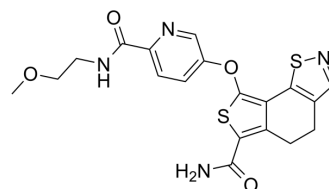


CDK8/19-IN-1

| | | | |
|---------------------------|--|-------|----------|
| Cat. No.: | HY-111427 | | |
| CAS No.: | 1818427-07-4 | | |
| Molecular Formula: | C ₁₉ H ₁₈ N ₄ O ₄ S ₂ | | |
| Molecular Weight: | 430.5 | | |
| Target: | CDK | | |
| Pathway: | Cell Cycle/DNA Damage | | |
| Storage: | Powder | -20°C | 3 years |
| | | 4°C | 2 years |
| | In solvent | -80°C | 6 months |
| | | -20°C | 1 month |



SOLVENT & SOLUBILITY

In Vitro

DMSO : 50 mg/mL (116.14 mM; Need ultrasonic)

| Concentration | Mass | | | |
|---------------|-----------|------------|------------|--|
| | 1 mg | 5 mg | 10 mg | |
| 1 mM | 2.3229 mL | 11.6144 mL | 23.2288 mL | |
| 5 mM | 0.4646 mL | 2.3229 mL | 4.6458 mL | |
| 10 mM | 0.2323 mL | 1.1614 mL | 2.3229 mL | |

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description

CDK8/19-IN-1 is a potent, selective and oral bioavailable CDK8/19 dual inhibitor, with IC₅₀s of 0.46 nM, 0.99 nM and 270 nM for CDK8, CDK19 and CDK9, respectively.

IC₅₀ & Target

| | | |
|--|---|------------------------------------|
| CDK8/CycC 0.46 nM (IC ₅₀) | CDK19/CycC 0.99 nM (IC ₅₀) | CDK9 270 nM (IC ₅₀) |
|--|---|------------------------------------|

In Vitro

CDK8/19-IN-1 (52h) is a potent CDK8/19 dual inhibitor, with IC₅₀s of 0.46 nM, 0.99 nM and 270 nM for CDK8, CDK19 and CDK9, respectively. CDK8/19-IN-1 also weakly inhibits CDK2, with 62% inhibition at 1 μM. CDK8/19-IN-1 (1 μM) shows >50% inhibition against GSK3β, PLK1, ASK1, CK1δ, PKA, ROCK1, PKCθ, CDC7. CDK8/19-IN-1 shows K_ds of 25, 46, 81, 86, 97, 160 and >3000 nM for CDK19, CDK8, DYRK1B, HASPIN, YSK4, HIPK1 and EPHA3, respectively. CDK8/19-IN-1 displays potent antitumor activity, with GI₅₀ of 0.43-2.5 nM for colon, multiple myeloma, acute myelogenous leukemia (AML), lung cancer cells^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

CDK8/19-IN-1 (52h; 1.25 mg/kg twice daily or 2.5 mg/kg once daily, p.o.) significantly suppresses tumor growth in mice bearing RPMI8226 human hematopoietic and lymphoid cells^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Ono K, et al. Design and synthesis of selective CDK8/19 dual inhibitors: Discovery of 4,5-dihydrothieno[3',4':3,4]benzo[1,2-d]isothiazole derivatives. *Bioorg Med Chem*. 2017 Apr 15;25(8):2336-2350.

Caution: Product has not been fully validated for medical applications. For research use only.

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