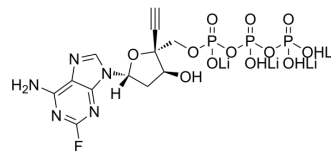


EFdA-TP tetralithium

Cat. No.:	HY-138561C
Molecular Formula:	C ₁₂ H ₁₄ FLi ₄ N ₅ O ₁₂ P ₃
Molecular Weight:	559.95
Target:	HIV; Reverse Transcriptase
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	EFdA-TP tetralithium is a potent nucleoside reverse transcriptase (RT) inhibitor. EFdA-TP tetralithium inhibits RT-catalyzed DNA synthesis as an effective immediate or delayed chain terminator (ICT or DCT). EFdA-TP tetralithium inhibits HIV-1 RT with multiple mechanisms ^[1] .
In Vitro	EFdA-TP (0.05-10 μM; for 15 min) tetralithium inhibits RT-catalyzed DNA synthesis as an ICT or DCT ^[1] . EFdA-TP tetralithium can block RT as a translocation-defective RT inhibitor that dramatically slows DNA synthesis, acting as a de facto immediate chain terminator ^[1] . EFdA-TP tetralithium can function as a delayed chain terminator, allowing incorporation of an additional dNTP before blocking DNA synthesis ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Eleftherios Michailidis, et al. 4'-Ethynyl-2-fluoro-2'-deoxyadenosine (EFdA) inhibits HIV-1 reverse transcriptase with multiple mechanisms. J Biol Chem. 2014 Aug 29;289(35):24533-48.

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

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