

Scientists describe 'multi-kingdom dialogue' between internal and external microbiota

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The microbiome is comprised of microorganisms that live in and on us and contribute to human health and disease. Credit: NHGRI

National Institutes of Health scientists and their collaborators have identified an internal communication network in mammals that may regulate tissue repair and inflammation, providing new insights on how diseases such as obesity and inflammatory skin disorders develop. The new research is published in *Cell*.

The billions of organisms living on body surfaces such as the skin of mammals—collectively called [microbiota](#)—communicate with each other and the host immune system in a sophisticated network. According to the study, viruses integrated in the host genome, remnants of previous infections called endogenous retroviruses, can control how the host immune system and the microbiota interact, affecting tissue repair and antimicrobial defenses. Endogenous retroviruses can comprise up to 10% of all genes.

The newly discovered role of endogenous retroviruses adds to the scientific community's understanding of certain diseases and

inflammatory states and opens new research avenues. "Together, our results support the idea that mammals may have co-opted their endogenous viromes as a means to communicate with their microbiota, resulting in a multi-kingdom dialogue that controls both immunity and inflammation," the authors state.

Scientists from NIH's National Institute of Allergy and Infectious Diseases led the project with collaborators from the NIH Center for Human Immunology, the National Cancer Institute, Stanford University and Scripps Research in California, University of Pennsylvania in Philadelphia, University of Oxford, and The Francis Crick Institute in England.

Building on a series of studies over the past decade showing that microbiota broadly promote immune protection, the NIAID scientists and collaborators sought to discover how this occurs. They used *Staphylococcus epidermidis*, a common skin bacterium with known helpful and harmful features, as a study model in laboratory and mouse experiments.

The models helped them identify the important roles of skin cells called keratinocytes and of endogenous retroviruses in communication between microbiota and the skin immune system. Keratinocytes are the primary interface between the host and its microbiota. Their study showed that *S. epidermidis* triggered an [antiviral response](#) in keratinocytes, and that finding led them to discover that endogenous retroviruses coordinate responses to the microbiota that stimulate the [immune system](#).

The mouse model also showed that a high-fat diet triggers an inflammatory immune response to *S. epidermidis* that can be controlled by providing antiretroviral treatment, suggesting a role for [endogenous retroviruses](#) in driving inflammatory responses caused by microbes under high-fat conditions. The researchers will continue exploring

how these ancient viruses control the beneficial role of the microbiota and how nutrition can change this dialogue toward pro-inflammatory responses.

More information: Djalma S. Lima-Junior et al, Endogenous retroviruses promote homeostatic and inflammatory responses to the microbiota, *Cell* (2021). DOI: [10.1016/j.cell.2021.05.020](https://doi.org/10.1016/j.cell.2021.05.020)

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