

Product Name	: Sorafenib
Synonyms	: Bay 43-9006; Bay 43-9006
Cat No.	: M13866
CAS Number	: 284461-73-0
Molecular Formula	: C ₂₁ H ₁₆ ClF ₃ N ₄ O ₃
Formula Weight	: 464.83
Chemical Name	: 2-Pyridinecarboxamide, 4-[4-[[[4-chloro-3-(trifluoromethyl)phenyl]amino]carbonyl]amino]phenoxy]-N-methyl-
Description	: A potent, orally available Raf inhibitor with IC ₅₀ of 6, 22, and 38 nM for Raf-1, wt Braf, and Braf V599E, respectively; Also demonstrates potent inhibition of certain proangiogenic RTKs, including VEGFR-2, PDGFR-β, VEGFR-3, Flt-3, c-Kit (IC ₅₀ <100 nM); Exhibits broad spectrum oral antitumor activity and targets the RAF/MEK/ERK pathway and receptor tyrosine kinases involved in tumor progression and angiogenesis. Kidney Cancer Approved
Pathway	: MAPK/ERK Signaling
Target	: Raf
Receptor	: B-Raf; B-Raf(V599E); Raf-1; VEGFR2/Flk1; VEGFR3
Solubility	: DMSO: ≥ 45 mg/mL
SMILES	: <chem>CNC(=O)C1=NC=CC(OC2=CC=C(NC(=O)NC3=CC(=C(Cl)C=C3)C(F)(F)F)C=C2)=C1</chem>
Storage	: (-20°C)
Stability	: ≥ 2 years
Reference	:

1. Wilhelm SM, et al. Cancer Res. 2004 Oct 1;64(19):7099-109. | 2. Carlomagno F, et al. J Natl Cancer Inst. 2006 Mar 1;98(5):326-34. | 4. Lyons JF, et al. Endocr Relat Cancer. 2001 Sep;8(3):219-25.