

Immune discovery could aid vaccine development for common cause of sore throat





Human challenge with Streptococcus pyogenes. Schematic of Streptococcus pyogenes human-challenge model. Details participant demographics, timepoints, outcomes and analyses performed. Figure created with BioRender.com. Credit: *Nature Communications* (2022). DOI: 10.1038/s41467-022-28335-3

Researchers have made a discovery that could aid in the development of



a vaccine for a common cause of children's sore throats.

The research, led by the Murdoch Children's Research Institute (MCRI), and published in *Nature Communications*, has identified an immune system signature associated with acute pharyngitis (<u>strep throat</u>).

MCRI researchers Dr. Joshua Osowicki, Professor Andrew Steer, Associate Professor Paul Licciardi, Associate Professor Daniel Pellicci, contributed to the study, which aimed to advance <u>vaccine research</u> into Strep A by better understanding the immune response to the bacteria.

Strep A causes at least 750 million infections and more than 500,000 deaths each year. The bacterium is the most frequent cause of pharyngitis (sore throat) and a very common cause of skin infections in children. These infections can lead to dangerous complications such as kidney damage, rheumatic fever and rheumatic heart disease.

Rheumatic fever and kidney inflammation after Strep A infections particularly affect school-aged children and causes accelerated <u>heart</u> <u>disease</u> and kidney disease in early adulthood.

"Strep A disease disproportionately affects young children, pregnant women and the elderly," Dr. Osowicki said. It's one of the most common reasons for a healthy child to be critically ill in ICU with an <u>infection</u> 24 hours later."

"No <u>vaccine</u> is currently available for Strep A, and greater efforts are required to develop new vaccines to prevent serious disease and death caused by this bacterium."

For the initial study, the researchers undertook comprehensive immune profiling of the early immune response in blood and saliva from 25 healthy adults who were challenged with Strep A (painted onto the back



of their throat), 19 of whom subsequently developed acute symptomatic pharyngitis.

Associate Professor Pellicci said they were able to identify an immune signature associated with acute pharyngitis, that would aid in the development of a future vaccine for Strep A.

"An immune signature is a marker in the blood that establishes whether a person has responded to an infection," he said. "We plan to look for this immune signature in people that receive vaccines to prevent Step A infections, which may fast-track research to identify effective vaccines that prevent serious diseases caused by this bacterium."

Professor Steer said critical knowledge gaps had hindered Strep A vaccine development but understanding the human immune response to the infection, via a human challenge study, could accelerate the development of effective vaccines.

A human challenge study is a type of clinical trial for a vaccine involving the intentional exposure of the test subject to the condition tested in a controlled environment. By understanding the immune response, researchers are then able to better understand what vaccines might be effective.

"As well as providing a platform to evaluate candidate vaccines, a humanchallenge model allows us to study the <u>immune response</u> to infection using samples collected before, during, and after development of disease," Associate Professor Paul Licciardi said.

Human-challenge models have been established for other infectious pathogens such as malaria and have enormous potential to successfully guide vaccine development.



More information: Jeremy Anderson et al, Immune signature of acute pharyngitis in a Streptococcus pyogenes human challenge trial, *Nature Communications* (2022). DOI: 10.1038/s41467-022-28335-3

Provided by Murdoch Children's Research Institute

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