

Product Datasheet

DS-9300

Catalog No. **BP-02372** · CAS **2259641-46-6** · Form **free base** · Physical state **other** · Color **White to off-white** · Version **1.0** · Revision **2026-09-11**

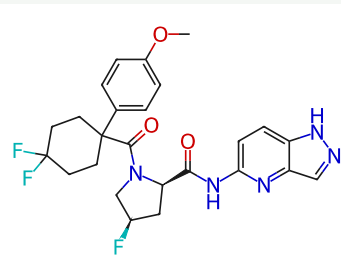
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Target Histone Acetyltransferase	Research area Cancer
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Molecular formula **C₂₅H₂₆F₃N₅O₃**

Molecular weight **501.5008**

STRUCTURE



DESCRIPTION

DS-9300 is a potent, orally active, selective EP300/CBP HAT inhibitor with an IC₅₀ value of 28 nM. DS-9300 has anticancer activity and can be used in prostate cancer disease research .

BIOLOGICAL ACTIVITY

In Vitro

DS-9300 inhibits the growth of prostate cancer cell lines with the IC₅₀ values of 0.6, 6.5, 3.4 and 287.2 nM for VCaP, 22Rv1, LNCaP and PC3 cells, respectively, indicating that it could effectively inhibit the growth of prostate cancer cell lines by inhibiting histone acetylation and down-regulating PSA expression

In Vivo

DS-9300 (p.o., 0.3-3 mg/kg, once daily, 33days) shows dose-dependent antitumor activity with 39%, 74% and 109% tumor growth inhibition at 0.3, 1 and 3 mg/kg, respectively, and no significant toxic effects in Castrated VCaP xenograft mouse model

SOLUBILITY

Soluble in DMSO

In vitro DMSO : 100 mg/mL

REFERENCES

[1].Ryutaro Kanada, et al. Discovery of DS-9300: A Highly Potent, Selective, and Once-Daily Oral EP300/CBP Histone Acetyltransferase Inhibitor. J Med Chem. 2023 Jan 12;66(1):695-715.