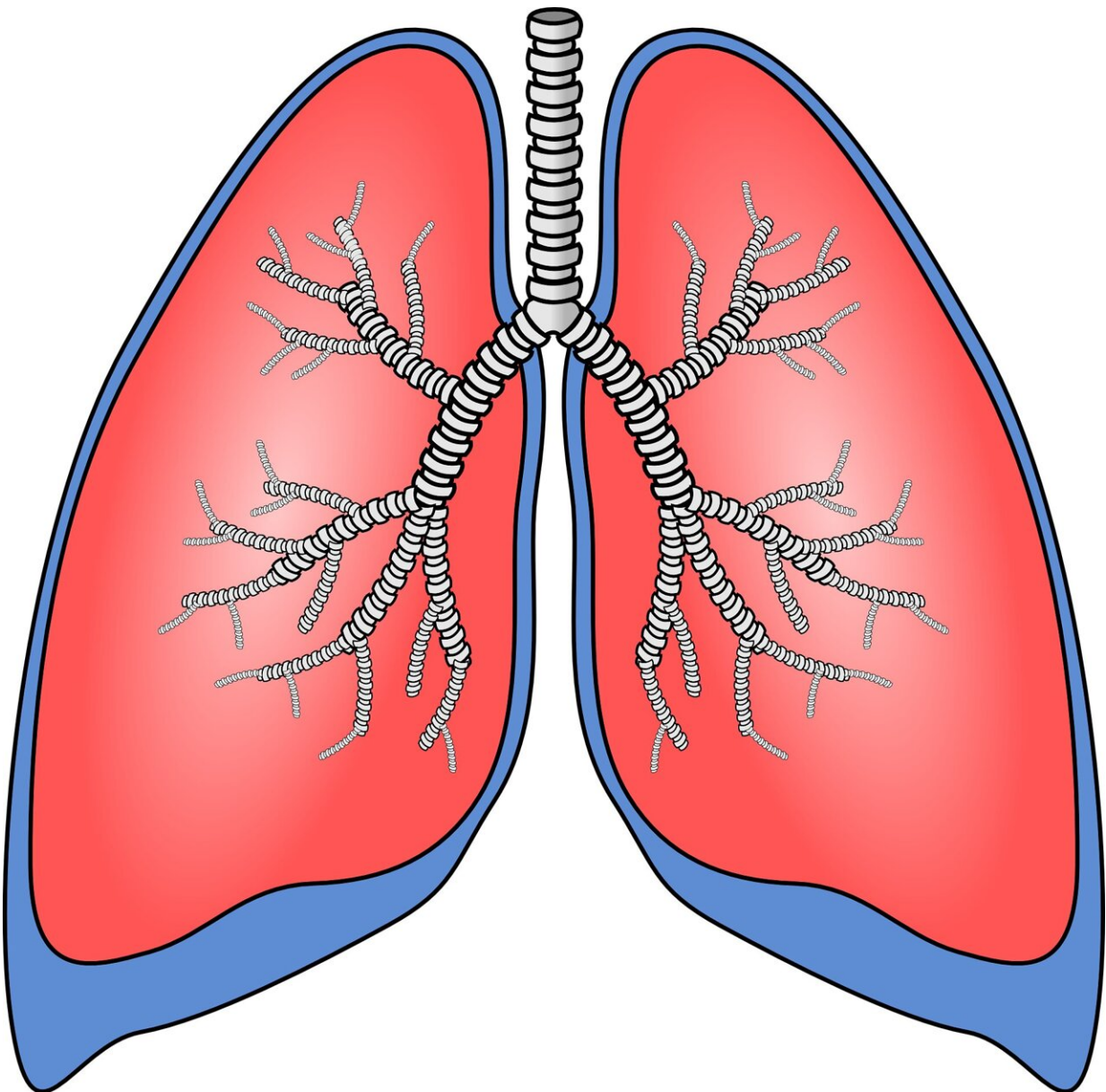


Zinc discovery holds promise for people with cystic fibrosis

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University of Queensland researchers have identified an opportunity to reduce infections in people living with cystic fibrosis (CF). Professor Matt Sweet, Dr. Kaustav Das Gupta and Dr. James Curson from UQ's Institute for Molecular Bioscience have discovered a fault in the bacteria-killing function of immune cells in people with CF and a potential way to get around it.

CF is a chronic disease in which defects in the CFTR ([cystic fibrosis](#) transmembrane conductance regulator) channel cause a build-up of mucus in the lungs, airways and digestive system, leading to recurring infections.

Professor Sweet said the team has found that in people with CF, immune cells called [macrophages](#) are defective in a zinc pathway that the body uses to kill bacteria. The study is [published](#) in the journal *Proceedings of the National Academy of Sciences*.

"One way that macrophages destroy bacteria is by poisoning them with toxic levels of metals such as zinc," Professor Sweet said.

"We discovered that the CFTR ion channel is crucial to the zinc pathway and because it doesn't work properly in people with CF, it may partly explain why they're more susceptible to bacterial infections."

Importantly, the researchers also identified a zinc transport protein that can restore the macrophages' ability to kill bacteria when the CFTR protein is not working.

"Our goal now is to deliver this [zinc](#) transport protein to macrophages in

people with CF with the expectation that it would reactivate their immune response and reduce infections," Professor Sweet said.

Around 3,600 Australians live with cystic fibrosis, which can reduce life expectancy to an average of 47 years.

Professor Peter Sly at UQ's Child Health Research Center, a pediatric respiratory physician and key collaborator on the project, said discovering more about how CF affects the immune system is key to patient care.

"People with CF have a hyper inflammatory state in their airways and are very susceptible to bacterial infections but frequent treatment with antibiotics can often lead to antibiotic-resistant infections," Professor Sly said.

"Current treatments can restore many aspects of CFTR function but they don't resolve or prevent lung infections so there is a need to restore immune functions."

The study was completed in collaboration with Professor Mark Schembri from IMB.

More information: Kaustav Das Gupta et al, CFTR is required for zinc-mediated antibacterial defense in human macrophages, *Proceedings of the National Academy of Sciences* (2024). [DOI: 10.1073/pnas.2315190121](https://doi.org/10.1073/pnas.2315190121)

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