



Background

Staphylococcus aureus is a commensal gram-positive bacterium which can cause a wide range of human diseases, including cellulitis, necrotizing fasciitis, pneumonia, infective endocarditis, bacteremia, and toxic shock syndrome. The ease with which *S. aureus* acquires antibiotic resistance means that treatment failure is increasingly common. The need for new clinical management approaches has significantly increased the investment and interest in alternate methods of treating and preventing *S. aureus* infections, including prophylactic vaccination.

Development of an effective vaccine for *S. aureus* has been a continuing effort over the past two decades, as yet with no success. The main strategy behind most vaccines for *S. aureus* that have failed in the clinic has been to elicit a humoral antibody response to clear the bacteria via opsonophagocytosis. It has become apparent that this approach **does not work**.

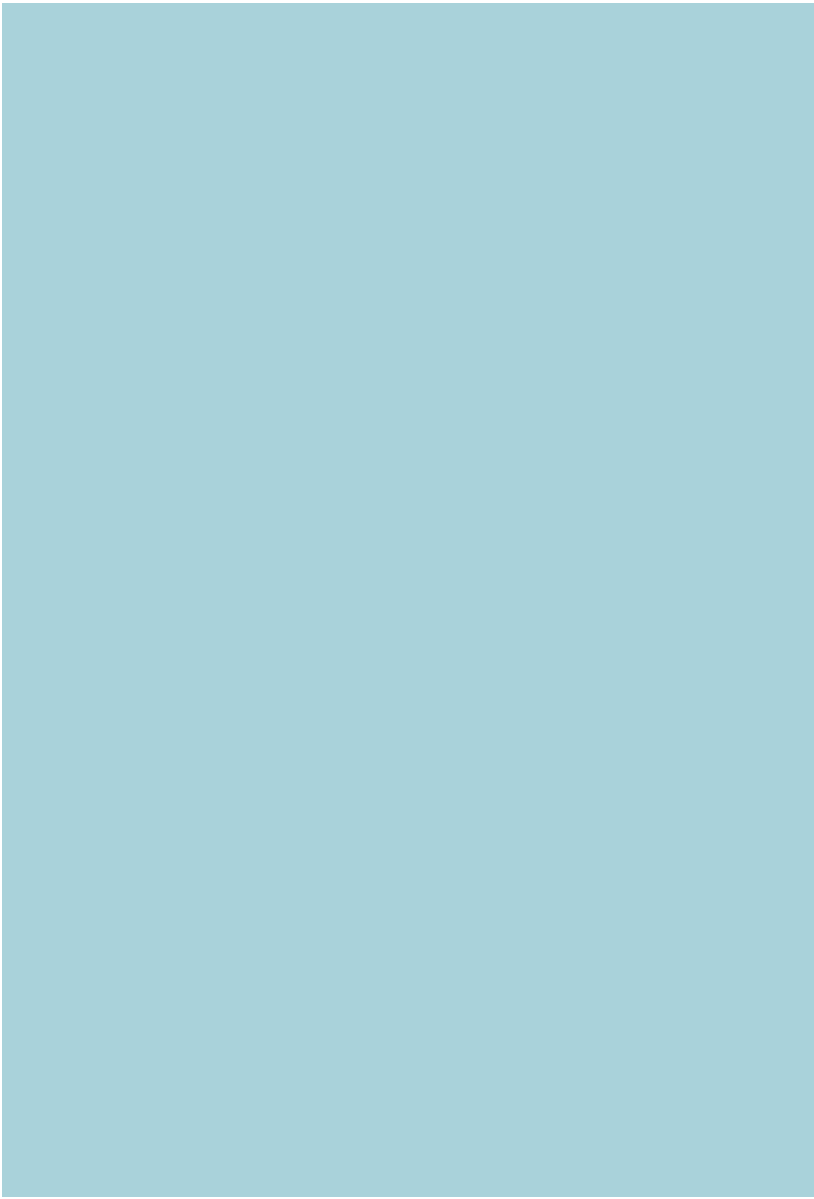
Technology

Our approach is to target staphylococcal superantigen-like (SSLs) proteins using a fusion protein comprising three key SSLs. SSLs are universally conserved virulence factors which are secreted by *S. aureus*. Our vaccine generates

neutralising antibodies which act on these secreted virulence factors, preventing *S. aureus* from evading the immune system and establishing infection.

Major advantages

- Targets secreted immune evasion factors – aims to neutralize staphylococcal virulence rather than providing sterilizing immunity.
- Focuses on virulence factors that are immune inhibiting (help the pathogen escape the immune



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