

Typhoid vaccine may protect against other infections

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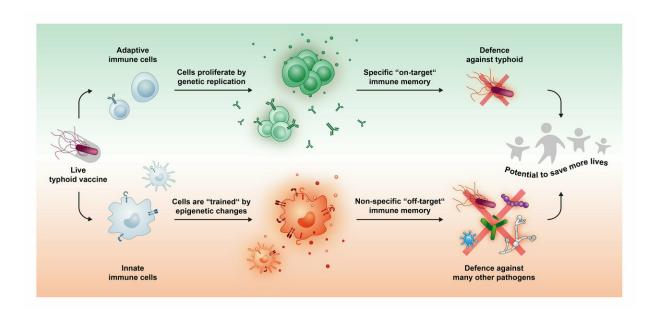


Illustration showing adaptive and innate immune responses to live typhoid vaccine. Credit: University of Liverpool/Liverpool School of Tropical Medicine

New research by the University of Liverpool and Liverpool School of Tropical Medicine shows that vaccination with weakened strains of *Salmonella* may also protect against other infections.

The researchers hope that the findings could impact vaccination strategy in the developing world, where infectious diseases are common and where broader <u>protection</u> could potentially save many lives.



Typhoid fever is a bacterial bloodstream infection caused by *Salmonella Typhi* that is estimated to affect between 11-18 million people and cause between than 128,000-190,000 deaths annually worldwide.

Published in the journal *Science Advances*, a new experimental study presents promising first data on the 'non-specific' immune response triggered by the live oral typhoid <u>vaccine</u> Ty21a.

"Live-attenuated *Salmonella* vaccines are low-cost, well-tolerated and easily administered. These vaccines could potentially be included in global vaccination programmes, not just for their impact on *Salmonella*, but also for their off-target, non-specific beneficial effects," says lead author Dr. Shaun Pennington from the Liverpool School of Tropical Medicine.

Previous evidence has suggested that some live-attenuated vaccines, such as those for measles and polio, can stimulate the human <u>immune system</u> to generate a wider protective response and lower all-cause mortality. In order to investigate whether *Salmonella* vaccines might offer similar protection, the researchers vaccinated a small group 16 healthy adults in the UK with the Ty21a vaccine and studied its impact on their immune system over the course of six months.

They looked at immune responses targeting *Salmonella* as well as those targeting a range of other pathogens. The changes they observed to levels of infection-fighting <u>white blood cells</u> (monocytes) and immune system messengers (cytokines) suggest that Ty21a can strengthen the <u>immune</u> <u>response</u> against subsequent, unrelated infections.

"The next step is to observe whether these responses also occur in children in low-income settings where their impact would be greatest. We'd like to conduct further <u>clinical studies</u>, where we will be able to assess the wider impact of our observations in conferring protection



against other common infections, not just *Salmonella*," says Professor Melita Gordon from the University of Liverpool and Malawi-Liverpool-Wellcome Trust Clinical Research Programme, who was the study's principal investigator.

The researchers add that the ability to manipulate live-attenuated *Salmonella* so that they express components of other pathogens could make their findings particularly exciting for future 'vector vaccine' design.

"*Salmonella* vector vaccines could provide *Salmonella*-specific protection, vectored-pathogen protection and non-specific protection, making live-attenuated *Salmonella* a hugely powerful 'triple threat' tool for global vaccine development," adds Professor Gordon.

The paper 'Nonspecific effects of oral vaccination with live-attenuated *Salmonella Typhi* strain Ty21a' is published in *Science Advances*.

More information: S.H. Pennington el al., "Nonspecific effects of oral vaccination with live-attenuated Salmonella Typhi strain Ty21a," *Science Advances* (2019).

advances.sciencemag.org/content/5/2/eaau6849

Provided by University of Liverpool

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