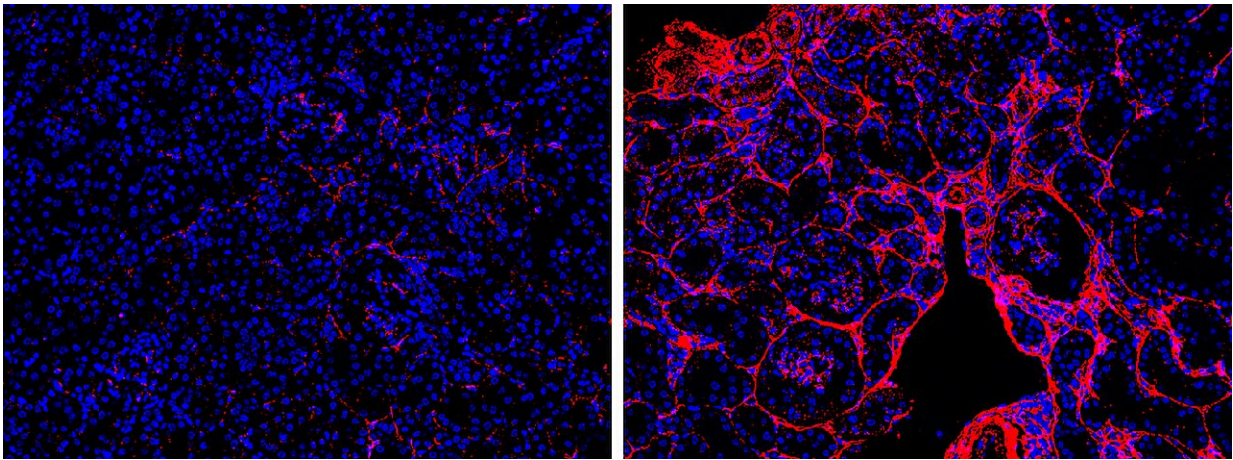


# Study shows glucagon is key for kidney health

February 23 2024

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The image on the left shows a normal kidney; the image on the right shows a kidney lacking the glucagon receptor and displaying a high degree of scarring. Scarring is visualized with a collagen III antibody shown at 20x magnification. Credit: UT Southwestern Medical Center

Glucagon, a hormone best known for promoting blood sugar production in the liver, also appears to play a key role in maintaining kidney health. When UT Southwestern Medical Center researchers removed receptors for this hormone from mouse kidneys, the animals developed symptoms akin to chronic kidney disease (CKD).

Their findings, [published](#) in *Cell Metabolism*, shed new light on

glucagon's physiological functions and provide new insights into CKD, a disease that affects hundreds of millions of people around the globe, according to the National Institute of Diabetes and Digestive and Kidney Diseases.

"Our study defines important protective effects of glucagon for kidney health and normal systemic metabolic well-being of the entire organism," said study leader Philipp Scherer, Ph.D., Professor of Internal Medicine and Cell Biology and Director of UTSW's Touchstone Center for Diabetes Research.

Over the past century, researchers have discovered that cells in the pancreas produce glucagon when blood sugar, or glucose, dips below a certain threshold. This hormone migrates through the bloodstream to receptors on the surface of liver cells, prompting the liver to produce glucose that fuels cells throughout the body. More recent research has shown that the kidneys also bear glucagon receptors, but besides stimulating production of a minor amount of glucose, their role has been unclear, Dr. Scherer explained.

To better understand the function of these kidney-based glucagon receptors, Dr. Scherer and his colleagues used [genetic techniques](#) to eliminate the receptors in mice and compared them to mice without genetic manipulations and others with glucagon receptors deleted in the liver.

Unlike the other two groups, the mice that had glucagon receptors removed in the kidneys showed a host of pathologies that plagued this organ. These included inflammation, scarring, and excess lipid deposits similar to what is seen in [fatty liver disease](#), as well as [high blood pressure](#) and associated kidney-related damage, alterations in the activity of energy-production genes, and signs of high oxidative stress.

Mice without kidney-based glucagon receptors also had a range of deficits stemming from kidney dysfunction that affected their entire bodies, such as a dysregulation of nitrogen, problems maintaining water and electrolyte balances, and heart problems.

These issues largely mimic those in patients with CKD, noted May-Yun Wang, Ph.D., Assistant Professor of Internal Medicine and first author of the study. Studies have shown that individuals with CKD have fewer kidney glucagon receptors, although it's unclear which occurred first—kidney pathology that decreased receptor numbers or pathology that arose from an insufficient amount of receptors. This question is a topic for future research, Dr. Wang said.

In the meantime, she added, newer drugs in late-stage clinical trials to treat obesity and diabetes are incorporating [glucagon](#), a strategy found to aid in weight loss that could also unexpectedly benefit patients with CKD.

"These drugs are already showing improvements to kidney health in [clinical trials](#), and our findings provide an explanation why," Dr. Scherer said.

**More information:** May-Yun Wang et al, Downregulation of the kidney glucagon receptor, essential for renal function and systemic homeostasis, contributes to chronic kidney disease, *Cell Metabolism* (2024). [DOI: 10.1016/j.cmet.2023.12.024](https://doi.org/10.1016/j.cmet.2023.12.024)

Provided by UT Southwestern Medical Center

Citation: Study shows glucagon is key for kidney health (2024, February 23) retrieved 12 May 2024 from <https://medicalxpress.com/news/2024-02-glucagon-key-kidney-health.html>

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