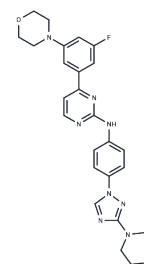


2-Pyridinamine, 4-[3-fluoro-5-(4-morpholinyl)phenyl]

Chemical Properties

| | |
|-------------------|--|
| CAS No. : | 1128096-64-9 |
| Formula: | C ₂₆ H ₂₇ FN ₈ O ₂ |
| Molecular Weight: | 502.54 |
| Storage: | Store at low temperature Powder: -20°C for 3 years In solvent: -80°C for 1 year <small>Actual storage temperature shall be subject to the COA.</small> |



Biological Description

| | |
|---------------|--|
| Description | 2-Pyridinamine, 4-[3-fluoro-5-(4-morpholinyl)phenyl] (Compound 9l) is an ATP-competitive JNK (c-Jun N-terminal kinase) inhibitor with IC ₅₀ values of 0.099 μM and 0.148 μM for JNK1 and JNK3 respectively, exhibiting the advantage of blood-brain barrier penetration. 2-Pyridinamine, 4-[3-fluoro-5-(4-morpholinyl)phenyl] inhibits c-Jun phosphorylation and reactive oxygen species (ROS) generation, making it suitable for neurodegenerative disease research. |
| Targets(IC50) | JNK,ROS |

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|-----------|------------|
| 1 mM | 1.9899 mL | 9.9495 mL | 19.8989 mL |
| 5 mM | 0.398 mL | 1.9899 mL | 3.9798 mL |
| 10 mM | 0.199 mL | 0.9949 mL | 1.9899 mL |
| 50 mM | 0.0398 mL | 0.199 mL | 0.398 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

Kamenecka T, et al. Synthesis, biological evaluation, X-ray structure, and pharmacokinetics of aminopyrimidine c-jun-N-terminal kinase (JNK) inhibitors. J Med Chem. 2010 Jan 14;53(1):419-31.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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