

Poster abstract submission

Approval Status

Not Started

Presenting author

Beckie Ingram

Presenting author's email

b.ingram@qub.ac.uk

Further authors (if any)

Rachael Bell
Julie Gormley

Affiliation(s)

VacTimmune

Country

UK

Type of organization

Industry / company

Poster titleA Dual-Use Vaccine Strategy Against *Pseudomonas aeruginosa***Poster abstract**

Pseudomonas aeruginosa is a WHO-designated priority antimicrobial-resistant pathogen and a major cause of chronic, life-limiting lung infection in people with cystic fibrosis, bronchiectasis and severe chronic obstructive pulmonary disease. Once established, infection is rarely cleared, even with repeated courses of high-dose antibiotics, driving progressive lung damage, recurrent hospitalisation and escalating antimicrobial resistance. Despite decades of effort, no licensed *P. aeruginosa* vaccine exists.

Here, we present a vaccine development strategy that challenges the prevailing assumption that antibacterial vaccines are exclusively prophylactic. Using a proprietary reverse-vaccinology pipeline, we identified conserved *P. aeruginosa* antigens and advanced a lead recombinant protein vaccine into biologically relevant preclinical infection models. Subcutaneous immunisation confers robust prophylactic protection against both systemic and mucosal infection, significantly reducing lung bacterial burden in wild-type mice and in a β ENaC transgenic model that recapitulates key features of cystic fibrosis lung disease.

Critically, we demonstrate that the same vaccine also disrupts established chronic infection. In a chronic lung infection model, vaccination administered after infection is established results in a marked reduction in bacterial burden following booster immunisation, providing clear evidence of intrinsic therapeutic activity. This dual prophylactic–therapeutic efficacy distinguishes our approach from historical vaccine strategies and supports a paradigm shift in antibacterial vaccinology.

The programme is now advancing toward translational development, including manufacturing scale-up, formulation optimisation and regulatory-aligned preclinical studies to support first-in-human evaluation. Together, these data support the progression of a first-in-class anti-*P. aeruginosa* vaccine capable of both preventing infection and reducing established disease burden, directly addressing a critical unmet need in AMR and chronic respiratory infection.

Research topic

Vaccines

If you wish to submit a graphic with your abstract you can upload it here.

