

Product Name: FRAX 597

Catalog No.: 6029

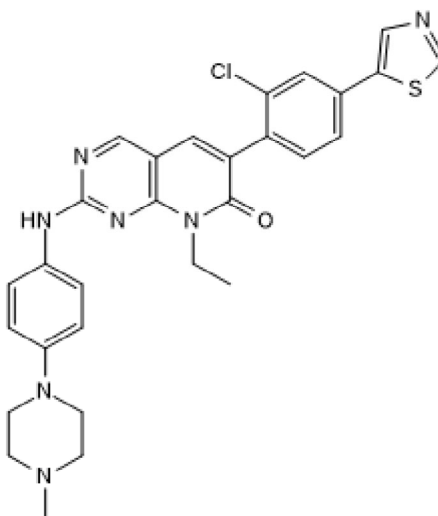
Batch No.: 2

CAS Number: 1286739-19-2

IUPAC Name: 6-[2-Chloro-4-(5-thiazolyl)phenyl]-8-ethyl-2-[[4-(4-methyl-1-piperazinyl)phenyl]amino]pyrido[2,3-*d*]pyrimidin-7-(8*H*)-one

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula:	C ₂₉ H ₂₈ ClN ₇ OS.
Batch Molecular Weight:	558.1
Physical Appearance:	Yellow solid
Solubility:	DMSO to 20 mM
Storage:	Store at -20°C
Batch Molecular Structure:	



2. ANALYTICAL DATA

HPLC:	Shows 99.3% purity
¹H NMR:	Consistent with structure
Mass Spectrum:	Consistent with structure

Microanalysis:	Carbon Hydrogen Nitrogen		
Theoretical	62.41	5.06	17.57
Found	62.17	5.01	17.32

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Description:

FRAX 597 is a potent group I PAK inhibitor (IC₅₀ values are 8, 13 and 19 nM for PAK1, 2 and 3, respectively). FRAX 597 exhibits significant inhibition of YES1, RET, CSF1R and TEK at 100 nM, but is inactive against group II PAK isoforms (IC₅₀ >10 μM for PAK4). FRAX 597 inhibits proliferation of pancreatic cancer and schwannoma cells in vitro and exhibits antitumor effects in mice.

Physical and Chemical Properties:

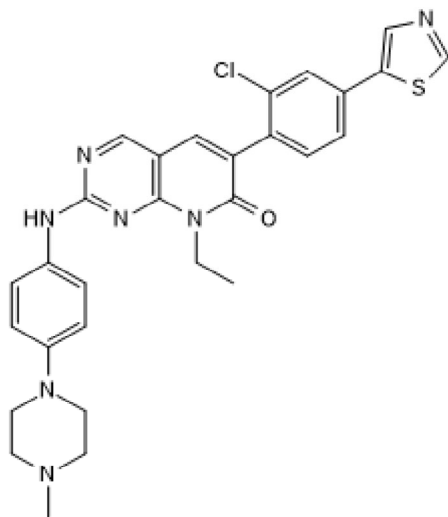
Batch Molecular Formula: C₂₉H₂₈ClN₇OS.

Batch Molecular Weight: 558.1

Physical Appearance: Yellow solid

Minimum Purity: ≥98%

Batch Molecular Structure:



Storage: Store at -20°C

Solubility & Usage Info:

DMSO to 20 mM

Stability and Solubility Advice:

Some solutions can be difficult to obtain and can be encouraged by rapid stirring, sonication or gentle warming (in a 45-60°C water bath).

Information concerning product stability, particularly in solution, has rarely been reported and in most cases we can only offer a general guide. *Unless contradicted by product-specific protocols or instructions, our standard recommendations apply:

SOLIDS: Provided storage is as stated on the product label and the vial is kept tightly sealed, the product can be stored for up to 6 months from date of receipt.

SOLUTIONS: We recommend that stock solutions, once prepared, are stored aliquoted in tightly sealed vials at -20°C or below and used within 1 month. Wherever possible solutions should be made up and used on the same day.

References:

Yeo et al (2016) FRAX597, a PAK1 inhibitor, synergistically reduces pancreatic cancer growth when combined with gemcit. *BMC Cancer* **16** 24. PMID: 26774265.

Licciulli et al (2013) FRAX597, a small molecule inhibitor of the p21-activated kinases, inhibits tumorigenesis of neurofibromatosis type 2 (NF2)-associated Schwannomas. *J.Biol.Chem.* **288** 29105. PMID: 23960073 .

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