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## JNJ-42041935

- 产品编号: D50942
  - CAS: 1193383-09-3
  - 分子式: C12H6ClF3N4O3
    - 纯度: ≥98%
  - InChi: InChI=1S/C12H6ClF3N4O3/c13-6-1-7-8(2-9(6)23-12(14,15)16)19-11(18-7)20-4-5(3-17-20)10(21) )22/h1-4H,(H,18,19)(H,21,22)
- InChi Key: FXHHASJVTYRJHH-UHFFFAOYSA-N

Smiles: OC(=O)C1C=NN(C=1)C1NC2C=C(OC(F)(F)F)C(Cl)=CC=2N=1

- 外观: 固体粉末
- 作用通路: HIF/HIF Prolyl-Hydroxylase
  - 溶解性: DMSO up to 100 mM
- 保存条件: Store in dry, dark place for one year.
- 产品介绍: JNJ-42041935 is a potent and selective HIF-PHD inhibitor. It is a 2-OG competitive and reversible inhibitor for PHD enzymes (pKis = 7.91, 7.29, and 7.65 for PHD1, 2, and 3, respectively). It is >100-fold selective for PHD compared to the related FIH (factor-inhibiting HIF) and a panel of various other enzymes. In an inflammation-induced anemia model in rats, 100 µM/kg/day JNJ-42041935 significantly increased the number of circulating reticulocytes and red blood cells, increased blood hemoglobin and hematocrit, and restored mean corpuscular volume and mean cell hemoglobin of red bloods cells. JNJ-42041935 is a new pharmacological tool, which can be used to investigate PHD inhibition and demonstrate that PHD inhibitors offer great promise for the treatment of inflammation-induced anemia.