

Can we reduce asthma attacks in children using exhaled nitric oxide measurements?

Submission date 10/04/2017	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 12/04/2017	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 28/05/2024	Condition category Respiratory	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Asthma is a long-term condition which affects the airways. When a person is suffering from asthma, the airways are extremely sensitive (hyperresponsive) to both natural chemicals the body produces and irritants outside the body, such as dust or pollen. Coming into contact with these substances can cause an asthma attack (also known as an exacerbation), which involves feelings of tightness in the chest as the airways become inflamed (swollen), causing coughing, wheezing, chest tightness and difficulty breathing. Every year in the UK, 150,000 children see their family doctor for an asthma exacerbation and 25,000 are hospitalised. One third of the £1 billion NHS budget for asthma is spent on provision for unscheduled care of which about one half is for childhood exacerbations. Exacerbations are relatively infrequent and short-lived but their importance to patients is emphasised in the Global Initiative for Asthma whose major goals include "to prevent asthma exacerbations". Everyone breathes out a gas called nitric oxide. Exhaled nitric oxide can be measured using a special breathing device. People with asthma breathe out more nitric oxide than people without asthma because nitric oxide is produced by the allergic cells which are present in the lungs of people with asthma. These allergic cells build up before an asthma attack. The aim of this study is to find out whether measuring fractional exhaled nitric oxide (FeNO) can guide asthma treatment and help prevent asthma attacks.

Who can participate?

Children aged between 6 and 16 who have been diagnosed with asthma and are currently being treated with inhaled steroids and who have had an asthma attack treated with steroid tablets.

What does the study involve?

Participants are randomly allocated to one of two groups. Those in the first group have their asthma treatment guided by symptoms and breath tests to measure FeNO. Those in the second group are treated according to the symptoms they are having alone. Participants in both groups are followed up after three, six, nine and twelve months in order to find out how many asthma exacerbations they have had and assessments of their quality of life. Participants are also offered additional allergy testing, which involves a blood test and a coughing up spit test. Additionally around 20 children are invited to take part in an interview at the end of the twelve month follow-up to explore attitudes to, and acceptability of the FeNO measurements. Five research nurses are also interviewed about the practicality of FeNO measurements.

What are the possible benefits and risks of participating?

All children will benefit from regular asthma assessments as part of the study. Asthma treatment is currently only guided by symptoms, and so children in the group where asthma treatment is guided by symptoms will continue to get current best asthma treatment. There is a risk that the optional allergy testing will cause itchiness on the participant's arm for a short time. The optional coughing up spit test may make the participant cough and possibly also wheeze, but this will only be conducted following a lung function check.

Where is the study run from?

25 hospitals around the UK

When is the study starting and how long is it expected to run for?

February 2017 to January 2021

Who is funding the study?

National Institute for Health Research (UK)

Who is the main contact?

1. Dr Stephen Turner (scientific)

s.w.turner@abdn.ac.uk

2. Mrs Jess Wood (public)

raaceno@abdn.ac.uk

Contact information

Type(s)

Scientific

Contact name

Dr Stephen Turner

ORCID ID

<https://orcid.org/0000-0001-8393-5060>

Contact details

Child Health

Royal Aberdeen Children's Hospital

Aberdeen

United Kingdom

AB25 2ZG

+44 1224 438470

s.w.turner@abdn.ac.uk

Type(s)

Public

Contact name

Mrs Jess Wood

Contact details

Centre for Healthcare Randomised Trials (CHaRT)
Health Services Research Unit
University of Aberdeen
Health Sciences Building
Foresterhill
Aberdeen
United Kingdom
AB51 2ZD
+44 1224 438179
raaceno@abdn.ac.uk

Additional identifiers

Protocol serial number

34390

Study information

Scientific Title

Reducing Asthma Attacks in Children using Exhaled Nitric Oxide as a biomarker to inform treatment strategy - a randomised trial (RAACENO)

Acronym

RAACENO

Study objectives

Current study hypothesis as of 08/02/2021:

Study aim:

The aim of the study is to compare treatment guided by exhaled nitric oxide and symptoms against treatment guided by symptoms alone (standard care), in children with asthma who are at risk of an asthma attack (ie have had an asthma attack in the previous 12 months), in terms of the presence of any asthma exacerbations over 12 months requiring prescription of OCS.

Hypothesis:

The proportion of children with ≥ 1 asthma exacerbation will be reduced when asthma treatment guided by FeNO plus symptoms is compared to treatment guided only by symptoms.

Previous study hypothesis:

Study aim:

The aim of the study is to compare treatment guided by exhaled nitric oxide and symptoms against treatment guided by symptoms alone (standard care), in children with asthma who are at risk of an asthma attack (ie have had an asthma attack in the previous 12 months), in terms of the presence of any asthma exacerbations over 12 months requiring prescription of OCS for 3-7 consecutive days.

Hypothesis:

The proportion of children with ≥ 1 asthma exacerbation will be reduced when asthma treatment guided by FeNO plus symptoms is compared to treatment guided only by symptoms.

Ethics approval required

Old ethics approval format

Ethics approval(s)

North of Scotland Research Ethics Committee 1, 17/10/2016, ref: 16/NS/0106

Study design

Randomized; Both; Design type: Screening, Diagnosis, Device, Active Monitoring, Qualitative

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Asthma

Interventions

Eligible and consenting participants will be randomised to one of the two groups (treatment decisions based on FeNO plus symptoms [interventional arm] or treatment decisions based on symptoms alone [standard care]) using a 24-hour telephone Interactive Voice Response randomisation application or via a web-based application, both hosted by CHaRT. Random allocation will use a minimisation algorithm (stratification by centre, age (<11 years, ≥ 11 years) sex and asthma severity as evidenced by BTS/SIGN treatment step (BTS step 2, BTS step 3, BTS step 4) including a random element (20%).

Experimental arm: Asthma treatment decisions will be guided by Fractional Exhaled Nitric Oxide (FeNO) and symptoms. The experimental intervention and subsequent adjustment of treatment steps are applied at recruitment, and at each of the follow-up visits (3, 6, 9 and 12 months).

FeNO guides treatment to either early escalation of anti-inflammatory medication (algorithm 1, elevated FeNO) or early intervention with bronchodilators (algorithm 2, FeNO not elevated).

Control arm: Asthma treatment decisions will be guided by symptoms alone. The control intervention and subsequent adjustment of treatment steps are applied at recruitment, and at each of the follow-up visits (3, 6, 9 and 12 months). Although FeNO will be measured in children in this arm at recruitment and at each of the follow-up visits, the results of the FeNO will not be used in treatment decisions for this arm of the trial. The FeNO will be used as outcome data only. For participants in this group, the FeNO result will not be available to the patient's GP or paediatrician (where appropriate) during the trial period.

In both arms, treatment decisions will be protocolised through the study website. Data required for step up/down decisions are entered into the study website which contains the algorithm for step up/down treatment decisions; this removes subjectivity from the treatment decisions. The trial algorithm applies to participants in both arms of the trial up to 12 month assessment.

Intervention Type

Other

Primary outcome(s)

Current primary outcome measure as of 08/02/2021:

Asthma exacerbation (attack) requiring prescription (and/or use of) one or more course of OCS in the 12 months after randomisation (yes/no). Outcomes will be assessed at the 3, 6, 9 and 12 months follow-up visits. In protocol version 3 (31 August 2017) we refined the definition of the primary outcome to reflect a change in treatment, where some exacerbations were being treated with a single dose of oral corticosteroid; and clarified that the primary source of information on exacerbations would be the child and parent or carer; with GP records used only where this information was missing.

Previous primary outcome measure:

Asthma exacerbation (attack) requiring prescription of OCS for 3-7 consecutive days determined from a patient held diary and verified by GP records where possible in the 12 months after randomisation (yes/no). Outcomes will be assessed at the 3, 6, 9 and 12 months follow-up visits.

Key secondary outcome(s))

1. Time to first exacerbation is determined from the child/parent or carer and from GP records in the 12 months after randomisation. Outcomes will be assessed at 3, 6, 9 and 12 months post randomisation
2. Number of exacerbations during follow up, based on prescribed oral corticosteroid, is assessed by responses to asthma attacks that have required oral steroid tablets on the follow-up CRF at 3,6,9, and 12 months follow-up determined from the child/parent or carer and from GP records in the 12 months after randomisation
3. Need for unscheduled healthcare assessment during follow up (yes/no) is measured by responses to health service use on the follow-up CRF at 3,6,9 and 12 months follow-up determined from the child/parent or carer and from GP records in the 12 months after randomisation
4. Number of unscheduled health assessments is measured by responses on the follow-up CRF to health service use at 3,6,9 and 12 months follow-up determined from the child/parent or carer and from GP records in the 12 months after randomisation
5. Asthma control during follow up (i.e. age-appropriate Asthma Control Test score) is assessed by responses to the Asthma Control Test (ACT)/Children's Asthma Control Test (CACT) - questionnaires completed by child/parent or carer at baseline, and at 3, 6, 9 and 12 month follow-up appointments
6. Asthma severity (%FEV1) is measured using spirometry at baseline and at 3, 6, 9 and 12 month follow-up appointments.
7. Fractional exhaled nitric oxide (FeNO) is measured by Niox Vero device at baseline, and 3, 6, 9 and 12 month follow-up appointments
8. Dose of ICS during the 12 months follow up (i.e. daily dose of budesonide equivalent averaged over 3 months) is assessed from current asthma treatment and from adherence to treatment (measured using adherence monitors); this data is collected at 3, 6, 9 and 12 months follow-up
9. Health status is assessed using the Paediatric Asthma Quality of Life Questionnaire (PAQLQ); a questionnaire completed by the child/parent or carer at baseline and 12 months
10. Exploration of experiences and ascertainment of acceptability of the intervention will be

collected through participant interviews following 12 month follow-up

11. Health economic evaluation is undertaken using data from GP records and participant reported data at baseline, 3, 6, 9 and 12 months follow-up

Completion date

31/01/2021

Eligibility

Key inclusion criteria

1. Asthma diagnosed or confirmed by consultant paediatrician (or Read code for asthma if recruited in primary care)
2. Aged 6 years or older and not yet reached the date of their 16th birthday
3. Currently prescribed inhaled corticosteroids (<1000mcg budesonide equivalent per day in those <12 years; <2000 mcg budesonide equivalent per day for older children)
4. At least one patient/parent reported asthma exacerbation treated with oral corticosteroids in the 12 months prior to recruitment

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

6 years

Upper age limit

15 years

Sex

All

Total final enrolment

515

Key exclusion criteria

1. Unable to provide FeNO measurement at baseline assessment (expected prevalence <5%)
2. Other chronic respiratory conditions which also have exacerbations
3. Current treatment with maintenance oral steroids

Date of first enrolment

01/06/2017

Date of final enrolment

01/08/2019

Locations

Countries of recruitment

United Kingdom

England

Scotland

Study participating centre

Royal Aberdeen Children's Hospital

Westburn Road

Aberdeen

United Kingdom

AB25 2ZG

Study participating centre

Birmingham Children's Hospital

Steelhouse Lane

Birmingham

United Kingdom

B4 6NH

Study participating centre

Bradford Teaching Hospitals NHS Foundation Trust

Children's Services Directorate

Duckworth Lane

Bradford

United Kingdom

BD9 6RJ

Study participating centre

Royal Alexandra Children's Hospital

Eastern Road

Brighton

United Kingdom

BN2 5BE

Study participating centre

Bristol Royal Hospital for Children

Paul O'Gorman Building

Upper Maudlin Street
Bristol
United Kingdom
BS2 8BJ

Study participating centre
Derbyshire Children's Hospital
Royal Derby Hospital
Uttoxeter Road
Derby
United Kingdom
DE22 3NE

Study participating centre
Royal Hospital for Sick Children
NHS Lothian
Royal Hospital for Sick Children
9 Sciennes Road
Edinburgh
United Kingdom
EH9 1LF

Study participating centre
Royal Hospital for Children
NHS Greater Glasgow & Clyde
1345 Govan Road
Glasgow
United Kingdom
G51 4TF

Study participating centre
Royal Brompton Hospital
Sydney Street
Chelsea
London
United Kingdom
SW3 6NP

Study participating centre
University Hospital Crosshouse
Kilmarnock Road

Crosshouse
Kilmarnock
United Kingdom
KA2 0BE

Study participating centre
Leicester Children's Hospital
Leicester Royal Infirmary
Infirmary Square
Leicester
United Kingdom
LE1 5WW

Study participating centre
Alder Hey Children's Hospital
E Prescott Road
Liverpool
United Kingdom
L14 5AB

Study participating centre
Manchester Royal Infirmary
Oxford Road
Manchester
United Kingdom
M13 9WL

Study participating centre
Great North Children's Hospital
Victoria Wing
Royal Victoria Infirmary
Newcastle upon Tyne
United Kingdom
NE1 4LP

Study participating centre
Nottingham Children's Hospital
Queen's Medical Centre
Derby Road

Nottingham
United Kingdom
NG7 2UH

Study participating centre
Southampton Children's Hospital
Southampton General Hospital
Tremona Road
Southampton
United Kingdom
SO16 6YD

Study participating centre
Royal Stoke Children's Service
Royal Stoke University Hospital
Newcastle Road
Stoke-on-Trent
United Kingdom
ST4 6PQ

Study participating centre
New Cross Hospital
Wolverhampton Road
Heath Town
Wolverhampton
United Kingdom
WV10 0QP

Study participating centre
Tayside Children's Hospital
NHS Tayside
Dundee
United Kingdom
DD1 9SY

Study participating centre
Barnsley Hospital
Gawber Road
Barnsley
United Kingdom
S75 2EP

Study participating centre
Sheffield Children's Hospital
Western Bank
Sheffield
United Kingdom
S10 2TH

Study participating centre
Royal Devon & Exeter Hospital
Barrack Road
Exeter
United Kingdom
EX2 5DW

Study participating centre
Derriford Hospital
Derriford Road
Plymouth
United Kingdom
PL6 8DH

Study participating centre
Walsall Manor Hospital
Moat Road
Walsall
United Kingdom
WS2 9PS

Study participating centre
Leeds General Infirmary
Great George Street
Leeds
United Kingdom
LS1 3EX

Study participating centre
King's College Hospital
Denmark Hill

London
United Kingdom
SE5 9RS

Study participating centre

Royal London Hospital

Barts Health NHS Trust
Whitechapel Road
Whitechapel
London
United Kingdom
E1 1FR

Study participating centre

Wythenshawe Hospital

Southmoor Road
Manchester
United Kingdom
M23 9LT

Study participating centre

St George Hospital

Blackshaw Road
London
United Kingdom
SW17 0QT

Study participating centre

Royal Shrewsbury Hospital

Mytton Oak Road
Shrewsbury
United Kingdom
SY3 8XQ

Study participating centre

Countess of Chester Hospital

Liverpool Road
Chester
United Kingdom
CH2 1UL

Study participating centre
Harrogate District Hospital
Lancaster Park Road
Harrogate
United Kingdom
HG2 7SX

Study participating centre
Russells Hall Hospital
Pensnett Road
Dudley
United Kingdom
DY1 2HQ

Study participating centre
Wishaw General Hospital
50 Netherton Street
Wishaw
United Kingdom
ML2 0DP

Study participating centre
Royal Oldham Hospital
Rochdale Road
Oldham
United Kingdom
OL1 2JH

Study participating centre
Royal Preston Hospital
Sharoe Green Lane
Preston
United Kingdom
PR2 9HT

Study participating centre
Ormskirk District General Hospital
Wigan Road
Ormskirk

United Kingdom
L39 2AZ

Study participating centre

Arrowe Park Hospital

Arrowe Park Road
Upton
Wirral
United Kingdom
CH49 5PE

Study participating centre

Hanscombe House Surgery

52A St Andrew Street
Hertford
United Kingdom
SG14 1JA

Study participating centre

Parsonage Surgery

Cavell Drive
Haymeads Lane
Bishops Stortford
United Kingdom
CM23 5JH

Study participating centre

Barrack Lane Medical Centre

1 Barrack Lane
Ipswich
United Kingdom
IP1 3NQ

Study participating centre

Roundwell Medical Centre

25-27 Dr Torrens Way
Costessey
United Kingdom
NR5 0GB

Study participating centre**Ixworth Surgery**

Peddars Close
Bury St Edmunds
United Kingdom
IP31 2HD

Study participating centre**Lakenham Surgery**

1 Ninham Street
Norwich
United Kingdom
NR1 3JJ

Study participating centre**Castle Partnership**

40 Fishergate
Norwich
United Kingdom
NR3 1SE

Sponsor information

Organisation

University of Aberdeen and NHS Grampian

ROR

<https://ror.org/00ma0mg56>

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research

Alternative Name(s)

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

Funding Body Type
Government organisation

Funding Body Subtype
National government

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 Dr Steve Turner (s.w.turner@abdn.ac.uk)

IPD sharing plan summary
Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	protocol	28/01/2022	02/02/2022	Yes	No
Results article		01/05/2022	07/06/2023	Yes	No
Protocol article		04/10/2019	07/10/2019	Yes	No
HRA research summary	Qualitative process evaluation		28/06/2023	No	No
Other publications		05/01/2023	07/06/2023	Yes	No
Other publications		27/05/2024	28/05/2024	Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes
Study website	Study website	11/11/2025	11/11/2025	No	Yes