

AIR-NET- Testing anti-inflammatories for the treatment of bronchiectasis

Submission date 02/08/2024	Recruitment status Recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 23/10/2024	Overall study status Ongoing	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 24/02/2026	Condition category Respiratory	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

AIR-NET- Testing anti-inflammatories for the treatment of bronchiectasis

Who can participate?

Adults over 18 years, with bronchiectasis.

What does the study involve?

Not provided at time of registration

What are the possible benefits and risks of participating?

Benefits:

Not provided at time of registration

Risks:

This trial is categorised as: Type B, Somewhat higher than the risk of standard medical care. The trial treatments are all repurposed drugs already approved for use in conditions other than bronchiectasis.

Arm 1: Usual care

Arm 2: Disulfiram

Disulfiram is an alcohol deterrent compound licensed for use as an adjuvant in the treatment of drinking problems. The most common adverse reaction (AR) reported during treatment is alcohol reaction. Disulfiram–ethanol reactions often develop within 15 minutes after exposure to ethanol; symptoms usually peak within 30 minutes to 1 hour, and then gradually subside over the next few hours. Symptoms may be severe and life-threatening. Participants will be informed of the reaction risk and must agree to abstain from alcohol during treatment and for up to 14 days after discontinuation.

Arm 3: Dipyridamole

Dipyridamole is licensed for use in secondary prevention of ischaemic stroke and transient ischaemic attacks and as an adjunct to oral anti-coagulation for prophylaxis of thromboembolism associated with prosthetic heart valves. The most commonly reported adverse effects are headache, dizziness, diarrhoea, nausea (very common, $\geq 1/10$) and angina pectoris, vomiting, rash, myalgia (common, $\geq 1/100 < 1/10$).

Arm 4: Doxycycline

Doxycycline is licensed for use in the treatment of a variety of infections caused by susceptible strains of Gram-positive and Gram-negative bacteria and certain other micro-organisms. The most common adverse effects observed in patients receiving tetracyclines, including doxycycline, are hypersensitivity, headache, nausea/vomiting, photosensitivity reaction rash including maculopapular and erythematous rashes (common, $\geq 1/100 < 1/10$).

Adverse events, medications, exacerbations and vital signs will be reviewed at each visit. Regular safety bloods will be performed to assess full blood count, urea and electrolytes and liver function tests.

The data monitoring committee will perform a safety review of each active treatment arm. Each safety review will take place after 10 participants have completed treatment in the active arm.

Where is the study run from?
University of Dundee (UK)

When is the study starting and how long is it expected to run for?
July 2024 to February 2027

Who is funding the study?
LifeArc (UK)

Who is the main contact?
airnet-tm@dundee.ac.uk
Professor James Chalmers, j.chalmers@dundee.ac.uk

Contact information

Type(s)

Principal investigator

Contact name

Prof James Chalmers

Contact details

Ninewells Hospital
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DD1 9SY
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Type(s)

Public, Scientific

Contact name

Dr . Study Team

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Integrated Research Application System (IRAS)

1010124

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

1-027-24, CPMS 62256

Study information

Scientific Title

A randomised, open-label, multifactorial, multicentre, platform trial using a range of repurposed anti-inflammatory treatments to improve outcomes in patients with bronchiectasis not due to cystic fibrosis, within the EMBARC international clinical research network.

Acronym

AIR-NET

Study objectives

Primary objective:

To evaluate the effect of a range of interventions compared to usual care on the activity of NE in sputum

Secondary objectives:

1. To evaluate the effect of a range of interventions compared to usual care on the activity of NE in sputum
2. To evaluate the effect of a range of interventions compared to usual care on time to onset of first bronchiectasis exacerbation
3. To evaluate the effect of a range of interventions compared to usual care on quality of life
4. To evaluate the effect of a range of interventions compared to usual care on clinical benefits
5. To evaluate the effect of a range of interventions compared to usual care on walking distance
6. To evaluate the safety of a range of interventions compared with usual care
7. To evaluate the effect of a range of interventions on peripheral blood neutrophil function

Ethics approval required

Ethics approval required

Ethics approval(s)

approved 21/10/2024, London-Central Research Ethics Committee (3rd Floor 3 picadilly Place, London Road, Manchester, M1 3BN, United Kingdom; +44 (0)207 104 8061; londoncentral.rec@hra.nhs.uk), ref: 24/LO/0679

Study design

Interventional double blind randomized parallel group controlled trial

Primary study design

Interventional

Study type(s)

Efficacy, Safety

Health condition(s) or problem(s) studied

Bronchiectasis

Interventions

Participants will be randomised using the online Tayside Randomisation System (TRuST) to one of the trial arms:

Arm 1: Usual care

Arm 2: Disulfiram 200 mg, two tablets taken orally, once daily

Arm 3: Dipyridamole 200 mg, one capsule taken orally, twice daily

Arm 4: Doxycycline 100 mg, one capsule taken orally, once daily

Trial treatment will be for 28 days, with a 28 day follow up period.

Intervention Type

Drug

Phase

Phase II

Drug/device/biological/vaccine name(s)

Disulfiram, dipyridamole, doxycycline

Primary outcome(s)

Activity of sputum Neutrophil elastase. Day 0 and 28

Key secondary outcome(s)

Current secondary outcome measures as of 06/11/2024:

1. Activity of sputum Neutrophil elastase. Days 0, 7, 14 and 56
2. Time to first pulmonary exacerbation (EMBARC definition). Days 0 to 28
3. Quality of life-bronchiectasis (QOL-B) respiratory symptom scale, Bronchiectasis Impact Measure (BIM) questionnaire. Days 0, 7, 14, 28 and 56
4. Distance covered during a 6-minute walk. Days 0 to 28
5. Frequency of adverse events (AEs) and serious adverse events (SAEs). Day 0, 28
6. Phagocytosis of bacteria; Reactive oxygen species generation; Degranulation; Ex-vivo formation of neutrophil extracellular traps; Mass cytometry (endpoints may vary depending on the experimental arm). Days 0 to 56

Tertiary outcome measures:

1. Frequency of pulmonary exacerbations (EMBARC definition). Days 0 to 28; Days 0 to 56
2. Measure the concentration of MMPs and NETs in sputum, as well as other biomarkers e.g. proteomics, bacterial load, microbiome. Days 0, 7, 14, 28 and 56
3. Measure serum biomarkers of inflammation and redox status. Days 0, 7, 14, 28 and 56
4. Peripheral blood neutrophil proteomics performed on isolated cells. Days 0 and 28

Previous secondary outcome measures:

1. Activity of sputum Neutrophil elastase. Days 0, 7, 14 and 56
2. Time to first pulmonary exacerbation (EMBARC definition). Days 0 to 28
3. Quality of life-bronchiectasis (QOL-B) respiratory symptom scale, Bronchiectasis Impact Measure (BIM) questionnaire. Days 0, 7, 14, 28 and 56
4. Distance covered during a 6-minute walk. Days 0 to 28
5. Frequency of adverse events (AEs) and serious adverse events (SAEs). Day 0, 28
6. Phagocytosis of bacteria; Reactive oxygen species generation; Degranulation; Ex-vivo formation of neutrophil extracellular traps; Mass cytometry (endpoints may vary depending on the experimental arm). Days 0 to 56

Tertiary outcome measures:

1. Frequency of pulmonary exacerbations (EMBARC definition). Days 0 to 28; Days 0 to 56
2. Measure the concentration of MMPs and NETs in sputum, as well as other biomarkers e.g. proteomics, bacterial load, microbiome. Days 0, 7, 14, 28 and 56
3. Measure serum biomarkers of inflammation and redox status. Days 0, 7, 14, 28 and 56
4. Peripheral blood neutrophil proteomics performed on isolated cells. Days 0 and 28
5. Label-free liquid chromatography/mass spectrometry sputum identification and relative quantification of proteins. Day 0 and 28
6. Peripheral blood transcriptomics. Day 0 and 28
7. Nasal brushing transcriptomics. Day 0 and 28

Sub-study outcome measures:

1. Change in skin perfusion with iontophoresis of acetylcholine and sodium nitroprusside using laser Doppler perfusion imaging. Days 0, 28 and 56
2. Change in arterial stiffness index. Days 0, 28 and 56
3. Change in pulse wave velocity. Days 0, 28 and 56

Completion date

28/02/2027

Eligibility

Key inclusion criteria

1. ≥ 18 years
2. Able to provide informed consent.
3. Capable of complying with all trial procedures and of completing the trial, in the opinion of the investigator.
4. Bronchiectasis, confirmed by computed tomography (CT), showing bronchiectasis in 1 or more lobes.
5. Normally produces sputum daily.
6. Able to provide a sputum sample at the screening visit or between screening and

randomisation.

7. Active neutrophilic inflammation at screening/baseline indicated by a positive NEATstik (Neutrophil Elastase Airways Test) result.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Mixed

Lower age limit

18 years

Upper age limit

90 years

Sex

All

Total final enrolment

0

Key exclusion criteria

1. Enrolled previously in the trial 3 times.
2. Respiratory infection or bronchiectasis exacerbation 4 weeks prior to screening and/or between screening and randomisation
3. Antibiotic or corticosteroid 4 weeks prior to screening and/or between screening and randomisation
4. Active allergic bronchopulmonary aspergillosis (defined by International Society for Human and Animal Mycology criteria) on steroids and/or anti-fungals,
5. Nontuberculous mycobacterial infection on antibiotic therapy
6. Immunodeficiency on immunoglobulin replacement
7. A primary diagnosis of COPD or asthma (a secondary diagnosis of COPD or asthma is permitted)
8. Cystic fibrosis
9. Active malignancy except non-melanoma skin cancer
10. Currently taking brensocatib
11. Use of any investigational drugs within five times of the elimination half-life after the last dose or within 30 days, whichever is longer. Current enrolment in non-interventional, observational studies will be allowed
12. Currently pregnant or breast-feeding
13. Women of childbearing age and not practicing an acceptable method of birth control
14. Additional exclusion criteria for individual treatment arms are described in the protocol

Date of first enrolment

18/11/2024

Date of final enrolment

01/09/2026

Locations

Countries of recruitment

United Kingdom

England

Northern Ireland

Scotland

Study participating centre

NHS Tayside Ninewells Hospital

Clinical Research Centre

James Arnott Drive

Ninewells Hospital

Dundee

Scotland

DD1 9SY

Study participating centre

Royal Infirmary of Edinburgh at Little France

51 Little France Crescent

Old Dalkeith Road

Edinburgh

Lothian

Scotland

EH16 4SA

Study participating centre

Liverpool Heart & Chest Hospital

Broadgreen Hospital

Thomas Drive

Liverpool

England

L14 3PE

Study participating centre

Royal Hallamshire Hospital

Glossop Road

Sheffield

England
S10 2JF

Study participating centre

The Leeds Teaching Hospitals NHS Trust

Adult Cystic Fibrosis Unit j6, Level 6
Gledhow Wing
St James' University Hospital
Leeds
England
LS97TF

Study participating centre

Royal Devon University Healthcare NHS Foundation Trust

Royal Devon University NHS Ft
Barrack Road
Exeter
England
EX2 5DW

Study participating centre

Royal Papworth Hospital (papworth Everard)

Papworth Road
Cambridge Biomedical Campus
Cambridge
England
CB2 0AY

Study participating centre

Guy's and St Thomas' NHS Foundation Trust

Respiratory Clinic Research Facility
Level 4 1st Floor Fulham Wing
Royal Brompton Hospital
London
England
SW3 6HP

Study participating centre

Belfast Health and Social Care Trust

Northern Ireland Clinical Research Facility
U Floor - Belfast City Hospital

Lisburn Road
Belfast
England
BT9 7AB

Study participating centre
Oxford University Hospitals NHS Foundation Trust
Oxford Special Airways Clinic
John Radcliffe Hospital
Headley Way
Headington
Oxford
England
OX3 9DU

Study participating centre
University Hospital Southampton NHS Foundation Trust
NIHR Clinical Research Facility
Southampton University Hospital
Tremona Road
Southampton
England
SO16 6YD

Sponsor information

Organisation
University of Dundee

ROR
<https://ror.org/03h2bxq36>

Funder(s)

Funder type
Research organisation

Funder Name
LifeArc

Alternative Name(s)**Funding Body Type**

Private sector organisation

Funding Body Subtype

Other non-profit organizations

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from the Chief Investigator (Professor James Chalmers, j.chalmers@dundee.ac.uk). Approval of requests will be at the discretion of the Chief Investigator. Data will be available after the primary publication of the relevant experimental arm; there is no end date for requests. Data will be transferred using an appropriate secure method (e.g. encrypted email, file transfer system, or similar), and will be limited to only data required to perform the proposed analysis. Data will be anonymised or pseudonymised as appropriate. Patient consent for data sharing will be obtained at trial entry. The full details and terms of the transfer will be set out in a data sharing agreement.

IPD sharing plan summary

Available on request