

# Certificate of Analysis

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**Product Name:** Linifanib

**Catalog No.:** 7743

**Batch No.:** 1

CAS Number: 796967-16-3

IUPAC Name: *N*-[4-(3-Amino-1*H*-indazol-4-yl)phenyl]-*N'*-(2-fluoro-5-methylphenyl)urea

## 1. PHYSICAL AND CHEMICAL PROPERTIES

**Batch Molecular Formula:** C<sub>21</sub>H<sub>18</sub>FN<sub>5</sub>O.¼H<sub>2</sub>O

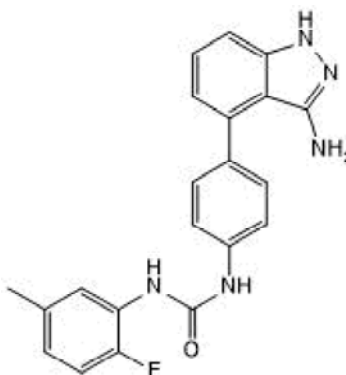
**Batch Molecular Weight:** 379.9

**Physical Appearance:** Off-white solid

**Solubility:** ethanol to 20 mM  
DMSO to 100 mM

**Storage:** Store at -20°C

**Batch Molecular Structure:**



## 2. ANALYTICAL DATA

**HPLC:** Shows 98.1% purity

**<sup>1</sup>H NMR:** Consistent with structure

**Mass Spectrum:** Consistent with structure

**Microanalysis:**

	Carbon	Hydrogen	Nitrogen
Theoretical	66.39	4.91	18.43
Found	66.65	4.86	18.48

Caution - Not Fully Tested • Research Use Only • Not For Human or Veterinary Use

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**Description:**

Linifanib is a potent receptor tyrosine kinase (RTK) inhibitor. It inhibits PDGFR $\beta$ , KDR, FLT3, CSF-1R (IC<sub>50</sub> values are 2, 4, 4 and 7 nM, respectively). Linifanib inhibits angiogenesis in vascularized micro-organs and potently inhibits VEGF-stimulated endothelial cell proliferation (IC<sub>50</sub> = 0.2 nM). Linifanib also promotes the generation and reprogramming of iPSCs from somatic cells.

**Physical and Chemical Properties:**

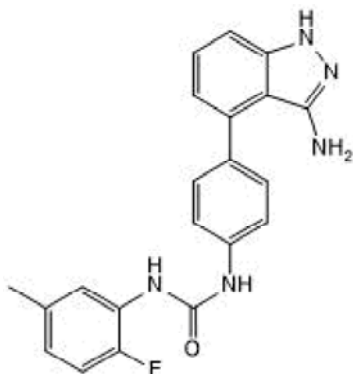
Batch Molecular Formula: C<sub>21</sub>H<sub>18</sub>FN<sub>5</sub>O.½H<sub>2</sub>O

Batch Molecular Weight: 379.9

Physical Appearance: Off-white solid

**Minimum Purity:** ≥98%

**Batch Molecular Structure:**



**Storage:** Store at -20°C

**Solubility & Usage Info:**

ethanol to 20 mM

DMSO to 100 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. Our standard recommendations are:

**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:**

**Guan *et al*** (2022) Chemical reprogramming of human somatic cells to pluripotent stem cells. *Nature* **605** 325. PMID: 35418683.

**Jahid *et al*** (2022) Structure-based design of CDC42 effector interaction inhibitors for the treatment of cancer. *Cell Rep.* **39** 110641. PMID: 35385746.

**Albert *et al*** (2006) Preclinical activity of ABT-869, a multitargeted receptor tyrosine kinase inhibitor. *Mol.Cancer Ther.* **5** 995. PMID: 16648571.

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