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|--------------------------|---------------------------------------------------------------------------------------------------------|
| <b>Product Name</b>      | : SU3327                                                                                                |
| <b>Synonyms</b>          | : —                                                                                                     |
| <b>Cat No.</b>           | : M24361                                                                                                |
| <b>CAS Number</b>        | : 40045-50-9                                                                                            |
| <b>Molecular Formula</b> | : C <sub>5</sub> H <sub>3</sub> N <sub>5</sub> O <sub>2</sub> S <sub>3</sub>                            |
| <b>Formula Weight</b>    | : 261.3                                                                                                 |
| <b>Chemical Name</b>     | : —                                                                                                     |
| <b>Description</b>       | : SU3327 is a potent, selective and substrate-competitive inhibitor of JNK(IC <sub>50</sub> of 0.7 μM). |
| <b>Pathway</b>           | : MAPK/ERK Signaling                                                                                    |
| <b>Target</b>            | : JNK                                                                                                   |
| <b>Receptor</b>          | : JNK;JNK-JIP interactions                                                                              |
| <b>Solubility</b>        | : DMSO:62.5 mg/mL (239.19 mM; Need ultrasonic)                                                          |
| <b>SMILES</b>            | : <chem>Nc1nnc(Sc2ncc([N+](O-)=O)s2)s1</chem>                                                           |
| <b>Storage</b>           | : (-20°C)                                                                                               |
| <b>Stability</b>         | : ≥ 2 years                                                                                             |
| <b>Reference</b>         | :                                                                                                       |

1.De SK, et al. Design, synthesis, and structure-activity relationship of substrate competitive, selective, and in vivo active triazole and thiadiazole inhibitors of the c-Jun N-terminal kinase. J Med Chem. 2009 Apr 9;52(7):1943-52.