

Gut bacteria may hold key to treating autoimmune disease

December 19 2016



Foxp3-mutant mice suffer inflammatory damage in their lung tissue (left), but this is prevented by treatment with the bacterium *Lactobacillus reuteri* (right). Credit: He et al., 2017

Defects in the body's regulatory T cells (T reg cells) cause inflammation and autoimmune disease by altering the type of bacteria living in the gut, researchers from The University of Texas Health Science Center at Houston have discovered. The study, "Resetting microbiota by *Lactobacillus reuteri* inhibits T reg deficiency-induced autoimmunity via adenosine A2A receptors," which will be published online December 19 in *The Journal of Experimental Medicine*, suggests that replacing the missing gut bacteria, or restoring a key metabolite called inosine, could



help treat children with a rare and often fatal autoimmune disease called IPEX syndrome.

T reg cells suppress the immune system and prevent it from attacking the body's own tissues by mistake. Defects in T reg cells therefore lead to various types of autoimmune disease. Mutations in the transcription factor Foxp3, for example, disrupt T reg function and cause IPEX syndrome. This inherited autoimmune disorder is characterized by a variety of inflammatory conditions including eczema, type I diabetes, and severe enteropathy. Without a stem cell transplant from a suitable donor, IPEX syndrome patients usually die before the age of two.

Autoimmune diseases can also be caused by changes in the <u>gut</u> <u>microbiome</u>, the population of bacteria that reside within the gastrointestinal tract. In the study, the team led by Yuying Liu and J. Marc Rhoads at The University of Texas Health Science Center at Houston McGovern Medical School find that mice carrying a mutant version of the Foxp3 gene show changes in their gut microbiome at around the same time that they develop autoimmune symptoms. In particular, the mice have lower levels of bacteria from the genus *Lactobacillus*. The researchers discovered that by feeding the mice with *Lactobacillus reuteri*, they could "reset" the gut bacterial community and reduce the levels of inflammation, significantly extending the animals' survival.

Bacteria can secrete metabolic molecules that have large effects on their hosts. The levels of a metabolite called inosine were reduced in mice lacking Foxp3 but were restored to normal after resetting the gut microbiome with *L. reuteri*. The researchers found that, by binding to cell surface proteins called adenosine A2A receptors, inosine inhibits the production of Th1 and Th2 cells. These pro-inflammatory T cell types are elevated in Foxp3-deficient mice, but their numbers are diminished by treatment with either *L. reuteri* or inosine itself, reducing



inflammation and extending the animals' life span.

"Our findings suggest that probiotic *L. reuteri*, inosine, or other A2A receptor agonists could be used therapeutically to control T cell-mediated autoimmunity," says Yuying Liu.

More information: He, B., et al. 2017. *J. Exp. Med.* DOI: <u>10.1084/jem.20160961</u>

Provided by Rockefeller University Press

Citation: Gut bacteria may hold key to treating autoimmune disease (2016, December 19) retrieved 5 May 2024 from https://medicalxpress.com/news/2016-12-gut-bacteria-key-autoimmune-disease.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.