

PD173955

**SYN-1061** 

6-(2,6-dichlorophenyl)-8-methyl-2-(3-(methylthio)phenylamino)pyrido[2,3-d]pyrimidin-7(8H)-one

CAS Registry No.: 260415-63-2

Smiles String:

Cn1c2c(cc(c1=0)c3c(cccc3Cl)Cl)cnc(n2)Nc4ccc(c4)SC

Molecular Weight: 443.35

Molecular Formula: C21H16Cl2N4OS

Lot Number: Refer to vial

1H-NMR: Available on request

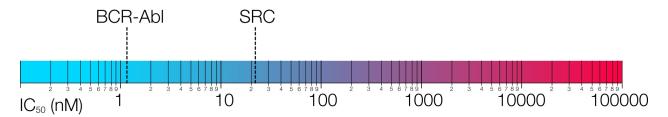
HPLC (Purity): > 95.0% @ 254 nm

ES-MS: Available on request

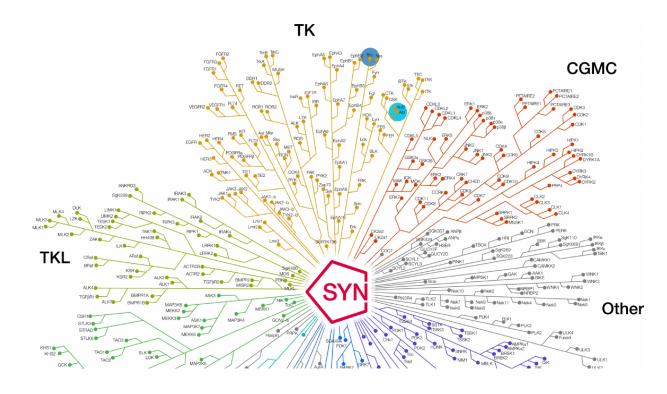
### **Description:**

PD173955 is an ATP-competitive, dual Src/Abl kinase inhibitor. It is a potent inhibitor of Bcr-Abl with an IC50 of 1-2 nM in kinase inhibition assays. In cellular growth assays it inhibits Bcr-Abl-dependent substrate tyrosine phosphorylation. PD173955 also inhibits Src kinase activity in vitro with an IC50 of 22 nM and could represent a novel class of anti-mitotic drug.

### **Biological Activity**



# Kinome Mapping



## **Shipping and Storage Temperature**

### Shipping:

Ambient

#### Storage:

2 years -20C, Powder 1 month, -4C in DMSO, More than one month -80C in DMSO

# Solubility

DMSO 112mg/mL, Ethanol 4mg/mL

# **Preparing Stock Solutions**

Stock Solution (1ml DMSO)	1mM	10mM	20mM	50mM
Mass(mg)	0.4433	4.4335	8.8670	22.1675

#### References

1. Nagar B, Bornmann WG, Pellicena P, Schindler T, Veach DR, Miller WT, Clarkson B, Kuriyan J. Crystal structures of the kinase domain of c-Abl in complex with the small molecule inhibitors PD173955 and imatinib (STI-571). Cancer Res. 2002 Aug 1;62(15):4236-43.

### **Ordering Information**

To order more of this or any other SYNkinase compound, go to synkinase.com, Call us Toll Free (US Only) at 1-877-854-6273 or email orders@synkinase.com.

Product Datasheet (Rev. 1.1)