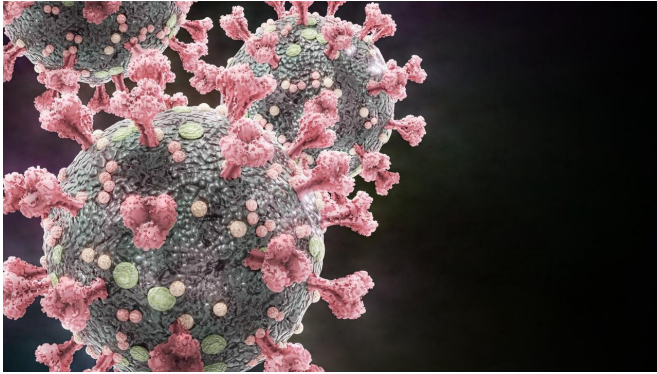


COVID-19: Scientists identify human genes that fight infection

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Microscopic view of coronavirus. Credit: Dotted Yeti

Scientists at Sanford Burnham Prebys have identified a set of human genes that fight SARS-CoV-2 infection, the virus that causes COVID-19. Knowing which genes help control viral infection can greatly assist researchers' understanding of factors that affect disease severity and also suggest possible therapeutic options. The genes in question are related to interferons, the body's frontline virus fighters.

The study was published in the journal *Molecular Cell*.

"We wanted to gain a better understanding of the cellular response to SARS-CoV-2, including what drives a strong or weak response to infection," says Sumit K. Chanda, Ph.D., professor and director of the Immunity and Pathogenesis Program at Sanford Burnham Prebys and lead author of the study. "We've gained new insights into how the virus exploits the [human cells](#) it invades, but we are still searching for its Achilles' heel so that we can develop optimal antivirals."

Soon after the start of the pandemic, clinicians found that a weak interferon response to SARS-

CoV-2 infection resulted in some of the more severe cases of COVID-19. This knowledge led Chanda and his collaborators to search for the [human genes](#) that are triggered by interferons, known as interferon-stimulated genes (ISGs), which act to limit SARS-CoV-2 infection.

Based on knowledge gleaned from SARS-CoV-1, the virus that caused a deadly, but relatively brief, outbreak of disease from 2002 to 2004, and knowing that it was similar to SARS-CoV-2, the investigators were able to develop laboratory experiments to identify the ISGs that control [viral replication](#) in COVID-19.

"We found that 65 ISGs controlled SARS-CoV-2 infection, including some that inhibited the virus' ability to enter cells, some that suppressed manufacture of the RNA that is the virus's lifeblood, and a cluster of genes that inhibited assembly of the virus," says Chanda. "What was also of great interest was the fact that some of the ISGs exhibited control across unrelated viruses, such as seasonal flu, West Nile and HIV, which leads to AIDS."

"We identified eight ISGs that inhibited both SARS-CoV-1 and CoV-2 replication in the subcellular compartment responsible for protein packaging, suggesting this vulnerable site could be exploited to clear viral [infection](#)," says Laura Martin-Sancho, Ph.D., a senior postdoctoral associate in the Chanda lab and first author of this study. "This is important information, but we still need to learn more about the biology of the virus and investigate if genetic variability within these ISGs correlates with COVID-19 severity."

As a next step, the researchers will look at the biology of SARS-CoV-2 variants that continue to evolve and threaten vaccine efficacy. Martin-Sancho notes that they have already started gathering variants for laboratory investigation,

"It's vitally important that we don't take our foot off the pedal of basic research efforts now that vaccines are helping control the pandemic," concludes Chanda. "We've come so far so fast because of investment in [fundamental research](#) at Sanford Burnham Prebys and elsewhere, and our continued efforts will be especially important when, not if, another viral outbreak occurs."

More information: Laura Martin-Sancho et al. Functional Landscape of SARS-CoV-2 Cellular Restriction, *Molecular Cell* (2021). [DOI: 10.1016/j.molcel.2021.04.008](#)

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