Phone: 888-558-5227

651-644-8424

Fax: 888-558-7329

Email: getinfo@lktlabs.com Web: lktlabs.com

Product Information

Product ID R0243

CAS No. 82640-04-8

Chemical Name [6-hydroxy-2-(4-hydroxyphenyl)benzo[b]thien-3-yl]- [4-[2-(1-

piperidinyl) ethoxy] phenyl]methanone hydrochloride

Synonym Evista

Formula C₂₈H₂₇NO₄S • HCl

Formula Wt. 510.05 Melting Point 258°C Purity ≥98%

Solubility Very slightly soluble in water. Soluble in DMSO at 28 mg/mL.

Pricing and Availability

Bulk quanitites available upon request

Product ID	Size	List Price
R0243	50 mg	\$82.10
R0243	250 mg	\$156.20
R0243	500 mg	\$242.70
R0243	1 g	\$433.30

Store Temp Ambient Ship Temp Ambient

Description Raloxifene is a selective estrogen receptor modulator (SERM) that exhibits agonist activity in bone and antagonist activity in uterine and breast tissues. Raloxifene is clinically used to prevent osteoporosis in post-menopausal women and to treat ER+ breast cancer. This compound exhibits anti-osteoporotic, anti-parasitic, neuromodulatory, and anticancer chemotherapeutic activities. Raloxifene decreases fracture risk and enhances bone strength in clinical settings. In vitro and in vivo, raloxifene inhibits growth of Leishmania and parasitic burden. Additionally, raloxifene increases uptake of glutamate and expression of GLT-1 in astrocytes. In animal models of breast cancer, this compound decreases expression of EGFR, vascular density, and tumor growth.

References Karki P, Webb A, Zerguine A, et al. Mechanism of raloxifene-induced upregulation of glutamate transporters in rat primary astrocytes. Glia. 2014 Aug;62(8):1270-83. PMID: 24782323.

> Reimão JQ, Miguel DC, Taniwaki NN, et al. Antileishmanial activity of the estrogen receptor modulator raloxifene. PLoS Negl Trop Dis. 2014 May 8;8(5):e2842. PMID: 24810565.

Gallant MA, Brown DM, Hammond M, et al. Bone cell-independent benefits of raloxifene on the skeleton: a novel mechanism for improving bone material properties. Bone. 2014 Apr;61:191-200. PMID: 24468719.

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Caution: This product is intended for laboratory and research use only. It is not for human or drug use.