Controlled human infection with *Neisseria lactamica* in late pregnancy to measure horizontal transmission and microbiome changes in mother–neonate pairs: a single-arm interventional pilot study protocol

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**ABSTRACT**

**Introduction** Infant upper respiratory microbiota are derived partly from the maternal respiratory tract, and certain microbiota are associated with altered risk of infections and respiratory disease. *Neisseria lactamica* is a common pharyngeal commensal in young children and is associated with reduced carriage and invasive disease by *Neisseria meningitidis*. Nasal inoculation with *N. lactamica* safely and reproducibly reduces *N. meningitidis* colonisation in healthy adults. We propose nasal inoculation of pregnant women with *N. lactamica* to establish if neonatal pharyngeal colonisation occurs after birth, and to characterise microbiome evolution in mother–infant pairs over 1 month post partum.

**Methods and analysis** 20 healthy pregnant women will receive nasal inoculation with *N. lactamica* (wild type strain Y92-1009) at 36–38 weeks gestation. Upper respiratory samples, as well as optional breastmilk, umbilical cord blood and infant venous blood samples, will be collected from mother–infant pairs over 1 month post partum. We will assess safety, *N. lactamica* colonisation (by targeted PCR) and longitudinal microevolution (by whole genome sequencing), and microbiome evolution (by 16S rRNA gene sequencing).

**Ethics and dissemination** This study has been approved by the London Central Research Ethics Committee (21/PR/0373). Findings will be published in peer-reviewed open-access journals as soon as possible.

**Trial registration number** NCT04784845.

**INTRODUCTION**

Upper respiratory pathobionts and infant disease

Upper respiratory tract (URT) pathobionts are common colonising bacteria capable of causing disease in immunocompetent individuals (eg, *Neisseria meningitidis*, *Streptococcus pneumoniae*, *Haemophilus influenzae*). Although harmless in many hosts, these remain a leading cause of global childhood death and disease, especially in the first year of life: globally, 492 000 and 89 000 infants die each year due to respiratory tract infection (RTI) and meningitis, respectively.1 While many infant infections are vaccine-preventable, global access to and acceptance of vaccines remains incomplete, such that 1.5 million vaccine-preventable childhood deaths occur each year.2 Furthermore, existing vaccines do not protect against all important childhood respiratory pathogens (such as respiratory syncytial virus and some serotypes of *S. pneumoniae*), and vaccinations are not routinely given in the first 2 months post partum, when risk of invasive infection is highest. Rising global antimicrobial resistance compounds these challenges, with 214 000 neonates dying of resistant infections each year.3

Recent research has highlighted the relationship between URT pathobionts, childhood disease and the evolving infant
microbiome (the site-specific total microbial community). Taxonomic profiling (eg, by bacterial 16S rRNA gene sequencing) demonstrates distinct URT microbiota during acute RTI, and that URT microbiome perturbation is associated with RTI severity and recurrence, childhood chronic wheeze and early allergic sensitisation.

Infant RTIs are often preceded by micro changes, including increased relative abundance of URT pathobionts and loss of topography (blurring of distinction between microbiomes at adjacent anatomical sites), suggesting a possible causal role in developing infection. However, as up to 95% of infants are colonised with at least one URT pathobiont at any time, the presence of pathobionts may indicate a state of dysbiosis (microbial dysregulation), in which disease-causing bacteria and viruses can cause infection in a previously resilient ecological niche.

Longitudinal cohort microbiome studies suggest that infant URT flora are acquired at least in part from their mothers: there is greater overlap between a neonate’s oral flora and that of its own mother than unrelated mothers, and shared bacterial strains (detected by whole genome sequencing) are more long-lived in an infant’s mouth than non-maternal strains. Moreover, a subset of maternal oral strains account for a disproportionately large share of their infants’ oral flora.

Can upper respiratory microbiota be manipulated to prevent infant disease?
In the future, it is possible that simple methods of modifying the infant nasopharyngeal microbiome could provide useful adjunct protection in the early months of life. Previously, neonatal nasal inoculation with low-virulence *Staphylococcus aureus* strain 502A has been shown to reduce colonisation by high-virulence *S. aureus* strain 80/81 in response to an outbreak, while pharyngeal inoculation with strain 215 alpha-haemolytic streptococci reduced pharyngeal pathobionts (*Escherichia coli, Klebsiella pneumoniae, S. aureus*) in neonatal intensive care inpatients. Furthermore, a variety of probiotics (living microorganisms administered to confer a health benefit) have been trialled in children and adults, with some reporting a reduction in upper RTI incidence and duration. However, high quality efficacy and microbiome data is lacking, with significant inter-study variation in the bacterial strain (usually lactobacilli, streptococci and bifidobacteria), preparation (usually capsules or dairy supplement), dose and regimen. That being said, there is evidence that many probiotics are safe in pregnancy, neonates and lactating mothers, with some already available over-the-counter to these groups.

Controlled human infection with *Neisseria lactamica*
Unlike probiotic studies (which often employ proprietary food supplements with uncontrolled concentrations of bacteria), controlled human infection offers a robust model for studying the impact of a defined bacterial inoculum on a human host. Our research group has previously developed a safe and reliable controlled human infection model using nasal inoculation with $10^5–10^6$ colony-forming units (CFU) *N. lactamica* strain Y92-1009 in healthy adults. In randomised blinded placebo-controlled studies, we have characterised *N. lactamica* colonisation density, duration, cellular and humoral immune responses, and genomic microevolution, observing colonisation rates of up to 85% and no serious adverse reactions following over 400 inoculations.

*N. lactamica* is a common non-capsulated pharyngeal commensal that does not cause invasive disease in immunocompetent hosts. Colonisation peaks at over 40% in 1–2-year-old children, before falling to less than 10% in adulthood, and *N. lactamica* carriage correlates inversely with *N. meningitidis* colonisation. Indeed, *N. lactamica* controlled human infection in healthy adults reduced naturally-acquired *N. meningitidis* carriage from 18% to 8% (both by displacing existing colonisation and preventing new acquisition), exceeding the typical impact of glycoconjugate ACWY vaccination. This displacement effect is not specific to a particular serogroup, unlike available glycoconjugate meningococcal vaccines. Furthermore, natural colonisation with *N. lactamica* has been observed to correlate inversely with invasive meningococcal disease.

**Rationale for this study**
The aim of this study is to establish if nasal inoculation with *N. lactamica* in pregnancy results in neonatal colonisation after birth, and to characterise the impact on the developing neonatal URT microbiome. If successful, this will provide a novel model for inducing and studying person-to-person commensal acquisition in neonates (unlike traditional controlled human infection models, which capture inoculation-induced colonisation). Looking ahead, this may provide a new strategy for reducing *N. meningitidis* colonisation and disease in neonates, and of modifying neonatal microbiome development with a view to improving health outcomes.

METHODS AND ANALYSIS

**Study overview**
In this prospective controlled human infection study, 20 healthy pregnant women will receive nasal inoculation with $10^5$ CFU *N. lactamica* (wild type strain Y92-1009) at 36+0 to 37+6 weeks gestation (figure 1). Upper respiratory samples and breastmilk will be collected from mother–infant pairs 1 day, 1 week and 1 month post partum, as well as an umbilical cord blood sample at delivery and an infant venous blood sample at 1 month post partum. Adverse events and relevant clinical data (including mode of delivery, infant feeding and use of antimicrobials) will be recorded. Samples will be analysed by microbiological culture, targeted PCR and sequencing to determine *N. lactamica* Y92-1009 presence, colonisation density and longitudinal microevolution, as well as 16S rRNA gene
sequencing to characterise microbiome evolution in mother–neonate pairs.

**Study hypothesis, objective and endpoints**

We hypothesise that nasal inoculation of pregnant women with $10^5$ CFU *N. lactamica* strain Y92-1009 will result in maternal upper respiratory *N. lactamica* colonisation; and that horizontal transmission of upper respiratory *N. lactamica* will occur from experimentally-colonised women to their infants by 4 weeks post partum.

This study aims to investigate this hypothesis, and to characterise the safety, colonisation kinetics and microbiome impact of nasal *N. lactamica* inoculation in pregnancy on mother–neonate pairs (table 1).

**N. lactamica inoculum**

Vials of lyophilised (freeze-dried) *N. lactamica* (LyoNlac) will be produced and stored under Good Manufacturing Practice (GMP)-like standards at the University of Southampton. The strain used for lyophilisation (Y92-1009, sequence type 3493, clonal complex 613) originates from the current GMP-compliant pharmaceutical manufacturing facilities at Public Health England (Porton Down, UK). Lyophilised inoculum stocks can be stored

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**Figure 1** Study overview flowchart. API, analytical profile index; CFU, colony forming units; h, hours; m, months; MALDI, matrix-assisted laser desorption/ionisation; Nlac, *Neisseria lactamica*; w, weeks.

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**Table 1** Primary, secondary and safety study endpoints

<table>
<thead>
<tr>
<th>Primary endpoint</th>
<th>Secondary endpoints</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Confirmation of neonatal (0–31 days) <em>Neisseria lactamica</em> colonisation by selective culture of nasopharyngeal and saliva samples.</td>
<td>1. Confirmation of neonatal <em>N. lactamica</em> Y92-1009 colonisation by strain-specific PCR.</td>
</tr>
<tr>
<td>Safety endpoints</td>
<td>2. <em>N. lactamica</em> colonisation density in inoculated volunteers compared with their infants across study time points and sample niche.</td>
</tr>
<tr>
<td>1. Percentage of inoculated volunteers with adverse reactions or serious adverse events within the study period.</td>
<td>3. Microbiome composition across study time points and sample niche in inoculated volunteers compared with their infants, and in colonised compared with uncolonised infants.</td>
</tr>
<tr>
<td>2. Percentage of neonates with serious adverse events within the study period.</td>
<td>4. <em>N. lactamica</em> genome sequence for isolates derived from inoculated volunteers compared with their infants across study time points and sample niche.</td>
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at 4°C (unlike frozen stocks requiring storage at −80°C). LyoNlac will be reconstituted in phosphate buffered saline and diluted to a concentration of 10^5 CFU/mL for inoculation. In our previous work, this dose resulted in _N. lactamica_ colonisation in 100% of 10 adults receiving lyophilised inoculum and 85% of 20 adults using frozen inoculum (ie, 90% overall). The inoculum will be administered immediately following reconstitution and serial dilution, and after no more than 30 min delay; recovery of viable bacteria remains over 70% after 30 min incubation in room air (unpublished data).

**Recruitment of study volunteers**

All potentially eligible healthy adult volunteers in the second and third trimesters of pregnancy will be identified by research midwives using a clinical database, and will be sent an ethically-approved study advertisement and a Participant Information Sheet. Individuals who express an interest will be invited to attend a screening visit. Participants will be offered reimbursement for their time, travel and inconvenience.

**Study visits and procedures**

**Screening**

Interested individuals will be reviewed by the study doctor at a screening visit (34+0 to 36+6 weeks gestation), to discuss the study timeline and procedures (table 2) and assess eligibility (box 1). Eligible individuals who provide informed consent at screening will be considered enrolled. A detailed medical history will be taken, followed by pharyngeal and saliva sampling to assess for _N. lactamica_ carriage; enrolled volunteers who are already naturally colonised with _N. lactamica_ will not be inoculated. However, they will be invited to remain in the study as an opportunistic comparison cohort, and will complete all postpartum visits and sampling as for the inoculated mother–neonate pairs (table 2).

**Inoculation**

Eligible consenting volunteers who are not already naturally colonised with _N. lactamica_ will receive nasal inoculation with 10^5 CFU _N. lactamica_ Y92-1009, using a sterile pipette (0.5 mL inoculum per nostril).

**Postpartum follow-up**

Mother–neonate pairs will be reviewed 1 day, 1 week and 1 month post partum. At each visit, maternal consent for neonatal sampling will be re-affirmed, and respiratory samples will be collected (table 2). An optional umbilical cord blood sample (visit 3) and neonatal venous blood sample (visit 5) will be collected, as well as relevant clinical data (including mode of delivery, infant feeding and use of antimicrobials). Optional breastmilk samples will also be collected at each postpartum visit (up to 5 mL each at visits 3 to 5). Volunteers may decline optional blood and breastmilk sampling without affecting their enrolment in or completion of the rest of the study.

Volunteers’ experiences of participation (including acceptability and tolerability of inoculation and maternal and infant sampling) will be captured using optional questionnaires, to help us identify areas for improving volunteer experience in current and future studies.

**Optional visit 6 (15+/-2 weeks post partum)**

At the time of writing, a study amendment is in progress to include an optional study visit at 15±2 weeks post partum (visit 6). This visit will involve maternal and infant pharyngeal, saliva and venous blood sampling, as well as saliva and oropharyngeal swabs from household contacts under 5 years old. In a recent _N. lactamica_ controlled human infection study involving non-pregnant adults, anti- _N. lactamica_ immunoglobulin G (IgG) (peak titre and titre increment) correlated inversely with subsequent pharyngeal colonisation density at 28 days post-inoculation (unpublished data), suggesting that IgG may play a role in controlling _N. lactamica_ carriage. Total IgG is up to 1.5 times more concentrated in umbilical cord blood compared with maternal blood, and the half-life of transplacental maternal IgG is approximately 30 days; thus, infant anti- _N. lactamica_ IgG is predicted to be less than 10%–20% that of maternal IgG by 15±2 weeks (3 to 4 half-lives). Visit 6 therefore offers a valuable opportunity to investigate the association between anti- _N. lactamica_ IgG and _N. lactamica_ carriage in mother–infant pairs.

**Sample processing**

_N. lactamica_ identification

Working in a class 2 microbiological safety cabinet in a Containment Level 2 laboratory, all respiratory samples will be suspended in 1 mL storage medium (10% glycerol in 0.1% diethylpyrocarbonate (DEPC)-treated water). Agar selective for _N. lactamica_ (GC agar supplemented with 5-bromo-4-chloro-3-indolyl-β-D-galactopyranoside (X-gal); Thermo Fisher Scientific, Basingstoke, UK) will be inoculated with sample solution, and all remaining sample solution will be split into 200 µL aliquots for storage in a secure, temperature-monitored −80°C freezer. Following 24–48 hours incubation, colonies with typical _N. lactamica_ morphology (blue on GC X-gal agar) will undergo identification by oxidase reaction, Gram stain, microscopy and analytical profile index (API-NH, bioMérieux, Lyon, France) or matrix-assisted laser desorption/ionisation time-of-flight (MALDI-TOF; Bruker, Coventry, UK). _N. lactamica_ density will be quantified and isolates will be stored at −80°C. Breastmilk will also be plated onto selective agar to examine for evidence of _N. lactamica_, and residual samples will be stored at −80°C for microbiome analysis.

_N. lactamica_ Y92-1009 strain-specific PCR

We have identified two chromosomal loci present in _N. lactamica_ strain Y92-1009 that, when amplified in conventional multiplex PCR and analysed by agarose gel electrophoresis, produce amplicons with a characteristic band pattern. Using the same multiplex PCR on isolates of naturally circulating strains of _N. lactamica_, this band pattern has not been reproduced. A third locus, present...
## Table 2 Study visits and activities

<table>
<thead>
<tr>
<th>Timeline</th>
<th>Screening (visit 1)</th>
<th>Inoculation (visit 2)</th>
<th>Birth (visit 3)</th>
<th>Follow-up (visits 4 and 5)</th>
<th>Optional (visit 6)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>34+0 to 36+6 weeks gestation</td>
<td>36+0 to 37+6 weeks gestation</td>
<td>0-24 hours post partum</td>
<td>7±3 days post partum</td>
<td>28±3 days post partum</td>
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<td>Participant Information Sheet and consent form</td>
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<td>Clinical review</td>
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<tr>
<td><em>Neisseria lactamica</em> nasal inoculation</td>
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<td>(Unless already colonised)</td>
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<tr>
<td>Nasopharyngeal swabs (mother)</td>
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<tr>
<td>Oropharyngeal swabs (mother)</td>
<td>+</td>
<td>+</td>
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<td>Saliva swabs (mother)</td>
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<td>Nasopharyngeal swabs (infant)</td>
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<td>Saliva swabs (infant)</td>
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<tr>
<td>Umbilical cord blood</td>
<td>Optional</td>
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<tr>
<td>Infant venous blood (2 mL)</td>
<td>Optional</td>
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<td>Optional</td>
<td>Optional</td>
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<tr>
<td>Maternal breast milk</td>
<td>Optional</td>
<td>Optional</td>
<td>Optional</td>
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<tr>
<td>Maternal venous blood (5 mL)</td>
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<tr>
<td>Oropharyngeal and saliva swabs (under 5 years)</td>
<td>Optional</td>
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</table>
Inflammation Research for 16S rRNA gene sequencing.

Transported to the University of Edinburgh Centre for Paired mother–infant frozen sample aliquots will be Microbiome analysis data).

**bacterial lysate as a template, with an optimum annealing assay has been validated using genomic DNA and boiled temperature 65°C and extension time 15 s (unpublished differences in **N. lactamica** microevolution in mother–infant pairs. Resulting sequences will be mapped to a closed **N. lactamica** Y92-1009 reference genome for comparison.**

**Sample size**

To our knowledge, this proof-of-concept represents the first respiratory controlled human infection study in pregnancy. As such, it is not possible to accurately define the sample size needed to conclusively rule out induced horizontal transfer of **N. lactamica**. A pragmatic sample size of 20 inoculated volunteers has been selected: based on our earlier work, **N. lactamica** colonisation is expected in 18 (90%) of volunteers inoculated with **10^5** CFU **N. lactamica**. This sample would therefore fail to detect horizontal transfer only if it occurs in less than 1 in 18 infants (<6%), in which case it would be impractical and even unethical to pursue horizontal **N. lactamica** controlled human infection as a method of studying and manipulating infant commensalisation. By 1 month of age, 2.5% of infants naturally carry **N. lactamica**. Based on a plot of lower confidence limits against observed infant **N. lactamica** colonisation for a sample of 18 experimentally-colonised volunteers, infant **N. lactamica** colonisation greater than 22% (95% CI (2.3% to 41.7%)) would demonstrate that inoculation-induced carriage exceeds natural **N. lactamica** carriage (figure 2).

We expect to screen at least 24 volunteers to reach our target of 20 inoculations, allowing for 10% dropout and 10% baseline natural **N. lactamica** carriage. It is not clear how maternal or neonatal antibiotics may influence **N. lactamica** carriage and microbiota. At the Principal Investigator’s discretion, an additional volunteer may be enrolled and inoculated for every volunteer (or

**Microbiome analysis**

Paired mother–infant frozen sample aliquots will be transported to the University of Edinburgh Centre for Inflammation Research for 16S rRNA gene sequencing. Resulting sequence reads will undergo taxonomic classification to compare mothers and infants, and (assuming some neonates become colonised with **N. lactamica** colonised and uncolonised infants, across different sample niches and study time points.

**N. lactamica sequencing**

If any neonates become colonised with **N. lactamica**, paired mother–infant **N. lactamica** isolates will undergo whole genome sequencing to assess for differences in **N. lactamica** microevolution in mother–infant pairs.

**Figure 2** Lower confidence limit plot for inoculation-induced *Neisseria lactamica* (Nlac) colonisation.

![Figure 2](http://bmjopen.bmj.com/)

neonate) who receives antibiotics after inoculation, up to a maximum of 10 additional volunteers.

Statistical analysis
The primary endpoint will be expressed as a proportion N/M, where:

- M=Number of mothers colonised with *N. lactamica*.
- N=Number of neonates (aged 0–31 days) colonised with the same strain of *N. lactamica* as their own mother.
- ‘Colonised’=evidence of *N. lactamica* by culture of upper respiratory swabs (using selective media and confirmation by API or MALDI-TOF) on at least one visit.
- ‘Same strain’=confirmed using Y92-specific *N. lactamica* PCR for inoculated volunteers, or whole genome sequencing for volunteers already naturally carrying *N. lactamica* at screening.

Subgroup analysis will be performed, to compare inoculated mother–infant pairs with naturally colonised mother–infant pairs, and to compare pairs where the mother or infant received antibiotics with pairs not receiving antibiotics.

Specific bioinformatic analysis strategies will depend on the quantity and quality of genomic data produced during the study, and on the rates of *N. lactamica* colonisation observed. Broadly speaking, microbiome bioinformatic pipelines (eg, QIIME2 and R packages) will be used to perform taxonomic classification, calculation of alpha and beta diversity and data visualisation.

Patient and public involvement
During early study design, an interview-based patient and public involvement project was conducted with 12 pregnant women (unpublished data). All 12 women expressed approval that the proposed study should go ahead, 11 of whom would personally have liked to participate. These discussions led to changes in study design (such as offering follow-up visits in the volunteer’s own home, and obtaining an umbilical cord rather than an infant venous blood sample at birth). During participant recruitment following study completion, the University of Southampton Public Engagement Research Unit will assist in showcasing the study at public engagement events.

ETHICS AND DISSEMINATION
Ethics and safety
Volunteer safety and ethical study conduct is of paramount importance. This study has been approved by the NHS Research Ethics committee and the Health Research Authority, and is sponsored by the University of Southampton. An External Safety Committee has been appointed to provide independent safety oversight. Safety of *N. lactamica* Y92-1009 inoculum is well-characterised: over 400 adults have previously been inoculated with this *N. lactamica* strain (of whom 30 received lyophilised inoculum, and 54 received 10⁵ CFU), with no serious adverse reactions reported to date (unpublished data). Due to the use of yeast-based and soya-based products in inoculum production, individuals reporting allergies to yeast or soya will not be eligible for enrolment. Inoculation will take place in a clinical area with access to adult, obstetric and paediatric emergency services. Volunteers will be reviewed by a study doctor immediately prior to and 30 min after inoculation.

Adverse events will be recorded throughout the study, and volunteers will be encouraged to report any concerns (including by telephone to the study doctor out-of-hours). Adverse events will also be recorded for any inoculated withdrawn volunteers, by telephone review at 4 weeks post partum. As per the current Good Clinical Practice guideline, adverse events are any untoward medical occurrence in a volunteer or their neonate, temporally, but not necessarily causally, related to study participation. Delivery of routine peripartum care is not considered an adverse event. Serious adverse events include hospitalisation (or prolongation of hospitalisation for routine peripartum care), persistent or significant disability, a life-threatening occurrence or death from any cause.

Mitigation strategies are in place to minimise the risk of SARS-CoV2 transmission to participants and study staff. Where possible, study visits will be conducted in the volunteer’s home, practicing social distancing and appropriate personal protective equipment for non-aerosol generating procedures. If participants report symptoms of or contact with anyone with suspected or confirmed SARS-CoV2 infection, study visits will be rescheduled until testing or self-isolation as necessary.

Data management and dissemination
All data will be stored in link-anonymised format using paper case report forms stored securely at the research site. Fully anonymised sequencing data will be uploaded to a curated online data repository. Research findings will be published in peer-reviewed journals as soon as is practicable, with an online link to the final approved study protocol. Participants will be provided with a lay summary of the study results once available.

Potential benefits and study impact
To our knowledge, this is the first respiratory controlled human infection study in pregnancy, and, if successful, would be the first to demonstrate induced person-to-person commensal transmission. This novel model could be used to characterise microbiome and immunological changes associated with induced commensalisation. Inducing *N. lactamica* colonisation in neonates may also provide a new strategy for reducing *N. meningitidis* colonisation and even disease in infants. Although many neonates are protected by trans-placental maternal antibodies, invasive meningococcal disease does rarely occur in infants, especially before vaccination. Looking ahead, improved understanding of microbiome evolution following controlled human infection is necessary before
inoculation in neonates (rather than in pregnant adults) can be considered.

Our team have recently trialled nasal inoculation in healthy non-pregnant adults using genetically-modified (GM) *N. lactamica* that expresses the meningococcal antigen *Neisseria* adhesin A (NadA). This GM inoculum has proved safe and effective at eliciting immune responses to NadA, providing a proof-of-concept that a harmless commensal may be used as a vehicle for prolonged mucosal exposure to an antigen of interest. Looking ahead, GM *N. lactamica* expressing antigens from other clinically-important paediatric respiratory pathogens could be explored, to complement existing childhood vaccination.

**REFERENCES**