



This occasional section within the journal surveys visions and achievements, often not on the main track of the developing biomedical sciences, but all relating to discoveries and developments of medicinal – both ancient and modern. What they have in common, in one way or another, is providing further background and glances around the edges of the core discipline of pharmacognosy, as it has been and continues to evolve within our times.

Croton olifandrus Pierre and Hutch Compounds Reverse HIV-1 Viral Latency

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Human Immunodeficiency Virus (HIV) infections are nowadays managed effectively using modern antiretroviral therapies (ART). Notably, these therapies do not cure the infection, and the virus may remain latent in reservoirs throughout the body. The latent virus can reactivate at any time, and disease progression may proceed. ART treatment must be ongoing, generally for the lifetime of the patient. However, even with correct treatment, the virus may still reactivate. Additionally, the latent virus in these reservoirs may cause chronic inflammation, as well as other comorbidities in some patients. Therefore, there is still no effective and readily accessible cure for HIV, and much research is still required in this field. Medicinal plants have potential for the treatment of viral diseases. Recently, substantial research has focussed on respiratory viruses due to covid-19 global

pandemic,¹⁻⁵ with hundreds of African plants identified to be effective against multiple viral respiratory infections.^{6,7}

A recent study by Cameroon researchers published in the Journal of Experimental Pharmacology investigated compounds isolated from *Croton olifandrus* Pierre and Hutch as potential latency-reversing drugs (LRD), which would allow the virus to be actively targeted and eliminated.⁸ Four of the isolated compounds displayed substantial reversal of viral latency. In particular, the authors highlighted β -stigmaterol and lupeol as particularly promising. Furthermore, some of the isolated compounds synergised the latency reversal effects of other existing latency reversing compounds when administered as combinations. The authors also reported that the compounds reversed latency by mechanisms distinct from the current protein kinase C activation and histone deacetylase inhibition target mechanism. Whilst this research is promising, the study examined the effects *in vitro*, and other issues such as bioavailability and toxicity need to be evaluated before these compounds can be considered for therapeutic use.



DOI: 10.5530/pc.2025.1.6

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CONFLICT OF INTEREST

The author declares that there is no conflict of interest.

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