

# New study reveals links between alcoholic liver disease and the circadian clock

January 17 2014

---



Researchers from the University of Notre Dame and the Indiana University School of Medicine have revealed a putative role for the circadian clock in the liver in the development of alcohol-induced hepatic steatosis, or fatty liver disease.

Hepatic steatosis is the abnormal accumulation of fats in the cells of the liver, and is linked to disturbed control of [fat metabolism](#). Alcohol-induced liver steatosis is produced by excessive alcohol consumption and is linked to hepatitis, or inflammation of the liver. It can be a precursor to an even more serious illness, [liver cirrhosis](#), which includes scarring of the liver. Ten percent to 35 percent of chronic heavy drinkers develop alcoholic hepatitis, and it is the main cause of liver disease in Western countries.

The team, led by associate professors Giles Duffield, from Notre Dame's Department of Biological Sciences and Eck Institute for Global Health, and Suthat Liangpunsakul from the Indiana University School of Medicine's Department of Medicine, Division of Gastroenterology and Hepatology, is interested in the molecular genetic basis for the molecular clock and liver steatosis.

The study, using molecular biological approaches and long-term alcohol feeding of experimental mice, reveals that the development of liver steatosis produced by alcohol abuse is intertwined with disturbances of the normal operation of the 24-hour clock system located in the cells of the liver. Importantly, this change in the liver clock seems to occur independently from the [master clock](#) system located in the brain.

The [circadian clock](#) regulates 24-hour rhythms in biochemistry, physiology and behavior, and its normal operation and appropriate synchronization to the external world, especially the alternating cycle of day and night, is critical to maintaining a normal healthy state. Disturbances of the clock have been linked to mental health disorders, to metabolic disease including obesity and diabetes, and to the development of cancer.

The liver plays many roles in the body, and includes control of metabolism, storage and release of energy molecules, and detoxification.

"Liver function changes daily in a rhythmic manner and is coordinated with cycles of feeding-fasting and to the energy demands of the body, such as activity and rest," Duffield said. "These daily rhythms are regulated by the circadian clock within those [liver cells](#), and disturbances to the molecular clock mechanism or poor temporal coordination of the clock with the timing of eating, or the sleep-wake and rest-activity cycle, can lead to illness."

The study suggests that either the circadian clock is important in the actual development of the [liver disease](#) or that the development of steatosis disrupts the normal pattern of the clock mechanism. The researchers' findings also offer novel insights into how the disease might be manipulated for clinical purposes.

Interestingly, the mechanism by which chronic alcohol intake is thought to alter the control of fat metabolism in the [liver](#) is also a shared signal to the circadian clock mechanism, this being the ratio of production of reduced nicotinamide adenine dinucleotide, or NADH, to NAD<sup>+</sup>. The authors suggest that this may be a key to the shared disturbance to the two biological mechanisms of lipid metabolism and the circadian clock.

The study, Zhou et al., was published this week in the journal *Scientific Reports*.

**More information:** [www.nature.com/srep/2014/14011.../full/srep03725.html](http://www.nature.com/srep/2014/14011.../full/srep03725.html)

Provided by University of Notre Dame

Citation: New study reveals links between alcoholic liver disease and the circadian clock (2014, January 17) retrieved 19 April 2024 from <https://medicalxpress.com/news/2014-01-reveals-links->

[alcoholic-liver-disease.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.