

Exploring better fatty liver remedies

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Flinders Emeritus Professor Greg Barritt, College of Medicine and Public Health, at work in his lab. Credit: Flinders University

Obesity often leads to increased accumulation of fat in the liver, which can, over time, result in diabetes, poor liver function, and liver cancer. It's increasingly a problem affecting obese children and young people who need exercise and good diet—but are not likely to embrace those solutions, wanting a pill instead to remedy the fatty liver problem. The key problem being that no such pill currently exists.

Flinders University medical researchers are now investigating a different type of drug targeting cellular/biochemical pathways that may reduce or prevent fat/lipid accumulation in liver, to ultimately reduce risk of cancer and Type 2 diabetes.

Non-alcoholic fatty liver disease (NAFLD) is the most common cause of paediatric liver disease, affecting 10% of [school-aged children](#) and 44–70% of obese children and young people (CYP) in the western world. Fat accumulation in the liver at an early age seems to be a critical factor in pre-disposing a young obese person to develop

serious liver disease, including [liver cancer](#), later in life.

"The best way for patients to reduce fat is the correct diet and good exercise, but most people do not want to do this, so drugs targeted to reduce fat in the liver are a sought-after alternative," says Flinders University Emeritus Professor Greg Barritt, who has researched the condition for many years.

"We are researching just one of these drug pathways which involves effects of altered calcium inside [liver cells](#) which seem to exacerbate fat accumulation."

The research team's initial review—"Deranged hepatocyte intracellular Ca²⁺ homeostasis and the progression of [non-alcoholic fatty liver disease](#) to [hepatocellular carcinoma](#)," by Eunos Ali, Grigori Rychkov and Greg Barritt, published in the journal *Cell Calcium*, is directed at better understanding the pathways involving calcium before moving forward to identify specific drug targets for this pathway.

"We are conducting experiments now and are in early stages, but we are confident that this is a significant path forward in battling a problem that continues to increase—especially among young people," says Professor Barritt.

This research adds to improved understanding about the cause and effect in the fat-to-cancer pathway, mutations and the time sequences involved in this process.

More information: Eunos S. Ali et al. Deranged hepatocyte intracellular Ca²⁺ homeostasis and the progression of non-alcoholic fatty liver disease to hepatocellular carcinoma, *Cell Calcium* (2019). [DOI: 10.1016/j.ceca.2019.102057](https://doi.org/10.1016/j.ceca.2019.102057)

Provided by Flinders University

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