

Product Name	: S49076
Synonyms	: S 49076; S-49076
Cat No.	: M11115
CAS Number	: 1265965-22-7
Molecular Formula	: C ₂₂ H ₂₂ N ₄ O ₄ S
Formula Weight	: 438.50
Chemical Name	: 2,4-Thiazolidinedione, 3-[[[2,3-dihydro-3-[[4-(4-morpholinylmethyl)-1H-pyrrol-2-yl]methylene]-2-oxo-1H-indol-5-yl]methyl]-
Description	: S49076 is a potent, ATP-competitive tyrosine kinase inhibitor of MET, AXL/MER, and FGFR1/2/3 with IC ₅₀ of <20 nM, also potently inhibits the kinase activity of mutated isoforms of MET (D1246N, Y1248D, Y1248H) and FGFR1/2; only inhibits 6% of kinases on a panel of 442 human wild-type and mutated kinases at 100 nM; inhibits the proliferation of MET- and FGFR2-dependent gastric cancer cells, blocks MET-driven migration of lung carcinoma cells, and inhibits colony formation of hepatocarcinoma cells expressing FGFR1/2 and AXL; causes tumor growth arrest in bevacizumab-resistant tumors in cancer xenograft models. Brain Cancer Phase 2 Clinical
Pathway	: Tyrosine Kinase
Target	: TAM Receptor
Receptor	: AXL; FGFR2; FGFR3; Mer; Met
Solubility	: DMSO: ≥ 31 mg/mL
SMILES	: <chem>O=C1NCC2=CC3=C(NC(=C3C=C4C(=CC(=CN4)C(=O)C=C2)SCC1=O)C=C5CCOCC5)</chem>
Storage	: (-20°C)
Stability	: ≥ 2 years
Reference	:

1. Burbridge MF, et al. Mol Cancer Ther. 2013 Sep;12(9):1749-62. | 2. Clémenson C, et al. Mol Cancer Ther. 2017 Oct;16(10):2107-2119. | 3. Rodon J, et al. Eur J Cancer. 2017 Aug;81:142-150.