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|--------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>Product Name</b>      | : Aurora Kinase Inhibitor III                                                                                                                                                    |
| <b>Synonyms</b>          | : —                                                                                                                                                                              |
| <b>Cat No.</b>           | : M20361                                                                                                                                                                         |
| <b>CAS Number</b>        | : 879127-16-9                                                                                                                                                                    |
| <b>Molecular Formula</b> | : C <sub>21</sub> H <sub>18</sub> F <sub>3</sub> N <sub>5</sub> O                                                                                                                |
| <b>Formula Weight</b>    | : 413.4                                                                                                                                                                          |
| <b>Chemical Name</b>     | : N-[3-[[4-[[3-(trifluoromethyl)phenyl]amino]-2-pyrimidinyl]amino]phenyl]-cyclopropanecarboxamide                                                                                |
| <b>Description</b>       | : Aurora kinase inhibitor III is a potent inhibitor of Aurora A kinase (IC <sub>50</sub> = 42 nM). It is selective for Aurora A over BMX/BTK, IGF-1R, c-Src, TRKB, SYK and EGFR. |
| <b>Pathway</b>           | : Cell Cycle/DNA Damage                                                                                                                                                          |
| <b>Target</b>            | : Aurora Kinase                                                                                                                                                                  |
| <b>Receptor</b>          | : Aurora A                                                                                                                                                                       |
| <b>Solubility</b>        | : DMSO: 50 mg/mL (120.9 mM)                                                                                                                                                      |
| <b>SMILES</b>            | : <chem>FC(F)(F)c1cccc(Nc2ccnc(Nc3cccc(NC(=O)C4CC4)c3)n2)c1</chem>                                                                                                               |
| <b>Storage</b>           | : (-20°C)                                                                                                                                                                        |
| <b>Stability</b>         | : ≥ 2 years                                                                                                                                                                      |
| <b>Reference</b>         | :                                                                                                                                                                                |

1. Zhang Q, Liu Y, Gao F et al. Discovery of EGFR Selective 4,6-Disubstituted Pyrimidines from a Combinatorial Kinase-Directed Heterocycle Library[J]. Journal of the American Chemical Society 2006 128(7):2182-2183.