



LKT Laboratories, Inc.
Canertinib Dihydrochloride

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

Product Information

Product ID C0252

CAS No. 289499-45-2

Chemical Name

Synonym CI1033

Formula $C_{24}H_{25}ClFN_5O_3 \cdot 2HCl$

Formula Wt. 558.86

Melting Point

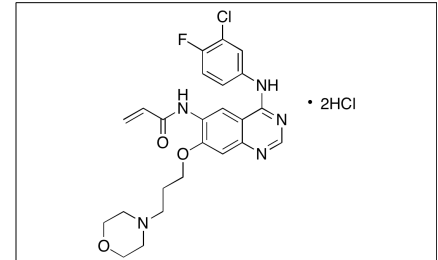
Purity $\geq 98\%$

Solubility

Store Temp Ambient

Ship Temp Ambient

Description Canertinib is an irreversible inhibitor of EGFR (HER2/ErbB). This compound is currently in clinical trials as an anticancer chemotherapeutic, displaying activity against a variety of cancers. In a cellular model of melanoma, canertinib exhibits cytostatic activity, stopping cell growth in the G1 phase, but not inducing apoptosis except at very high concentrations. In a related animal model, canertinib inhibits tumor growth. Separately, canertinib inhibits Akt and ERK1/2 in cells lacking ErbB receptors and induces caspase-mediated apoptosis.



Bulk quantities available upon request

Product ID	Size
C0252	25 mg
C0252	100 mg
C0252	250 mg

References Djerf Severinsson EA, Trinks C, Gréen H, et al. The pan-ErbB receptor tyrosine kinase inhibitor canertinib promotes apoptosis of malignant melanoma in vitro and displays anti-tumor activity in vivo. *Biochem Biophys Res Commun.* 2011 Oct 28;414(3):563-8. PMID: 21982771.

Trinks C, Severinsson EA, Holmlund B, et al. The pan-ErbB tyrosine kinase inhibitor canertinib induces caspase-mediated cell death in human T-cell leukemia (Jurkat) cells. *Biochem Biophys Res Commun.* 2011 Jul 8;410(3):422-7. PMID: 21669187.

Zinner RG, Nemunaitis J, Eiseman I, et al. Phase I clinical and pharmacodynamic evaluation of oral CI-1033 in patients with refractory cancer. *Clin Cancer Res.* 2007 May 15;13(10):3006-14. PMID: 17505003.

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