

Product Datasheet

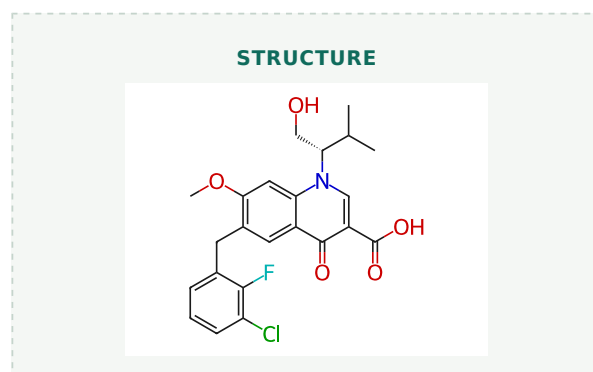
Elvitegravir

Catalog No. **BP-02494** · CAS **697761-98-1** · Form **free base** · Physical state **other** · Color **White to off-white** · Version **1.0** · Revision **2026-09-10**

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Target HIV IntegraseHIV	Research area Infection
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Molecular formula	C₂₃H₂₃ClFNO₅
Molecular weight	447.8840



DESCRIPTION

Elvitegravir (GS-9137; JTK-303; D06677) is an HIV integrase inhibitor for HIV-1IIB, HIV-2EHO and HIV-2ROD with IC50 of 0.7 nM, 2.8 nM and 1.4 nM, respectively.

BIOLOGICAL ACTIVITY

Elvitegravir also inhibits PBMC and PA with IC50 of 0.89 and 20 nM, respectively. Elvitegravir (also known as JTK-303/GS-9137) blocked the integration of HIV-1 cDNA through the inhibition of DNA strand transfer. EVG inhibited the replication of HIV-1, including various subtypes and multiple-drug-resistant clinical isolates, and HIV-2 strains with a 50% effective concentration in the subnanomolar to nanomolar range. The replication capacity of EVG-resistant variants was significantly reduced relative to both wild-type virus and other IN inhibitor-resistant variants selected by L-870810. Elvitegravir (GS-9137) is active against HIV-1 and HIV-2 and has a serum-free antiviral IC50 of 0.3-0.9 nM in peripheral blood mononuclear cells. According to the results of the phase II clinical trial, patients taking once-daily elvitegravir boosted by ritonavir had greater reductions in viral load after 24 weeks compared to individuals randomized to receive a ritonavir-boosted protease inhibitor. Elvitegravir is currently in a Phase III clinical trial in the treatment of HIV-1 Infection.

SOLUBILITY

Soluble in DMSO

REFERENCES

[1]. Shimura K, et al. Broad antiretroviral activity and resistance profile of the novel human immunodeficiency virus integrase inhibitor elvitegravir (JTK-303/GS-9137). J Virol. 2008 Jan;82(2):764-74.