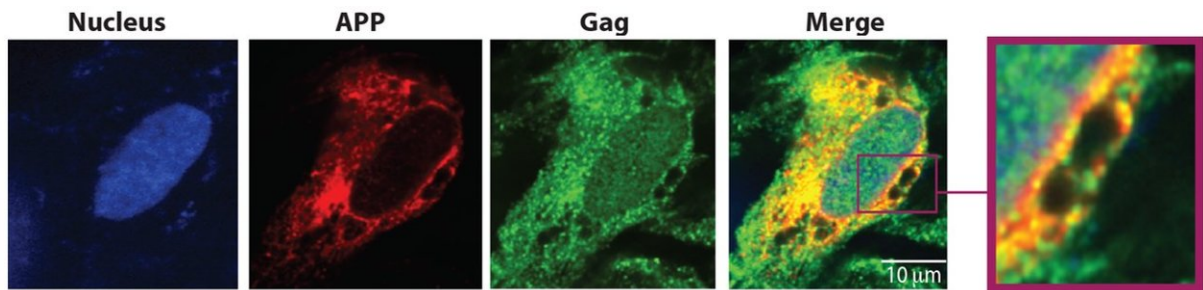


Molecular mechanism behind HIV-associated dementia revealed

January 5 2018, by Will Doss



The interaction between APP (red) and GAG (green) in the cell. Credit: Northwestern University

For the first time, scientists have identified and inhibited a molecular process that can lead to neurodegeneration in patients with HIV, according to a Northwestern Medicine study published in *Nature Communications*.

Mojgan Naghavi, Ph.D., associate professor of Microbiology-Immunology, was senior author on the study, and Qingqing Chai, Ph.D., a postdoctoral fellow in Naghavi's lab, was the lead author.

Previous studies have found elevated levels of toxic beta-amyloid protein in the brains of patients with human immunodeficiency virus (HIV), causing HIV-associated neurocognitive disorder (HAND). While

[combination antiretroviral therapy](#) can greatly slow down HIV progression, up to 50 percent of these patients exhibit milder forms of HAND. Buildup of beta-amyloid is thought to be a major contributor to neurodegeneration in a variety of dementia-associated diseases, especially Alzheimer's, but how and why beta-amyloid is produced in HIV-infected patients and its contribution to HAND was a mystery, Naghavi said.

"The HIV virus cannot infect neurons because they don't have the right receptors," said Naghavi, who is also a member of the Robert H. Lurie Comprehensive Cancer Center of Northwestern University. "We've known about elevated beta-amyloid levels for a long time, but nobody knew how or why the protein was overproduced in response to infection."

The first clue came when Naghavi and her colleagues performed a large-scale assay looking for unusual interactions between cellular proteins and Gag, an important HIV protein. A number of proteins were shown to bind with Gag, but one stood out—membrane-associated amyloid precursor protein (APP), a protein that can be processed to produce the toxic beta-amyloid protein.

"We put two and two together: We knew APP is a precursor to beta-amyloid, so we looked for APP and found increased processing of this protein into beta-amyloid in infected macrophages and microglia," Naghavi said.

Macrophages and microglia are two types of non-neuronal brain cells that HIV infect, often functioning as reservoirs of the disease in the brain, according to Naghavi. In those cells, APP interferes with HIV infection, and in response, HIV tries to circumvent that barrier.

During infection of a microglia or macrophage cell, HIV's Gag [protein](#)

promotes processing of APP, reducing resistance to cell takeover, which also has the side-effect of producing toxic beta-amyloid proteins, Naghavi said.

"By binding to APP, somehow Gag drags it into cell membrane regions called lipid rafts, where there are enzymes that promote processing of APP into beta-amyloid," she said. "This is the first time we've showed HIV is trying to overcome a block that is induced by APP in microglia and macrophages."

To reduce buildup of beta-amyloid, the scientists targeted the enzyme that cleaves APP, called gamma secretase. A simple inhibitor that blocked APP processing produced dual benefits by inhibiting infection and reducing beta-amyloid production, according to Naghavi.

When the drug blocks APP processing, it reduces beta amyloid and increases APP compared to non-drugged models. This results in the combined benefits of reduced neurodegeneration and sustaining APP's original function of blocking infection.

"We've opened the door for how these drugs might be used to block both HIV replication and treat HIV-associated dementia," Naghavi said. "It would also be interesting to investigate whether APP affects replication of other neurotropic viruses, for example Zika or Cytomegalovirus."

However, translating these findings into a drug that works in patients is still a ways off. Compounds that have the intended effect may also be toxic in humans or have other undesirable side effects, so finding non-toxic drugs that can modulate gamma-secretase is the next step towards an actionable treatment.

HIV patients are living longer thanks to more effective treatments, so there's a growing need to address the HIV-specific chronic health issues

they encounter as they age, said Naghavi.

"HIV patients on combination antiretroviral therapy have close to a normal life, but they are still suffering from diseases like HAND," Naghavi said. "This is just the beginning."

More information: Qingqing Chai et al. HIV-1 counteracts an innate restriction by amyloid precursor protein resulting in neurodegeneration, *Nature Communications* (2017). [DOI: 10.1038/s41467-017-01795-8](https://doi.org/10.1038/s41467-017-01795-8)

Provided by Northwestern University

Citation: Molecular mechanism behind HIV-associated dementia revealed (2018, January 5) retrieved 27 April 2024 from <https://medicalxpress.com/news/2018-01-molecular-mechanism-hiv-associated-dementia-revealed.html>

<p>This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.</p>
--