

<b>Product Name</b>	: eCF506
<b>Synonyms</b>	: —
<b>Cat No.</b>	: M21978
<b>CAS Number</b>	: 1914078-41-3
<b>Molecular Formula</b>	: C <sub>26</sub> H <sub>38</sub> N <sub>8</sub> O <sub>3</sub>
<b>Formula Weight</b>	: 510.63
<b>Chemical Name</b>	: —
<b>Description</b>	eCF506 is a potent and selective inhibitor of SRC (IC <sub>50</sub> < 0.5 nM). eCF506, the first small molecule with subnanomolar IC <sub>50</sub> for SRC that requires 3 orders of magnitude greater concentration to inhibit ABL. eCF506 exhibits excellent water solubility, an optimal DMPK profile and oral bioavailability, halts SRC-associated neuromast migration in zebrafish embryos without inducing life-threatening heart defects, and inhibits SRC phosphorylation in tumor xenografts in mice.
<b>Pathway</b>	: Tyrosine Kinase
<b>Target</b>	: Src
<b>Receptor</b>	: Src
<b>Solubility</b>	: DMSO: 62.5 mg/mL (122.40 mM; Need ultrasonic)
<b>SMILES</b>	: <chem>COc1cc(ccc1NC(=O)OC(C)(C)C)-c1nn(CCN2CCC(CC2)N(C)C)c2ncnc(N)c12</chem>
<b>Storage</b>	: (-20°C)
<b>Stability</b>	: ≥ 2 years
<b>Reference</b>	:

1. Fraser C, et al. Rapid Discovery and Structure-Activity Relationships of Pyrazolopyrimidines That Potently Suppress Breast Cancer Cell Growth via SRC Kinase Inhibition with Exceptional Selectivity over ABL Kinase. *J Med Chem*. 2016 May 26;59(10):4697-710.