



LKT Laboratories, Inc.

NMS-1286937

Phone: 888-558-5227

651-644-8424

Fax: 888-558-7329

Email: getinfo@lktlabs.com

Web: lktlabs.com

Product Information

Product ID N5072

CAS No. 1034616-18-6

Chemical Name

Synonym NMS1286937, NMS-P937

Formula $C_{24}H_{27}F_3N_8O_3$

Formula Wt. 532.52

Melting Point

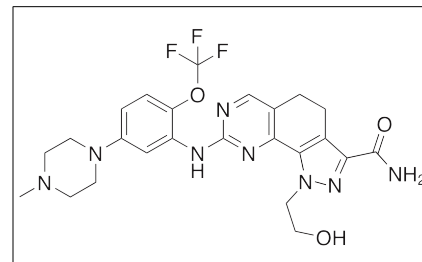
Purity $\geq 98\%$

Solubility DMSO 42 mg/mL warmed (78.87 mM)
Ethanol 10 mg/mL warmed (18.77 mM)
Water Insoluble

Store Temp -20°C

Ship Temp Ambient

Description NMS-1286937 is an inhibitor of polo-like kinase 1 (PLK1) that is currently in clinical trials as a potential treatment for various cancers; it exhibits anticancer chemotherapeutic activity. In osteosarcoma cells, NMS-1286937 induces cell cycle arrest and apoptosis and inhibits cell growth. In animal models of acute myelogenous leukemia (AML), this compound increases survival time and decreases tumor cell infiltration. Additionally, NMS-1286937 inhibits cell proliferation and induces apoptosis, cell cycle arrest, and tumor regression in cellular and animal models of colon cancer.



Pricing and Availability

Bulk quantities available upon request

Product ID	Size	List Price
N5072	1 mg	\$136.00
N5072	5 mg	\$217.20
N5072	10 mg	\$329.60

References Sero V, Tavanti E, Vella S, et al. Targeting polo-like kinase 1 by NMS-P937 in osteosarcoma cell lines inhibits tumor cell growth and partially overcomes drug resistance. *Invest New Drugs*. 2014 Dec;32(6):1167-80. PMID: 25193492.

Casolaro A, Golay J, Albanese C, et al. The Polo-Like Kinase 1 (PLK1) inhibitor NMS-P937 is effective in a new model of disseminated primary CD56+ acute monoblastic leukaemia. *PLoS One*. 2013;8(3):e58424. PMID: 23520509.

Valsasina B, Beria I, Alli C, et al. NMS-P937, an orally available, specific small-molecule polo-like kinase 1 inhibitor with antitumor activity in solid and hematologic malignancies. *Mol Cancer Ther*. 2012 Apr;11(4):1006-16. PMID: 22319201.

Beria I, Bossi RT, Brasca MG, et al. NMS-P937, a 4,5-dihydro-1H-pyrazolo[4,3-h]quinazoline derivative as potent and selective Polo-like kinase 1 inhibitor. *Bioorg Med Chem Lett*. 2011 May 15;21(10):2969-74. PMID: 21470862.

Caution: This product is intended for laboratory and research use only. It is not for human or drug use.