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## **Product Information**

Product ID D6957

CAS No. 82413-20-5

Chemical Name 3-Hydroxytamoxifen

Synonym E-Droloxifene; 3-HYDROXYTAMOXIFEN; Droloxifenum [Latin]; Droloxifeno

[Spanish]; FK-435

Formula C<sub>26</sub>H<sub>29</sub>NO<sub>2</sub> Formula Wt. 387.51 Melting Point 127-129°C

Purity ≥98% Solubility

## **Pricing and Availability**

Bulk quanitites available upon request

Product ID	Size	List Price
D6957	5 mg	\$71.70
D6957	25 mg	\$134.40
D6957	100 mg	\$413.60
D6957	250 mg	\$723.60

Store Temp Ambient Ship Temp Ambient

Description Droloxifene is an anti-estrogen triphenylthylene selective estrogen receptor modulator (SERM); it is an analog of tamoxifen. Droloxifene acts as an estrogen receptor agonist in bone and as an estrogen receptor antagonist in breast tissue. As a result of these actions, this compound exhibits anticancer chemotherapeutic, chemopreventive, anti-osteoporotic, and anti-resorptive benefits. In breast cancer cells, droloxifene decreases production of NO and increases levels of pro-apoptotic TGF-β. Droloxifene increases apoptosis in luteal cells in vivo, increasing levels of c-myc and increasing the Bax/Bcl-2 ratio. In ovariectomized rats that act as a post-menopausal animal model, droloxifene inhibits bone resorption and turnover as well as estrogen-related bone loss. This compound also decreases levels of E-selectin and increases levels if vascular cellular adhesion molecule 1 (VCAM-1) in clinical settings. TEST!!!!!!

References Shelly W, Draper MW, Krishnan V, et al. Selective estrogen receptor modulators: an update on recent clinical findings. Obstet Gynecol Surv. 2008 Mar;63(3):163-81. PMID: 18279543.

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Knabbe C, Zugmaier G, Schmahl M, et al. Induction of transforming growth factor beta by the antiestrogens droloxifene, tamoxifen, and toremifene in MCF-7 cells. Am J Clin Oncol. 1991;14 Suppl 2:S15-20. PMID: 1835818.

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