifene is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF), which should be purged with an inert gas. The solubility of lasofoxifene in ethanol is approximately 1.5 mg/ml and approximately 20 mg/ml in DMSO and DMF.

Lasofoxifene is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, lasofoxifene should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. Lasofoxifene has a solubility of approximately 0.5 mg/ml in a 1:1 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

### Description

Lasofoxifene is a third-generation, non-steroidal selective estrogen receptor modulator (SERM). It selectively binds to human ER $\alpha$  with an IC<sub>50</sub> value of 1.5 nM and inhibits bone loss in ovariectomized rats.<sup>1</sup> In clinical studies of postmenopausal osteoporosis, 0.5 mg/day lasofoxifene was associated with reduced risks of nonvertebral and vertebral fractures, ER-positive breast cancer, coronary heart disease, and stroke but an increased risk of venous thromboembolic events.<sup>2,3</sup> Lasofoxifene has also been shown to act as an inverse agonist at the CB2 cannabinoid receptor, indicating its potential to be repurposed as a therapeutic for indications wherein  $C\bar{B}_2$  is a target.<sup>4</sup>

#### References

- 1. Ke, H. Z., Paralkar, V. M., Grasser, W. A., et al. Effects of CP-336,156, a new, nonsteroidal estrogen agonist/ antagonist, on bone, serum cholesterol, uterus, and body composition in rat models. Endocrinology **139(4)**, 2068-2076 (1998).
- Cummings, S. R., Ensrud, K., Delmas, P. D., et al. Lasofoxifene in postmenopausal women with osteoporosis. N. Engl. J. Med. 362(8), 686-696 (2010).
- Gennari, L., Merlotti, D., and Nuti, R. Selective estrogen receptor modulator (SERM) for the treatment of osteoporosis in postmenopausal women: Focus on lasofoxifene. Clin. Interv. Aging 5, 19-29 (2010).
- Kumar, P. and Song, Z.H. CB2 cannabinoid receptor is a novel target for third-generation selective estrogen receptor modulators bazedoxifene and lasofoxifene. Biochem. Bioph. Res. Co. 443(1), 144-149 (2014).

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

al 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

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