



LKT Laboratories, Inc.

7-Ethyl-10-hydroxycamptothecin

Phone: 888-558-5227

651-644-8424

Fax: 888-558-7329

Email: getinfo@lktlabs.com

Web: lktlabs.com

Product Information

Product ID C0154

CAS No. 86639-52-3

Chemical Name 4,11-diethyl-4,9-dihydroxy-1H-Pyrano[3',4':6,7]-indolizino[1,2-b]quinoline-3,14(4H,12H)-dione

Synonym SN-38, 7-ethyl-10-hydroxy-CPT

Formula C₂₂H₂₀N₂O₅

Formula Wt. 392.40

Melting Point 228-235 °C

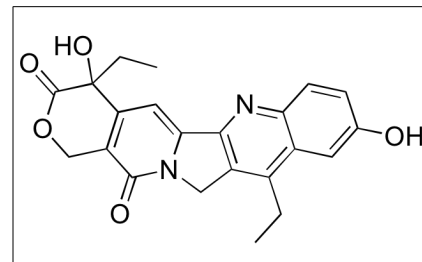
Purity ≥98%

Solubility

Store Temp -20 °C

Ship Temp Ambient

Description SN-38 is a camptothecin derivative. Camptothecin is a quinolone alkaloid precursor of irinotecan; it is originally found in *Camptotheca*. Camptothecin exhibits anticancer chemotherapeutic activity, inhibiting topoisomerase I and inducing double-stranded DNA breaks. Camptothecin is extremely cytotoxic and only its derivatives are clinically used as chemotherapeutics.



Bulk quantities available upon request

Product ID **Size**

C0154 5 mg

C0154 10 mg

C0154 50 mg

C0154 100 mg

References Rodríguez-Berna G, Mangas-Sanjuán V, Gonzalez-Alvarez M, et al. A promising camptothecin derivative: Semisynthesis, antitumor activity and intestinal permeability. *Eur J Med Chem.* 2014 Jun 25;83C:366-373. PMID: 24980118.

Berniak K, Rybak P, Bernas T, et al. Relationship between DNA damage response, initiated by camptothecin or oxidative stress, and DNA replication, analyzed by quantitative 3D image analysis. *Cytometry A.* 2013 Jul 11. [Epub ahead of print]. PMID: 23846844.

Redinbo MR, Stewart L, Kuhn P, et al. Crystal structures of human topoisomerase I in covalent and noncovalent complexes with DNA. *Science.* 1998 Mar 6;279(5356):1504-13. PMID: 9488644.

Blackwood E, Epler J, Yen I, et al. Combination drug scheduling defines a "window of opportunity" for chemopotiation of gemcitabine by an orally bioavailable, selective ChK1 inhibitor, GNE-900. *Mol Cancer Ther.* 2013 Oct;12(10):1968-1980. PMID: 23873850.

Walton MI, Eve PD, Hayes A, et al. The preclinical pharmacology and therapeutic activity of the novel CHK1 inhibitor SAR-020106. *Mol Cancer Ther.* 2010 Jan;9(1):89-100. PMID: 20053762.

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