



LKT Laboratories, Inc.

(Z)-4-Hydroxytamoxifen

Phone: 888-558-5227
651-644-8424
Fax: 888-558-7329
Email: getinfo@lktlabs.com
Web: lktlabs.com

Product Information

Product ID H9711

CAS No. 68047-06-3

Chemical Name

Synonym

Formula $C_{26}H_{29}NO_2$

Formula Wt. 387.51

Melting Point 142-144 °C

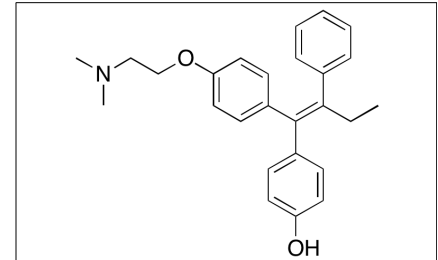
Purity ≥98%

Solubility Soluble in DMSO, ethanol
(20mg/ml) or methanol
(10mg/ml); insoluble water.

Store Temp -20 °C

Ship Temp Ambient

Description (Z)-4-Hydroxytamoxifen is the more active isomer of 4-hydroxytamoxifen. 4-Hydroxytamoxifen is the active metabolite of tamoxifen, a selective estrogen receptor modulator (SERM) that acts as an agonist or antagonist in various tissues. 4-Hydroxytamoxifen exhibits anticancer chemotherapeutic activity, inducing autophagy and vacuole formation as well as KRAS degradation in various cancer cell lines. In cardiac myocytes, 4-hydroxytamoxifen decreases Ca^{2+} amplitude, slowing relaxation and decreasing contractility.



Bulk quantities available upon request

Product ID	Size
H9711	5 mg
H9711	10 mg
H9711	25 mg

References Asp ML, Martindale JJ, Metzger JM. Direct, differential effects of tamoxifen, 4-hydroxytamoxifen, and raloxifene on cardiac myocyte contractility and calcium handling. *PLoS One*. 2013 Oct 24;8(10):e78768. PMID: 24205315.

Kohli L, Kaza N, Coric T, et al. 4-Hydroxytamoxifen induces autophagic death through K-Ras degradation. *Cancer Res*. 2013 Jul 15;73(14):4395-405. PMID: 23722551.

Schwartz JA, Zhong L, Deighton-Collins S, et al. Mutations targeted to a predicted helix in the extreme carboxyl-terminal region of the human estrogen receptor-alpha alter its response to estradiol and 4-hydroxytamoxifen. *J Biol Chem*. 2002 Apr 12;277(15):13202-9. Erratum in: *J Biol Chem* 2002 Jul 5;277(27):24842. PMID: 11823467.

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