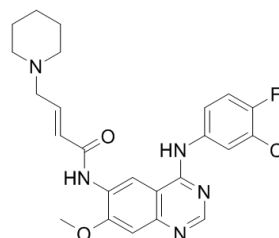


<b>Product Name</b>	: Dacomitinib
<b>Synonyms</b>	: PF-00299804;PF-299804;PF 299804;PF 00299804
<b>Cat No.</b>	: M10413
<b>CAS Number</b>	: 1110813-31-4
<b>Molecular Formula</b>	: C <sub>24</sub> H <sub>25</sub> ClFN <sub>5</sub> O <sub>2</sub>
<b>Formula Weight</b>	: 469.90
<b>Chemical Name</b>	: 2-Butenamide, N-[4-[(3-chloro-4-fluorophenyl)amino]-7-methoxy-6-quinazoliny]-4-(1-piperidinyl)-, (2E)-



**Description** : Dacomitinib (PF-00299804, PF-299804) is a potent, irreversible, orally active pan-ErbB receptor tyrosine kinase inhibitor with IC<sub>50</sub> of 6, 45.7 and 73.7 for EGFR, ERBB2 and ERBB4, respectively; irreversibly inhibits erbB tyrosine kinase activity through binding at the ATP site and covalent modification of nucleophilic cysteine residues in the catalytic domains of erbB family members inhibits erbB1 autophosphorylation in the A431 human squamous cell carcinoma line with IC<sub>50</sub> of 15.1 nM; causes significant antitumor activity, including marked tumor regressions in a variety of human tumor xenograft models that express and/or overexpress erbB family members or contain the double mutation (L858R/T790M) in EGFR associated with resistance to gefitinib and erlotinib. Lung Cancer Phase 3 Clinical

<b>Pathway</b>	: Angiogenesis
<b>Target</b>	: EGFR
<b>Receptor</b>	: EGFR;HER2/ErbB2
<b>Solubility</b>	: DMSO: ≥ 50 mg/mL (Need ultrasonic)
<b>SMILES</b>	: <chem>COC1=C(C=C2C(=C1)N=CN=C2NC3=CC(=C(C=C3)F)Cl)NC(=O)/C=C/CN4CCCCC4</chem>
<b>Storage</b>	: (-20°C)
<b>Stability</b>	: ≥ 2 years
<b>Reference</b>	:

1. Engelman JA, et al. Cancer Res. 2007 Dec 15;67(24):11924-32. | 2. Gonzales AJ, et al. Mol Cancer Ther. 2008 Jul; 7(7):1880-9. | 3. Nguyen KS, et al. Clin Lung Cancer. 2009 Jul;10(4):281-9. | 4. Kalous O, et al. Mol Cancer Ther. 2012 Sep; 11(9):1978-87.