Methicillin-resistant Staphylococcus aureus (MRSA) in nursing homes: can an improvement in infection control practices decrease MRSA prevalence?

| Recruitment status | Prospectively registered | | |
|----------------------|---|--|--|
| No longer recruiting | ☐ Protocol | | |
| Overall study status | Statistical analysis plan | | |
| Completed | [X] Results | | |
| Condition category | Individual participant data | | |
| | No longer recruiting Overall study status Completed | | |

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number EAT/2953/04

Study information

Scientific Title

Study objectives

- 1. What is the prevalence of MRSA in nursing homes in the Northern Health and Social Services Board?
- 2. Can an improvement in infection control standards result in a decrease in MRSA prevalence in nursing homes in this board area?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethical approval granted by the Office for Research Ethics Committees Northern Ireland (ORECNI) on 03/11/2005, reference number: 05/NIR03/154

Study design

Phase 1 is observational; Phase 2 is a cluster randomised controlled trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Methicillin-resistant Staphyloccus aureus (MRSA) colonisation

Interventions

Nasal and skin swab and urinary catheter sample where appropriate from residents in both phases 1 and 2. In phase 2, homes assigned to the intervention group will receive an audit of current infection control practices followed by infection control training and feedback. In control homes, routine care will continue as normal.

Intervention Type

Other

Phase

Phase I/II

Primary outcome(s)

In phase 1, determination of MRSA prevalence in nursing homes in Northern Health and Social Services (NHSSB). In phase 2, the prevalence of MRSA in intervention homes compared to control homes following infection control intervention, audit and feedback.

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/11/2008

Eligibility

Key inclusion criteria

Phase 1: all nursing homes in the Northern Board will be asked to participate. Within these homes, all residents and staff will be invited to participate.

Phase 2: For a nursing home to be considered in this phase, it must have at least 20 consenting adults.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Senior

Sex

All

Key exclusion criteria

Nursing home residents who are terminally ill, in a coma or not deemed fit to personally give consent and have no next-of-kin available to give consent, will be excluded from both phases of the trial. For staff, there are no specific exclusion criteria, other than failure to provide written informed consent.

Date of first enrolment

01/11/2005

Date of final enrolment

01/11/2008

Locations

Countries of recruitment

United Kingdom

Northern Ireland

Study participating centre Queen's University Belfast

Belfast United Kingdom BT9 7BL

Sponsor information

Organisation

Queens University Belfast (UK)

ROR

https://ror.org/00hswnk62

Funder(s)

Funder type

Government

Funder Name

Central Services Agency, a part of the Northern Ireland Health and Social Services (NIHSS), Reasearch and Development Office (EAT/2953/04)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

| Output type | Details | Date created Date adde | d Peer reviewed | ? Patient-facing? |
|-------------------------------|-------------------------------|------------------------|-----------------|-------------------|
| Results article | results | 01/09/2010 | Yes | No |
| Participant information sheet | Participant information sheet | 11/11/2025 11/11/202 | 5 No | Yes |