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## **Product Information**

Product ID CAS No.	R0243 82640-04-8	HO HO HO N		
Chemical Name	[6-hydroxy-2-(4-hydroxyphenyl)benzo[b]thien-3-yl]- [4-[ 2-(1- piperidinyl) ethoxy] phenyl]methanone hydrochloride			
Synonym	Evista			
Formula	C <sub>28</sub> H <sub>27</sub> NO <sub>4</sub> S • HCI			
Formula Wt.	510.05			
Melting Point	258°C	Bulk quanitites available upon request		
Purity		Product ID	Size	
Solubility	Very slightly soluble in water. Soluble in DMSO at	R0243	50 mg	
		R0243	250 mg	
		R0243	500 mg	
Store Temp	Ambient	R0243	1 g	
Ship Temp	Ambient			
Description	Relaxifene is a selective estrogen recentor modulator (SFRM) that exhibits agonist activity in bone and antagonist activity in			

**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.

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