

PUBLIC RELEASE: 6-OCT-2016

Newly discovered gut organism protects mice from bacterial infections

CELL PRESS



PRINT E-MAIL

While bacteria are often stars of the gut microbiome, emerging research depicts a more complex picture, where microorganisms from different kingdoms of life are actively working together or fighting against one another. In a study published October 6 in *Cell*, scientists reveal one example: a newly discovered protist that protects its host mice from intestinal bacterial infections.

"This was a serendipitous finding, but an important one," says senior author Miriam Merad, a Professor of Oncological Sciences and of Medicine at the Icahn School of Medicine at Mount Sinai. "This study shows how vital it is to go beyond bacteria when studying the microbiome."

The investigators made the discovery when they realized that mice that had been bred at their own facility had a greater number of immune cells in the gut than mice purchased from an outside vendor. Graduate student Aleksey Chudnovskiy, the study's first author, together with postdoctoral fellow Arthur Mortha, decided to figure out why that was the case. When they performed an intestinal cleanse on the two groups of mice, they were surprised to find

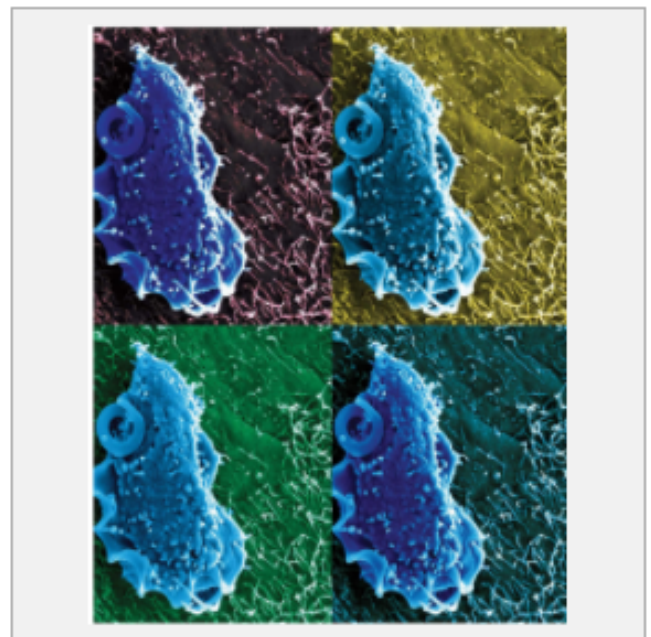


IMAGE: THIS IS A SCANNING ELECTRON MICROSCOPY (SEM) IMAGE OF T.MU IN THE MOUSE COLON. [view more >](#)

CREDIT: CHUDNOVSKIY MERAD

that the mice from the Mount Sinai facility had flagellated protozoa living in their guts. DNA sequencing revealed that the microorganism was a new protozoan parasite, which they named *Tritrichomonas musculus* (T. mu).

Further investigations showed that when this protist was given to the mice that didn't have it, they, too, had an increase in the number of immune cells in their guts and also increased inflammatory cytokines. The researchers set out to discover the underlying mechanism. They found that T. mu activates the inflammasome in the gut epithelial cells of the mice, which in turn led to the activation of cytokines. They also found that dendritic cells were required to induce inflammation.

To determine whether colonization of T. mu in the gut affected the mice's ability to fight off infection, they infected mice with *Salmonella* and found that the animals that had T. mu as part of their microbiome were very resistant to *Salmonella* infection. "The protective effect of this species is very striking," Chudnovskiy says.

T. mu was found to be an ortholog of *Dientamoeba fragilis*, a parasite that's found in the guts of many humans, but the researchers don't know if *D. fragilis* also has a protective effect. It's something they plan to study. "People from industrialized countries traveling to emerging countries are more susceptible to intestinal infection than the indigenous population," Merad explains. "It's possible that protists, which are known to be common in emerging countries, contribute to the protective effect against intestinal pathogenic infections."

She adds: "The fight against pathogens determined the survival of the human species, and those with stronger immune systems are the ones who survived. It is likely that the microbiome is a big part of the evolutionary process. Thus, identifying those commensals that confer immune strength in exposed communities should help identify novel therapeutics."

###

This research was funded by the National Institutes of Health and supported in part by the National Institute of Allergy and Infectious Diseases.

Cell, Chudnovskiy et al: "Host-protozoan interactions protect from mucosal infections through activation of the inflammasome" [http://www.cell.com/cell/fulltext/S0092-8674\(16\)31174-6](http://www.cell.com/cell/fulltext/S0092-8674(16)31174-6)

Cell (@CellCellPress), the flagship journal of Cell Press, is a bimonthly journal that publishes findings of unusual significance in any area of experimental biology, including but not limited to cell biology, molecular biology, neuroscience, immunology, virology and microbiology, cancer, human genetics, systems biology, signaling, and disease mechanisms and therapeutics. Visit: <http://www.cell.com/cell>. To receive Cell Press media alerts, contact press@cell.com

Disclaimer: AAAS and EurekAlert! are not responsible for the accuracy of news releases posted to EurekAlert! by contributing institutions or for the use of any information through the EurekAlert system.