

## Hidden epigenetic mechanisms behind chronic UTIs found

**b** 5,000 Culture in matrigel #2 4,000 #3 TER (ohm×cm<sup>2</sup> -3 c 3-5 d 3,000 #4 Dissociate spheroids, seed in transwells Switch to 5% CM 2,000 Isolate mouse urothelial cells for differentiation 1,000 10-15 passages before seeding n 12 20 8 16 Days of culture С K20 F-actin f Ecad DAP 20 µI d g p63 DAF Upk3a+/p63+ cell (intermediate cells) 20 µm h K5+/K14- cell

The culture of USCs of juvenile C3H/HeN mice regenerates differentiated urothelium in vitro. a, USCs isolated from 8-week-old C3H/HeN mice were expanded by spheroidal culture in matrigel with 50% L-WRN conditioned media (CM) including Y-27632, a ROCK inhibitor, and SB431542, a TGF  $\beta$  type 1

April 18 2023, by Justin Jackson



inhibitor. After 3 d of spheroid culture, cells were dissociated into a single-cell suspension and  $3-4 \times 10^5$  cells were seeded onto transwell membranes. The cells were cultured in 50% CM for 3–5 d, then cultured in 5% CM for 2–3 weeks until full differentiation. b, Cell cultures with a TER value >4,000 ohm  $\times$  cm<sup>2</sup> were then analyzed (5 transwells cultured from one juvenile C3H/HeN cell line). For consistency, TER was measured 1 d after media change. c,d, Differentiated urothelia on the transwells were fixed and imaged via (c) confocal microscopy and (d) SEM to show a top-down view of the urothelium at magnification  $500 \times$ (top panel) and 10,000× (bottom panel). In c, samples were stained for F-actin, the terminal differentiation marker K20 and nuclei (DAPI). e-h, The urothelia were also paraffin-embedded, sectioned and stained with hematoxylin and eosin (H&E) (e) and immunostained for K20, Ecad and DAPI (f), Upk3a, p63 and DAPI (g) or K5, K14 and DAPI (h). Representative images are shown. Data are from 2-3 independent experiments using USCs from 5 different juvenile C3H/HeN mice. Credit: Nature Microbiology (2023). DOI: 10.1038/s41564-023-01346-6

Published in *Nature Microbiology*, a study in mice finds that the microbial basis for urinary tract infections can include localized epigenetic alterations that make additional infections more likely.

Previous <u>urinary tract infections</u> (UTIs) are known to predispose patients to future infections, but the mechanisms behind the recurrences are not well understood. Researchers from the Washington University School of Medicine in St Louis followed up on their previous research, which found differential epithelial cellular changes after UTIs in mice bladders, leading to different outcomes.

The current study, "Uropathogenic Escherichia coli <u>infection</u>-induced epithelial trained immunity impacts urinary tract disease outcome," compared urothelial stem cell (USC) lines isolated from mice with a history of either resolved or chronic uropathogenic Escherichia coli



(UPEC) infection, and found epigenetic changes, including differences in chromatin accessibility, DNA methylation and histone modification. A News & Views article on the research was published in the same journal issue.

One clue the researchers followed was that previous infection results in urothelial stem cell changes that persist over many generations of cell culture, indicating a possible epigenetic modification within those cells. Epigenetic modifications are not changes to the DNA sequences of the cells but in how DNA is accessed and, therefore, which genes are expressed. These sorts of changes can alter normal cell function and, in this case, immunity responses.

They discovered that UPEC infection acts as an epi-mutagen, reprogramming the functional activity of urothelial cells, leading to remodeling and an alteration or trained innate response to subsequent infections. The research demonstrates the direct role of a mucosal bacterial infection in eliciting specific <u>epigenetic changes</u> in mucosal epithelial stem cells.

While not proposing a solution, the research on mice does identify previously unknown mechanisms that inform future research in devising solutions to chronic UTIs. This could be a significant and actionable elevation of our knowledge on the subject that currently only offers recommendations on how to wipe after using the toilet, suggestions to keep genitals dry and encouraging copious amounts of cranberry juice drinking as strategies to prevent reinfection.

Even without a current curative solution, the study may be welcome news for those suffering from chronically reoccurring infections in confirming that the problem is an epigenomic-based immunity issue and not somehow a failure on their part to follow instructions on basic hygiene or consume enough cranberry juice.



**More information:** Seongmi K. Russell et al, Uropathogenic Escherichia coli infection-induced epithelial trained immunity impacts urinary tract disease outcome, *Nature Microbiology* (2023). <u>DOI:</u> <u>10.1038/s41564-023-01346-6</u>

Soumitra Mohanty, John Kerr White and Annelie Brauner, Epigenetic modifications influence urinary tract infection outcome, *Nature Microbiology* (2023). DOI: 10.1038/s41564-023-01371-5

© 2023 Science X Network

Citation: Hidden epigenetic mechanisms behind chronic UTIs found (2023, April 18) retrieved 21 April 2023 from https://medicalxpress.com/news/2023-04-hidden-epigenetic-mechanisms-chronic-utis.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.