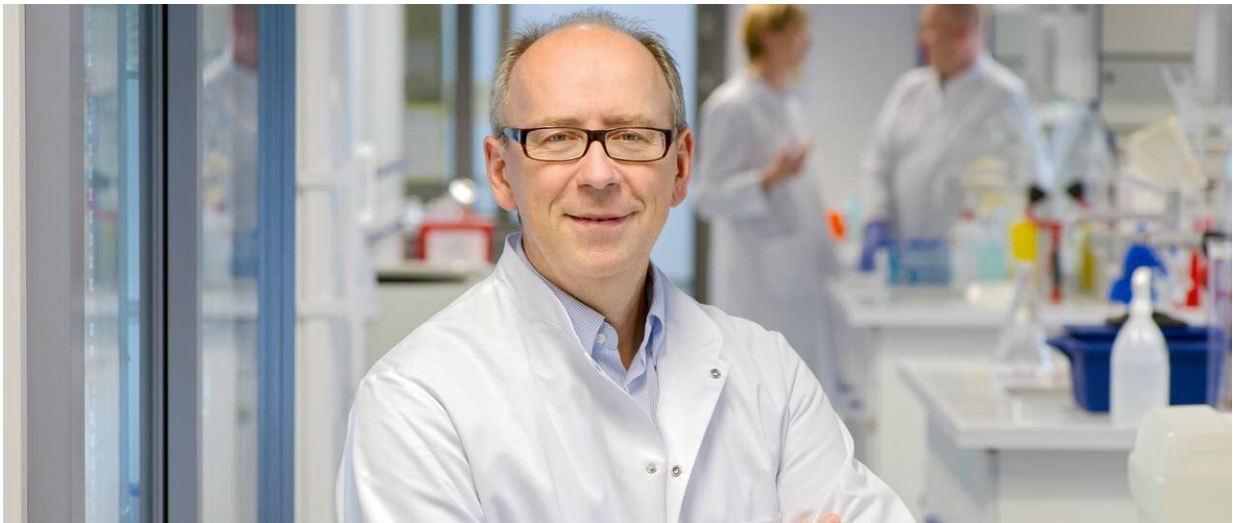


Fatty liver hepatitis is caused by auto-aggressive immune cells

March 24 2021



Prof. Percy Knolle. Credit: Andreas Heddergott

Non-alcoholic steatohepatitis (NASH), often called 'fatty liver hepatitis,' can lead to serious liver damage and liver cancer. A team of researchers at the Technical University of Munich (TUM) has discovered that this condition is caused by cells that attack healthy tissue—a phenomenon known as auto-aggression. Their results may help in the development of new therapies to avoid the consequences of NASH.

Fatty liver disease (NASH) is often associated with obesity. However, our understanding of the causes has been very limited. A team working

with the immunologist Prof. Percy Knolle of TUM has now explored this process step by step in model systems based on mice—and gained promising insights into the mechanisms causing NASH in humans. "We have seen all of the steps observed in the model systems in human patients," says Prof. Knolle. The team's results will be published in *Nature*.

Auto-aggressive immune cells destroy liver tissue

The [immune system](#) protects us against bacteria and viruses and the development of cancerous tumors. The so-called CD8 killer T [cells](#) play an important role here. They specifically recognize infected body cells and eliminate them. With fatty liver hepatitis, the CD8 T cells have lost this targeted deactivation ability. "We have discovered that, in NASH, the [immune cells](#) are not activated by certain pathogens, but rather by metabolic stimuli," says Michael Dudek, the first author of the study. "The T cells activated in this way then kill liver cells of all types."

Sequential activation of T cells

Until that point, the immune cells undergo a unique, step-by-step—and previously unknown—activation process. The T cells develop their auto-aggressive properties only when exposed to inflammation signals and products of fat metabolism in the right order. "Like when we use the combination to unlock a safe, the T cells are switched to 'deadly mode' only through the defined sequence of activation stimuli," says Prof. Knolle, a professor of molecular immunology at TUM. As the trigger for the killing of tissue cells, the international team of researchers identified a basically harmless metabolite: the presence of the energy-carrying molecule ATP outside cells. When auto-aggressive CD8 T cells in the liver reacted with ATP, they destroyed nearby cells, thus causing NASH.

Auto-aggression, but not an auto-immune disorder

The destruction of tissue through auto-aggressive immune cells, as discovered by the researchers, differs from familiar auto-immune disorders, in which immune system cells specifically attack certain cells in the body. The authors note, however, that the tissue-destroying auto-aggressive T cells may also play a role in auto-immune pathologies that has yet to be discovered.

New therapies for fatty liver hepatitis

Until now, the only way of reversing the effects of fatty [liver](#) hepatitis was to eliminate the underlying factors—namely obesity and a high-calorie diet. In other words, patients had to change their lifestyles. The realization that the disease is caused by activated immune cells now suggests possibilities for the development of new therapies. "The destructive auto-aggressive form of the immune response is fundamentally different from the protective T cell immune response to viruses and bacteria," says Prof. Knolle. He is confident that further research can identify targeted immunotherapies that simply prevent the destruction of tissue.

More information: Auto-aggressive CXCR6+ CD8 T cells cause liver immune pathology in NASH, *Nature* (2021). [DOI: 10.1038/s41586-021-03233-8](https://doi.org/10.1038/s41586-021-03233-8)

Provided by Technical University Munich

Citation: Fatty liver hepatitis is caused by auto-aggressive immune cells (2021, March 24) retrieved 25 April 2024 from <https://medicalxpress.com/news/2021-03-fatty-liver-hepatitis-auto-aggressive-immune.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.