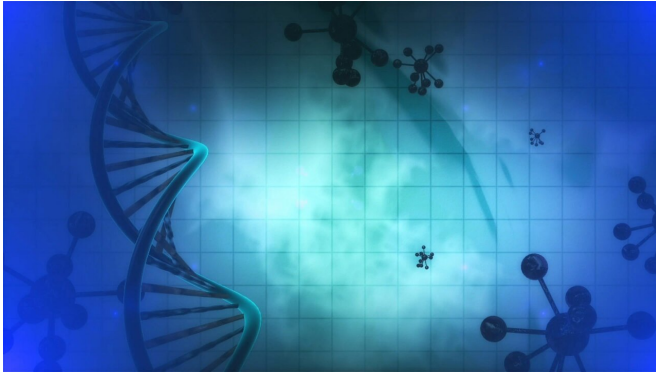


Genes could be key to new COVID-19 treatments, study finds

11 December 2020



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Potential treatments for COVID-19 have been identified after the discovery of five genes associated with the most severe form of the disease.

Genetic evidence is second only to [clinical trials](#) as a way to tell which treatments will be effective in a disease. Existing drugs that target the actions of the [genes](#) reveal which drugs should be repurposed to treat COVID-19 in clinical trials, experts say.

Genes involved in two molecular processes—antiviral immunity and lung inflammation—were pinpointed. The breakthrough will help doctors understand how COVID-19 damages lungs at a molecular level.

Researchers from the University of Edinburgh made the discovery by studying the DNA of 2,700 patients in 208 intensive care units (ICUs) in the UK.

Researchers from the GenOMICC consortium—a global collaboration to study genetics in [critical illness](#)—compared the genetic information of

COVID-19 patients in ICU with samples provided by healthy volunteers from other studies, such as UK Biobank, Generation Scotland and 100,000 Genomes.

The team found key differences in five genes of the ICU patients compared with samples provided by healthy volunteers. The genes—IFNAR2, TYK2, OAS1, DPP9 and CCR2—partially explain why some people become desperately sick with COVID-19, while others are not affected.

Having highlighted the genes, the team were then able to predict the effect of drug treatments on patients, because some genetic variants respond in a similar way to particular drugs.

For example, they showed that a reduction in the activity of the TYK2 gene protects against COVID-19. A class of anti-inflammatory drugs called JAK inhibitors, which includes the drug baricitinib, produces this effect.

They also discovered that a boost in the activity of the gene INFAR2 is also likely to create protection, because it is likely to mimic the effect of treatment with interferon—proteins released by cells of the immune system to defend against viruses. However, experts caution that to be effective, patients might need the treatment early in disease.

Based on the findings published in *Nature*, the researchers say that clinical trials should focus on drugs that target these specific antiviral and anti-inflammatory pathways.

Dr. Kenneth Baillie, the project's chief investigator and Academic Consultant in Critical Care Medicine and Senior Research Fellow at University of Edinburgh's Roslin Institute, said: "This is a stunning realisation of the promise of human genetics to help understand critical illness. Just like in sepsis and influenza, in COVID-19, damage to the lungs is caused by our own immune system,

rather than the virus itself. Our genetic results provide a roadmap through the complexity of immune signals, showing the route to key [drug](#) targets.

"Our results immediately highlight which drugs should be at the top of the list for clinical testing. We can only test a few drugs at a time, so making the right choices will save thousands of lives.

"This work is only possible because of the generous contribution of the patients themselves and their families, research teams in NHS hospitals across the country, and the generous funding we've received from the public and organisations."

GenOMICC (Genetics of Susceptibility and Mortality in Critical Care) started in 2015 as an open, global consortium of intensive care clinicians dedicated to understanding genetic factors that influence outcomes in intensive care from diseases such as SARS, influenza and sepsis. Throughout 2020 it has been focused on COVID-19 research in partnership with Genomics England.

More information: *Nature* (2020). [DOI: 10.1038/s41586-020-03065-y](#)

Provided by University of Edinburgh

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