

Study reveals why heart failure patients suffer depression, impaired thinking

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Professor Tami Martino, University of Guelph. Credit: University of Guelph

Heart failure patients often have trouble with thinking and depression.

A new study by University of Guelph researchers explains why and

points to ways to prevent and treat both [heart](#) and brain maladies through the emerging field of circadian medicine.

Published recently in Nature's *Scientific Reports*, the study is the first to reveal how cognition and mood in mice are regulated by the [body clock](#) and how pertinent brain regions are impaired in heart failure, said Tami Martino, a professor in U of G's Department of Biomedical Sciences and director of the Centre for Cardiovascular Investigations.

"Neurosurgeons always look in the brain; cardiologists always look in the heart. This new study looked at both," said Martino, whose work in the emerging field of circadian medicine is supported by funding from the Canadian Institutes of Health Research. She recently received a Mid-Career Investigator Award from the Heart and Stroke Foundation of Canada.

Coronary heart disease, the most common cause of heart failure, causes one in three deaths in Canada, according to the Heart and Stroke Foundation.

Human patients with heart failure often have neurological conditions such as cognitive impairment and depression, said Martino. She worked on the study with master's student Austin Duong and Ph.D. student Cristine Reitz—both co-first authors—and neuroscientists including U of G psychology professor Boyer Winters and biomedical sciences professor Craig Bailey.

Martino suspected the heart-brain connection involved the circadian mechanism molecule, called "clock."

Circadian rhythms in humans and other organisms follow Earth's 24-hour cycle of light and darkness, signalling when to sleep and when to be awake.

Martino's earlier research showed how disrupting circadian rhythms—as with shift workers, jet-lagged travellers and patients disturbed in intensive-care units—can trigger changes that worsen heart disease and impair overall health and well-being.

For this new study, the researchers compared normal mice with mice carrying a mutation in their circadian mechanism (called "clock mice"). They found that the mutation affected the structure of neurons in brain areas important for cognition and mood.

Working with University of Toronto colleagues, the team also found differences in clock regulation of blood vessels in the brains of the clock mice.

After inducing heart failure in mice to simulate human heart failure, they used microarray profiling to identify key genes in the brain that were altered in neural growth, stress and metabolism pathways.

The results show that the circadian mechanism influences neural effects of heart failure, said Martino. Pointing out that no cure exists for the heart condition, she said understanding how the circadian mechanism works in the [brain](#) may lead to new strategies to improve patients' quality of life.

Patients recovering from heart attacks often experience disturbed circadian rhythms from light, noise and interactions with hospital staff at night. "Maintaining [circadian rhythms](#) especially for patients with heart disease could lead to better health outcomes."

More generally, the findings point to potential health benefits for people in general. Avoiding shift work for people with underlying heart conditions or [sleep disorders](#), reducing light at night or avoiding [social jet lag](#) (going to bed late and waking up later than usual on weekends)

could all help reduce neurobiological impairments.

Those problems—and potential solutions—involve not just hearts but brains, she said. "If we're not yet able to cure [heart failure](#), we should at least be focusing on how we can improve quality of life for patients."

More information: Austin T. H. Duong et al, The Clock Mechanism Influences Neurobiology and Adaptations to Heart Failure in Clock Δ 19/ Δ 19 Mice With Implications for Circadian Medicine, *Scientific Reports* (2019). [DOI: 10.1038/s41598-019-41469-7](https://doi.org/10.1038/s41598-019-41469-7)

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