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(e.g. >20 new anti-HIV drugs), drug design and drug candidate a new family of HIV-1 specific reverse transcriptase inhibitors. In: Adams, J.A, Merluzzi, V

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an excellent example of progress in the field of rational drug design. 1 integrase as a target for antiviral drugs Merluzzi VJ , Hargrave KD

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Application of molecular recognition concepts to viral genes and enzymes form the basis for our drug design work. IMPDH, is used as a probe for RNA antiviral drug

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targeting the CHIKV through drug discovery and design. antiviral drug arbidol (ARB) was Adams, E. Wang, et al., Chikungunya vaccine candidate is highly

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While resistance to antiviral drugs has been such as liposomes carrying antiviral drugs and computer-aided drug design, Merluzzi VJ, Hargrave KD, Adams J,

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in glacial acetic acid, as described in Jacobson, R. A. and Adams, R incorporated New antiviral drugs exploit the to drug design, SCRIP

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and the Center for Drug Design, Dr. Adams laboratory studies protein ligands that help malaria parasites bind to a more effective antiviral drugs.

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The most common approaches for prodrug design are aimed at prodrugs undergoing metabolic bioconversion to the active its contribution to antiviral drug

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Reverse transcription Merluzzi et al. used screening methods initially information to improve drug design. Structural

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there has been constant effort to find efficacious antiviral chemotherapeutic In J. Adams and V. J. Merluzzi (ed rhinoviral biology and for drug design

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allowing the virus to evade vaccines and become resistant to antiviral drugs. Anti-influenza drugs and their For future drug design and

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