



LKT Laboratories, Inc.

Gefitinib

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Product Information

Product ID G1721

CAS No. 184475-35-2

Chemical Name N-(3-chloro-4-fluorophenyl)-7-methoxy-6-(3-morpholin-4-ylpropoxy)quinazolin-4-amine

Synonym

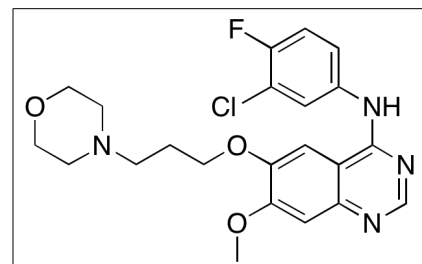
Formula C₂₂H₂₄ClFN₄O₃

Formula Wt. 446.90

Melting Point 212-214°C

Purity ≥98%

Solubility DMSO (20 mg/ml), DMF (20 mg/ml)



Pricing and Availability

Bulk quantities available upon request

Product ID	Size	List Price
G1721	100 mg	\$171.30
G1721	250 mg	\$301.70
G1721	1 g	\$880.30

Store Temp Ambient

Ship Temp Ambient

Description Gefitinib is an anticancer chemotherapeutic compound that inhibits mutant isoforms of EGFR. Across a variety of cancer cell lines, gefitinib increases poly(ADP)-ribose polymerase (PARP) cleavage and apoptosis, resulting in inhibition of cell growth. Gefitinib also induces expression of PPAR γ in cellular models, likely through activation of transcription factor CCAT/enhancer binding protein β (CEBP- β). Gefitinib's inhibition of EGFR also results in inhibition of heat shock protein 70 (HSP70), which can exacerbate pulmonary fibrosis. Additionally, gefitinib increases phosphorylation of p38 MAPK and JNK, inducing apoptosis in keratinocytes. This compound also promotes differentiation of acute myelogenous leukemia cells in vitro.

References Mansure JJ, Nassim R, Chevalier S, et al. A novel mechanism of PPAR gamma induction via EGFR signalling constitutes rational for combination therapy in bladder cancer. PLoS One. 2013;8(2):e55997. doi: 10.1371/journal.pone.0055997. Erratum in: PLoS One. 2013; 8(5). PMID: 23409107.

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