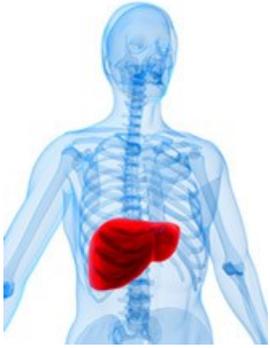


Straight from the gut: Microbes can cause obesity

By [Helen Dodson](#)

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Obesity and chronic liver disease can be triggered by a family of proteins that alter populations of microbes in the stomach, a discovery that suggests the condition may be infectious, Yale scientists report. The study, in the advance online publication of [Nature](#), expands on earlier Yale research that showed how similar microbial imbalances caused by the same family of proteins increases the risk of intestinal diseases such as colitis.



The Yale scientists' most extraordinary finding, they said, was that the altered intestinal environment that led to obesity and liver disease was infectious among the community of mice. "When healthy mice were co-housed with mice that had altered gut microbes, the healthy mice also developed a susceptibility for development of liver disease and obesity," said senior author Richard A. Flavell, professor of immunobiology at [Yale School of Medicine](#) and a Howard Hughes Medical Institute investigator.

The proteins in question are called inflammasomes. They are responsible for launching the immune system's inflammatory response. Inflammasomes act as sensors and regulators of the microbial environment of the intestines.

The Yale team found that a deficiency in components of two particular inflammasomes in mice resulted in the development of an altered microbial community associated with increased bacteria. This determined the severity of non-alcoholic fatty liver disease (NAFLD) and obesity in the mice.

NAFLD is the result of metabolic syndrome, a collection of disorders that includes obesity and diabetes, and is the leading cause of chronic liver disease in the western world. It is estimated that up to 30 million people suffer from NAFLD in the United States alone. Twenty percent of people with NAFLD develop chronic liver inflammation, placing them at risk for cirrhosis and liver cancer, but the causes have been unclear.

The next step, Flavell said, is extending this research to humans and to identify more precisely the bacteria involved in the progression to liver disease. "We found, in mice, that targeted antibiotic treatment brought the microbial composition back to normal, and thus eased the liver disease. Our hope is that our findings may eventually lead to a treatment for humans."

The researchers who led this project in Flavell's laboratory are Jorge Henao-Mejia, Eran Elinav, and Chengcheng Jin. Other participating researchers were Liming Hao, Wajahat Z. Mehal, Till Strowig, Christoph A. Thaiss, Stephanie C. Eisenbarth, Michael J. Jurczak, Joao-Paulo Camporez, and Gerald I. Shulman of Yale; Andrew L. Kau and Jeffrey I. Gordon of Washington University School of Medicine; and Hal M. Hoffman of the University of California at San Diego.

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For more information on Flavell's research, visit his [website](#).

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