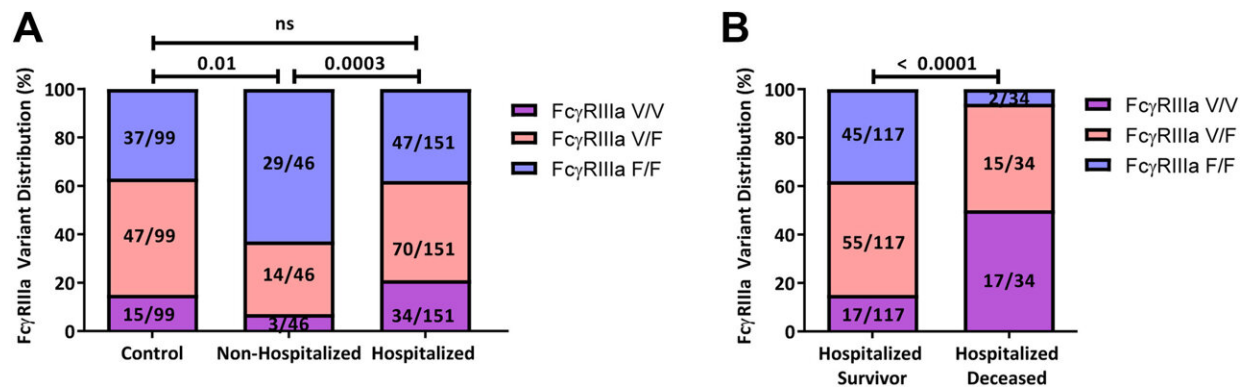


Distinct cellular receptor mutations influence COVID-19 disease severity

May 12 2022, by Johannes Angerer



FcγRIIIa-158-V/F variant distribution. A. Distribution of FcγRIIIa-158-V/F variants between control persons (n = 99), nonhospitalized (n = 46), and hospitalized (N = 151) patients with COVID-19. B. Distribution of FcγRIIIa-158-V/F variants between hospitalized survivors (N = 117) and hospitalized deceased (N = 34) patients with COVID-19. Bars represent the relative frequency of FcγRIIIa-158-V/V, FcγRIIIa-158-V/F, and FcγRIIIa-158-F/F genotypes. χ^2 test was used for statistical comparison between variants. FcγR, Fcγ-receptor. Credit: *Genetics in Medicine* (2022). DOI: 10.1016/j.gim.2022.04.005

Distinct genetic variants significantly influence an individual's immune response to the SARS-CoV-2 virus and may influence the COVID-19 disease severity. A research group led by MedUni Vienna was able to demonstrate that distinct genetic variants of the CD16a antibody

receptor are associated with the risk of severe COVID-19. This affects about 15% of the population. The results are now published in the journal *Genetics in Medicine*.

Natural killer [cells](#) (NK cells) play a significant role in combating the [viral replication](#) in the initial stages of viral infections. NK cells have specialized receptors on their surface that bind to antibodies that are specifically produced against viruses. This enables the antibody-dependent activation of killer cells (ADCC), which leads to the destruction of virus-infected cells and triggers the release of pro-inflammatory factors.

This interaction between antibodies and the NK cell surface receptor is influenced by certain [genetic factors](#), resulting in either strongly (high-affinity) or weakly (low-affinity) binding genetic receptor variants. Working in collaboration with Alexander Zoufaly from the Favoriten hospital, a research group led by Hannes Vietzen and Elisabeth Puchhammer-Stöckl from the Center for Virology of the Medical University of Vienna has now shown that certain genetic variants of the CD16a antibody receptor are associated with the risk of severe COVID-19.

In their study, recently published in the journal *Genetics in Medicine*, the authors show that people who had to be hospitalized with severe COVID-19 were significantly more likely to have the high-affinity [variant](#) of the CD16a receptor. This high-affinity variant only occurs in around 15% of the population, and carriers of this variant have a significantly higher risk of developing severe COVID-19. This high-affinity variant was particularly common in COVID-19 patients who had to be treated in intensive care units or who died from COVID-19.

In subsequent cell culture experiments, the research team was able to show that this high-affinity variant of the antibody receptor results in a

significantly elevated antibody-dependent activation of NK cells and in a particularly strong release of pro-inflammatory factors.

Hannes Vietzen explains: "The antibody-dependent activation of NK cells is a delayed immune response. It now appears that this particular immune response no longer helps to control the SARS-CoV-2 viral replication but aggravates the course of COVID-19 disease by triggering an exaggerated immune response."

The tests involved are special scientific assays. Routine laboratory testing for these parameters is not being considered, since there are currently no therapeutic or preventive options targeting this [genetic predisposition](#) to reduce the risk of severe COVID-19. Genetic predisposition is further only one of several factors that influence the severity of the disease.

More information: Hannes Vietzen et al, High-affinity FcγRIIIa genetic variants and potent NK cell-mediated antibody-dependent cellular cytotoxicity (ADCC) responses contributing to severe COVID-19, *Genetics in Medicine* (2022). [DOI: 10.1016/j.gim.2022.04.005](#)

Provided by Medical University of Vienna

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