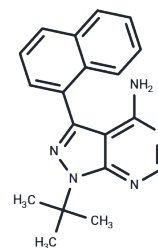


## 1-Naphthyl PP1

## Chemical Properties

|                   |   |
|-------------------|---|
| CAS No. :         | 221243-82-9   |
| Formula:          | C <sub>19</sub> H <sub>19</sub> N <sub>5</sub>  |
| Molecular Weight: | 317.39  |
| Storage:          | Powder: -20°C for 3 years   In solvent: -80°C for 1 year<br>Actual storage temperature shall be subject to the COA. |



## Biological Description

|               |  |
|---------------|--|
| Description   | 1-Naphthyl PP1 (1-NA-PP 1) is a selective Src inhibitor, targeting v-Src and c-Fyn, c-Abl, CDK2, and CAMK II with IC50 values of 1.0, 0.6, 0.6, 18, and 22 μM, respectively. |
| Targets(IC50) | Serine/threonin kinase,Src   |

## Solubility Information

|                     |  |
|---------------------|--|
| Solubility          | DMSO: 12 mg/mL (37.81 mM),Sonication is recommended.<br>(< 1 mg/ml refers to the product slightly soluble or insoluble)  |
| In vivo Formulation | 10% DMSO+40% PEG300+5% Tween 80+45% Saline: 1 mg/mL (3.15 mM),Sonication is recommended.<br><i>Please add the solvents sequentially, clarifying the solution as much as possible before adding the next one. Dissolve by heating and/or sonication if necessary. Working solution is recommended to be prepared and used immediately. The formulation provided above is for reference purposes only. In vivo formulations may vary and should be modified based on specific experimental conditions.</i> |

## Preparing Stock Solutions

|       | 1mg       | 5mg        | 10mg      |
|-------|-----------|------------|-----------|
| 1 mM  | 3.1507 mL | 15.7535 mL | 31.507 mL |
| 5 mM  | 0.6301 mL | 3.1507 mL  | 6.3014 mL |
| 10 mM | 0.3151 mL | 1.5753 mL  | 3.1507 mL |
| 50 mM | 0.063 mL  | 0.3151 mL  | 0.6301 mL |

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Tandon M, et al. New pyrazolopyrimidine inhibitors of protein kinase d as potent anticancer agents for prostate cancer cells. PLoS One. 2013 Sep 23;8(9):e75601.

Bishop AC, et al. A chemical switch for inhibitor-sensitive alleles of any protein kinase. Nature. 2000 Sep 21;407(6802):395-401.

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