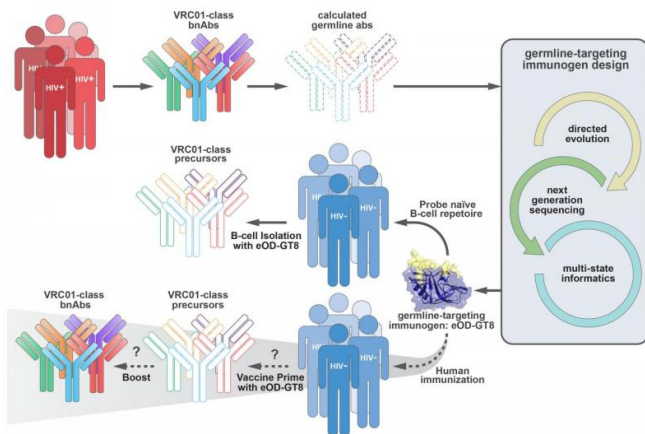


Findings in humans provide encouraging foundation for upcoming AIDS vaccine clinical trial

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An immunogen named eOD-GT8 was designed to bind and activate the predicted human germline precursor B cells of the VRC01 class of HIV broadly neutralizing antibodies (bnAbs). As a test, eOD-GT8 was used as a probe to isolate human naive B cells from HIV-uninfected individuals. This process pulled out VRC01-class human precursor B cells, confirming the design and suggesting that eOD-GT8 immunogens should be further tested in a human vaccine setting to determine if eOD-GT8 can activate the targeted VRC01-class B cells as a first step toward the ultimate elicitation of VRC01-class bnAbs that could protect people against HIV. Credit: J.G. Jardine et al. / Science (2016)

Some people infected with HIV naturally produce antibodies that effectively neutralize many strains of the rapidly mutating virus, and scientists are working to develop a vaccine capable of inducing such "broadly neutralizing" antibodies that can prevent HIV infection.

An emerging [vaccine](#) strategy involves immunizing people with a series of different engineered HIV proteins as immunogens to teach the immune system to produce broadly [neutralizing antibodies](#)

against HIV. This strategy depends on the ability of the first immunogen to bind and activate special cells, known as broadly neutralizing antibody precursor B cells, which have the potential to develop into broadly neutralizing antibody-producing B cells.

A research team has now found that the right precursor ("germline") cells for one kind of HIV broadly neutralizing antibody are present in most people, and has described the design of an HIV vaccine germline-targeting immunogen capable of binding those B cells. The findings by scientists from The Scripps Research Institute (TSRI), the International AIDS Vaccine Initiative (IAVI) and the La Jolla Institute for Allergy and Immunology were published in *Science* on March 25.

"We found that almost everybody has these broadly neutralizing antibody precursors, and that a precisely engineered protein can bind to these cells that have potential to develop into HIV broadly neutralizing antibody-producing cells, even in the presence of competition from other [immune cells](#)," said the study's lead author, William Schief, TSRI professor and director, Vaccine Design of the IAVI Neutralizing Antibody Center at TSRI, in whose lab the engineered HIV vaccine protein was developed.

The body's immune system contains a large pool of different precursor B cells so it can respond to a wide variety of pathogens. But that also means that precursor B cells able to recognize a specific feature on a virus surface are exceedingly rare within the total pool of B cells.

"The challenge for vaccine developers is to determine if an immunogen can present a particular viral surface in a way that distinct B cells can be activated, proliferate and be useful," said study co-author Shane Crotty, professor at the La Jolla

Institute. "Using a new technique, we were able to show—well in advance of clinical trials—that most humans actually have the right B cells that will bind to this vaccine candidate. It is remarkable that protein design can be so specific as to 'find' one in a million cells, demonstrating the feasibility of this new vaccine strategy."

The work offers encouraging insights for a planned Phase 1 clinical trial to test a nanoparticle version of the engineered HIV vaccine protein, the "eOD-GT8 60mer." "The goal of the clinical study will be to test safety and the ability of this engineered protein to elicit the desired immune response in humans that would look like the start of broadly neutralizing antibody development," Schief said. "Data from this new study was also important for designing the clinical trial, including the size and the methods of analysis."

In June, scientists from TSRI, IAVI and The Rockefeller University reported that the eOD-GT8 60mer produced antibody responses in mice that showed some of the traits necessary to recognize and inhibit HIV. If the eOD-GT8 60mer performs similarly in humans, additional boost immunogens are thought to be needed to ultimately induce broadly neutralizing antibodies that can block HIV.

The new work also provides a method for researchers to assess whether other new vaccine proteins can bind their intended precursor B cells. This method is a valuable tool in the design of more targeted and effective vaccines against AIDS, providing the ability to vet germline-targeting immunogens before testing them in large, time-consuming and costly clinical trials.

Looking at blood donated by healthy volunteers, the scientists found B cells that were capable of creating "VRC01-class" antibodies that recognized a critical surface patch, or epitope, of HIV. VRC01-class [broadly neutralizing antibodies](#) are a group of antibodies isolated from different individuals that appear to have developed in a very similar way, and it has been hypothesized that the starting VRC01-class B cells were very similar in the different people. The eOD-GT8 60mer is designed to engage these precursor B [cells](#) to initiate HIV broadly neutralizing antibody

development.

More information: "HIV-1 broadly neutralizing antibody precursor B cells revealed by germline-targeting immunogen," *Science*, [DOI: 10.1126/science.aad9195](https://doi.org/10.1126/science.aad9195)

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