The value of measuring airway inflammation in the management of asthma

Submission date Recruitment status Prospectively registered 23/01/2004 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 23/01/2004 Completed [X] Results Individual participant data **Last Edited** Condition category 08/11/2022 Respiratory

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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

RBF 00XX2

Study information

Scientific Title

The value of measuring airway inflammation in the management of asthma

Study objectives

Test the hypothesis that managing patients with asthma by effectively suppressing airway inflammation, using a standardised protocol, will improve the control of mild exacerbations, airway responsiveness and symptoms compared with standard clinical approach.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Asthma

Interventions

- 1. Treatment of airways inflammation
- 2. Standard care

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Improved asthma control; control of airway inflammation; change in spirometry over one-year, severe exacerbations; total use of inhaled corticosteroids in each group; proportion of patients screened with a diagnosis of asthma without objective evidence.

Secondary outcome measures

Not provided at time of registration

Overall study start date

19/06/2000

Completion date

18/05/2002

Eligibility

Key inclusion criteria

80 patients with asthma

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

80

Total final enrolment

74

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

19/06/2000

Date of final enrolment

18/05/2002

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Glenfield Hospital

Leicester United Kingdom LE3 9QP

Sponsor information

Organisation

NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details

The Department of Health Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.doh.gov.uk

Funder(s)

Funder type

Government

Funder Name

NHS Executive Trent (UK)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

Not provided at time of registration

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output typeDetailsDate createdDate addedPeer reviewed?Patient-facing?Results article30/11/2002YesNo