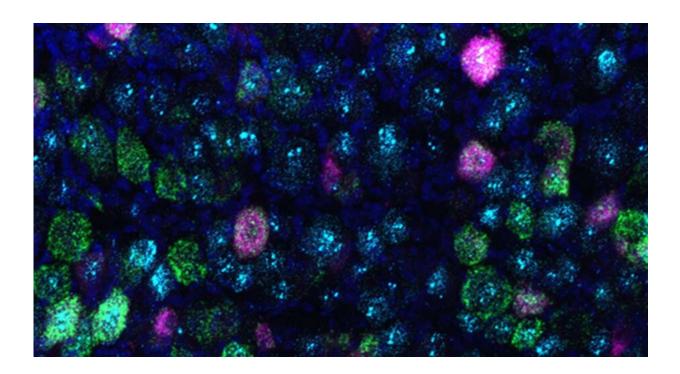


Scientists identify sensor protein that underlies bladder control

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An image of sensory neurons, with those that detect signals in the bladder labeled in magenta and the "mechanosensor" protein PIEZO2 in cyan. Credit: Courtesy of Adam Coombs.

A team co-led by scientists at Scripps Research has found that the main sensor protein enabling our sense of touch also underlies the feeling of having a full bladder and makes normal bladder function possible.



The discovery, published Oct. 14 in *Nature*, marks a key advance in basic neurobiology and may also lead to better treatments for <u>bladder</u> control and urination problems, which are common especially among the elderly.

"We tend to take urination for granted, and it has been under-studied, yet it's a huge burden when something goes wrong with this system," says the study's lead author Kara Marshall, Ph.D., a postdoctoral research associate in the Department of Neuroscience at Scripps Research. "Now we've identified a crucial part of how urination normally works."

Marshall and her colleagues focused in this study on the PIEZO2 protein , a "mechanosensor" that detects the physical stretching of tissues where it resides. They found that PIEZO2 is expressed in cells of the bladder and is necessary for normal urinary continence and functioning in both mice and humans.

"Who would have imagined that the same mechanosensor protein enabling our <u>sense of touch</u> also alerts us that our bladder is full?" says co-senior author Ardem Patapoutian, Ph.D., Professor and Presidential Endowed Chair in Neurobiology at the Dorris Neuroscience Center at Scripps Research, and a Howard Hughes Medical Institute investigator.

In 2010, Patapoutian and his lab first identified PIEZO2 and its sister protein PIEZO1 as mechanosensors that sense mechanical distortions of tissues. For that feat, among others, Patapoutian was a co-recipient of the prestigious 2020 Kavli Prize in Neuroscience.

Like most sensor proteins, the PIEZOs are ion-channel proteins, which are embedded in their host cell's outer membrane and, when triggered by a stimulus, allow a flow of charged atoms into the cell. Sensor ionchannel proteins are usually found in <u>sensory neurons</u> in the skin, joints and other organs. On a given neuron, when enough of these channels



open to admit ion flows, the neuron will fire a nerve signal to the brain.

For PIEZOs, the stimulus that triggers the opening of the ion channel is the stretching of the cell membrane due to mechanical forces on the local tissue. In studies over the past decade, Patapoutian and his colleagues have shown that PIEZO2 is expressed is different organs and tissues throughout the body. For example, they exist in lung tissues to sense lung stretch and help regulate breathing, in blood vessels to sense blood pressure and in the skin to mediate the sense of touch.

The new study was a collaboration with Alexander Chesler, Ph.D., and Carsten Bönnemann, MD, senior investigators at the National Institutes of Health. Chesler and Bönnemann, and their colleagues, have been studying people born with genetic mutations that result in the functional loss of PIEZO2. These individuals suffer various impairments in sensory pathways known to be PIEZO2-related.

For the study, NIH investigators found that these PIEZO2-deficient individuals, in addition to their other sensory deficits, lack the normal sense of having a full bladder. They typically urinate on a schedule to avoid incontinence and have trouble completely emptying their bladder when they do urinate.

Patapoutian, Marshall and their colleagues showed in experiments that the loss of PIEZO2 has similar effects in mice. The urinary tract uses PIEZO2 protein in both bladder sensory neurons and in bladder-lining cells called umbrella cells to detect stretch and facilitate urination, indicating a two-part sensor system. As they determined in experiments, bladder neurons in mice normally respond robustly with nerve signals when the bladder is filled but are almost completely silent during bladder filling if they lack PIEZO2.

The mice lacking PIEZO2 in their lower urinary tracts also showed



abnormal urination reflexes in the muscles controlling the urethra, the duct in which urine flows from the bladder. That suggests that in mice and most likely in people, the mechanosensor protein is needed both for normal bladder-stretch sensation and for normal urination.

The team is currently following up with research on the distinct roles of bladder neurons and umbrella cells, and how they signal to each other. They are also investigating the possible roles of other mechanosensors, such as PIEZO1, in bladder control and urination.

"Mice without PIEZO2 had clear urination deficits, but ultimately were still able to urinate, so that suggests another mechanosensory protein may be involved," Marshall says.

More information: Kara L. Marshall et al, PIEZO2 in sensory neurons and urothelial cells coordinates urination, *Nature* (2020). DOI: 10.1038/s41586-020-2830-7

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