

Study suggests some gut microbes may be keystones of health

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A University of Oregon study finds abundance does not dictate the immune response to bacteria in the guts of zebrafish. This graphic depicts immune response (neutrophils, shown in green) to separate bacterial species, and the response when the two bacteria are placed into a germ-free fish gut at the same time. Credit: Courtesy of Annah S. Rolig

University of Oregon scientists have found that strength in numbers



doesn't hold true for microbes in the intestines. A minority population of the right type might hold the key to regulating good health.

The <u>discovery</u>, based on research using zebrafish raised completely germ free, is reported in a paper published in the Nov. 11 issue of *Cell Host & Microbe*. The findings provide a path to study the function of each bacterial species in the gut and to eventually, perhaps, predict and prevent disease, says lead author Annah S. Rolig, a postdoctoral researcher in the UO's Institute of Molecular Biology.

In the project, researchers watched for immune response as isolates of species of bacteria, normally associated with healthy zebrafish, were introduced one at a time and in combination into previously germ-free <u>intestines</u> of the fish.

In a telling sequence, one bacterial species, Vibrio, drew numerous neutrophils, which indicated a rapid inflammatory response in one fish. Another species, Shewanella, inserted into a separate germ-free fish barely attracted an immune response. In a third germ-free fish, both species were introduced together and assembled with a ratio of 90percent Vibrio to 10-percent Shewanella.

The inflammatory response in the third fish was completely controlled by the low-abundance species.

"Until now, we've only been able to capture proportional information, like you'd see displayed in a pie graph, of the makeup of various microbiota, in percentages of their abundance," Rolig said. "Biologists in this field have typically assumed an equal contribution based on that makeup."

Low counts of a bacterial species generally have been discounted in importance, but slight shifts in the ratios of abundant microbe



populations have been thought to have roles in obesity, diabetes and inflammatory bowel diseases such as Crohn's disease.



Annah S. Rolig, a postdoctoral researcher at the University of Oregon, led a project that focused on the individual contributions of bacterial species in the guts of germ-free zebrafish. Her team found that an abundance of a species is not what drives immune response. Species in very low abundance can be the important regulators. Credit: University of Oregon

That thinking is now changing, Rolig said. "The contribution of each



bacterium is not equal. There is a per-capita effect that needs to be considered."

The keystone - an important participant that functions to regulate a healthy microbiota - may reside in low-abundant <u>bacterial species</u>. The research team found through additional scrutiny that these species secreted molecules - for now unidentified - that allowed them to dampen the <u>immune response</u> to the whole community.

"Now we've shown that these minor members can have a major impact. If we can identify these keystone species, and find that in a disease state one species may be missing, we might be able to go in with a specific probiotic to restore healthy functioning," said Rolig, who also is a scientist in the National Institutes of Health-funded Microbial Ecology and Theory of Animals Center for Systems Biology, known as the META Center, at the UO.

To develop a model to capture per-capita contributions of <u>microbes</u> in a population, Rolig and her co-authors—biology graduate student Adam R. Burns, microbiologist Brendan Bohannan of the Institute of Ecology and Evolution and biologist Karen Guillemin, director of the META Center—turned to UO physicist Raghuveer Parthasarathy. His math-driven model, detailed in the paper, provides formulas that predict collective inflammatory responses of combinations of bacteria.

"I'm really proud of this paper because it exemplifies an achievement of one of the major goals of the META Center for Systems Biology, namely to provide a predictive model of how host-microbe systems function," Guillemin said. "This experimental and modeling framework could be readily generalized to more complex systems such as humans, for example to predict disease severity in individuals with inflammatory bowel disease based on the pro-inflammatory capacity of their gut microbes as assayed in cell culture."



More information: *Cell Host & Microbe*, <u>dx.doi.org/10.1016/j.chom.2015.10.009</u>

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