

Scientists show potent new compound virtually eliminates HIV in cell culture

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A new study by scientists on the Florida campus of The Scripps Research Institute shows, in cell culture, a natural compound can virtually eliminate human immunodeficiency virus (HIV) in infected cells. The compound defines a novel class of HIV anti-viral drugs endowed with the capacity to repress viral replication in acutely and chronically infected cells.

The HIV/AIDS pandemic continues to affect 34 million individuals worldwide, including more than 3 million children, according to the <u>World Health Organization</u>. Current treatment involves the use of several antiretroviral drugs, termed Highly <u>Active Antiretroviral Therapy</u> (HAART), which can extend the life expectancy of HIV-positive individuals and decrease viral load without, however, eradicating the virus.

"We know that there are reservoirs of HIV that aren't being eliminated by current treatment and that keep replenishing the infection," said Susana Valente, a Scripps Research biologist who led the study. "Viral production from these cellular reservoirs that harbor an integrated <u>viral</u> <u>genome</u> is not affected by current <u>antiretroviral drugs</u>, which only stop novel rounds of infection. The compound in the current study virtually eliminates all viral replication from already-infected cells where HIV hides."

The new study, published in the July 20, 2012 issue of the journal <u>Cell</u> <u>Host and Microbe</u>, focused on a medically promising compound known



as Cortistatin A. This natural product was isolated in 2006 from a marine sponge, Corticium simplex, discovered more than 100 years ago. In 2008, Scripps Research chemist Phil Baran and his team won the global race to synthesize the compound, presenting an efficient and economical method.

In the new study, Valente and her colleagues collaborated with the Baran lab, using a synthetic version of the compound, didehydro-Cortistatin A, to study the compound's effect on two strains of HIV. The strains were HIV-1, the most common form of the virus, and HIV-2, which is concentrated in West Africa and some parts of Europe.

The results showed that the compound reduced viral production by 99.7 percent from primary CD4+T cells (a type of immune cell) isolated from patients without levels of the virus in their bloodstream and who had been under HAART treatment for a long period of time. When the compound was added to other antiviral treatments, it further reduced by 20 percent viral replication from CD4+T cells isolated from patients with detectable amounts of virus in their bloodstreams.

The inhibitor works by binding tightly to the viral protein known as Tat, a potent activator of HIV gene expression, effectively preventing the virus from replicating even at miniscule concentrations—making it the most potent anti-Tat inhibitor described to date, Valente said. Another interesting feature of this compound is that withdrawal of the drug from cell culture does not result in virus rebound, which is normally observed with other antiretrovirals.

While most antiretroviral <u>compounds</u> block only new infections, didehydro-Cortistatin A reduces <u>viral replication</u> from already-infected cells, potentially limiting cell-to-cell transmission.

The new inhibitor already has a drug-like structure, is effective at very



low concentrations, and has no toxicity associated with it, at least at the cellular level, the study noted.

More information: "Potent Suppression of Tat-dependent HIV Transcription by didehydro-Cortistatin A" *Cell Host and Microbe*.

Provided by Scripps Research Institute

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