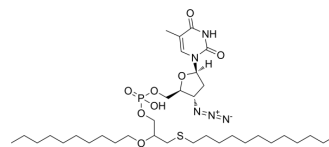


Fozivudine tidoxil

Cat. No.:	HY-126781
CAS No.:	141790-23-0
Molecular Formula:	C ₃₅ H ₆₄ N ₅ O ₈ PS
Molecular Weight:	745.95
Target:	HIV; DNA/RNA Synthesis
Pathway:	Anti-infection; Cell Cycle/DNA Damage
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Fozivudine tidoxil (BM-211290) is an orally active thioether lipid-zidovudine (ZDV) conjugate with anti-HIV activity. Fozivudine tidoxil, a member of the NRTI family of agent, is incorporated into the newly synthesized strand of DNA during intracellular viral replication and irreversibly binds viral RT which disrupts viral reverse-transcription ^{[1][2]} . Fozivudine tidoxil is a click chemistry reagent, it contains an Azide group and can undergo copper-catalyzed azide-alkyne cycloaddition reaction (CuAAC) with molecules containing Alkyne groups. Strain-promoted alkyne-azide cycloaddition (SPAAC) can also occur with molecules containing DBCO or BCN groups.								
IC₅₀ & Target	HIV								
In Vitro	Fozivudine tidoxil (BM-211290) is a member of the nucleoside analogue reverse transcriptase inhibitor (NRTI) family ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.								
In Vivo	<p>Fozivudine tidoxil (BM-211290; 45 mg/kg; PO; twice daily; one day before FIV challenge for a total of six weeks) is effective at lowering plasma- and cell-associated viremia at two weeks post-FIV infection^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 30%;">Animal Model:</td> <td>Specific pathogen-free cats at 6 months of age^[1]</td> </tr> <tr> <td>Dosage:</td> <td>45 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>PO; twice daily; one day before FIV challenge for a total of six weeks</td> </tr> <tr> <td>Result:</td> <td>Effective at lowering plasma- and cell-associated viremia at two weeks post-feline immunodeficiency virus (FIV) infection with a trend toward lower plasma- and cell-associated viremia at four and six weeks post-infection (PI).</td> </tr> </table>	Animal Model:	Specific pathogen-free cats at 6 months of age ^[1]	Dosage:	45 mg/kg	Administration:	PO; twice daily; one day before FIV challenge for a total of six weeks	Result:	Effective at lowering plasma- and cell-associated viremia at two weeks post-feline immunodeficiency virus (FIV) infection with a trend toward lower plasma- and cell-associated viremia at four and six weeks post-infection (PI).
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REFERENCES

[1]. Michelle M Miller, et al. Administration of Fozivudine Tidoxil as a Single-Agent Therapeutic During Acute Feline Immunodeficiency Virus Infection Does Not Alter Chronic Infection. *Viruses*. 2012 Jun;4(6):954-62.

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

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