

Scientists uncover secret behind molecule that blocks HIV infection

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A simulation shows how the proteins assemble themselves gradually into a net around the invading HIV capsid. Credit: Alvin Yu



Rhesus macaques don't monkey around when it comes to HIV; they have a protein that effectively disables invading HIV particles.

A group of University of Chicago scientists announced an innovative study that explains how the macaques' immune protein, called TRIM5 α , works its magic. It also represents a significant step forward in the science of modeling how complex biological proteins assemble themselves, the scientists said.

"These proteins work together to encase the HIV capsid in a hexagonal net and restrict viral activity," said postdoctoral fellow Alvin Yu and lead author of the study, which was published in *Nature Communications*.

It turns out that part of the secret appears to be defects—irregularities in the <u>lattice</u> net that allow the TRIM5 α proteins the flexibility to encase any shape the invading <u>virus</u> takes.

Yu and Prof. Gregory Voth have been working to understand the physics underlying <u>biological processes</u> using computer simulations. But a <u>simulation</u> that actually represented every atom in an HIV virus can easily overwhelm even the largest supercomputers, so there's a delicate methodology to picking which parts are vital to include in a simulation, and which can be safely pixelated out. This is called "coarse-graining."





Rhesus monkeys have proteins that link up to encase invading HIV particles and disable them. UChicago researchers showed how the proteins assemble into a hexagonal net that can target the rapidly-evolving virus, even if it takes different shapes. Credit: Alvin Yu

They turned their attention to the mystery of the TRIM5 α protein. Yu and Voth knew that TRIM5 α could form two-dimensional lattices, but questions remained as to how the proteins collectively wrap around the three-dimensional capsid. So they ran simulations and modeled how the proteins interacted with each other as well as the invading virus, based on what is known from other experiments about its makeup.

The new models suggested several key points. One was that the protein uses an elaborate "hopping" mechanism to gradually accumulate atop the capsid until it reaches a tipping point. Then, as the lattice grows, the proteins squeeze together in such a way that irregularities start to appear



in the lattice.

Careful experiments by their collaborators verified what they had seen in the model. Yu surmised that these irregularities are important so the monkey TRIM5 α can adjust to differently shaped HIV capsules. "HIV capsids are notable for varying significantly in their structures—so these TRIM proteins also need the ability to adapt to their different structures," Yu said.

It's also important how strongly the proteins bind to each other and the virus. "There's also a fine balance in interactions between the <u>protein</u> and virus. The assembly of this lattice is a collective behavior that only occurs over very particular ranges of interaction strengths," Yu explained.

Such knowledge may someday help inform treatments for HIV.

There are also implications for computing, Voth said: "This is a big step forward in technology for <u>computer simulations</u> of very large molecular systems. It really effectively represents more than a billion atoms, which you could never do with a study at full atomic resolution. It's a revolution in that sense."

More information: Alvin Yu et al. TRIM5α self-assembly and compartmentalization of the HIV-1 viral capsid, *Nature Communications* (2020). DOI: 10.1038/s41467-020-15106-1

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