Does giving advice on avoiding allergies and triggers improve asthma control? A research trial

Submission date	Recruitment status No longer recruiting	Prospectively regist	
13/02/2008		[] Protocol	
Registration date	Overall study status Completed	[] Statistical analysis p	
26/02/2008		[X] Results	
Last Edited 21/06/2010	Condition category Respiratory	[] Individual participan	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s) Scientific

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers 04/016

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Study information

Scientific Title

Does structured advice on Allergy and Allergen Avoidance given by practice nurses improve control of asthma in primary care? A single blind randomised controlled trial

Acronym

AAA

Study objectives

The control of asthma can be improved by a structured allergy assessment followed by individualised avoidance advice, given by practice nurses in primary care.

Ethics approval required Old ethics approval format

Ethics approval(s) Ethics approval received from Bristol South NHS Ethics Committee on the 15th November 2004.

Study design Randomised controlled trial

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) Not specified

Study type(s) Quality of life

Participant information sheet

Health condition(s) or problem(s) studied Asthma

Interventions

Control patients:

Usual care asthma reviews (UC) consisted of assessment of symptoms, inhaler technique, and medication usage, and provision of self-management action plans.

Intervention patients:

Structured allergen and trigger avoidance advice reviews (AAA) comprised the elements of a usual review as above, supplemented by a structured asthma and allergy assessment consisting of:

- 1. Skin prick testing
- 2. Completion of the Structured Allergy Questionnaire and Asthma Trigger Inventory
- 3. Avoidance advice for identified triggers

Duration of these was about 30 - 45 minutes. There was no further contact with the nurse until follow up 4 months later.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Lung function/symptoms, assessed at 4 months.

Secondary outcome measures

Self efficacy, assessed at 4 months.

Overall study start date

01/01/2005

Completion date

01/01/2006

Eligibility

Key inclusion criteria

1. Diagnosis of asthma, confirmed by 20% peak expiratory flow [PEF] diurnal variation in medical records or 15% reversibility on spirometry (British Thoracic Society [BTS]/Scottish Intercollegiate Guidelines Network [SIGN] 2003)

2. Prescribed asthma medication within the past year

- 3. Aged between 16 to 55 years, either sex
- 4. Not had skin prick testing in the past 10 years

Participant type(s)

Patient

Age group

Adult

Sex Both

Target number of participants 200

Key exclusion criteria Unable to give informed consent. Date of first enrolment 01/01/2005

Date of final enrolment 01/01/2006

Locations

Countries of recruitment England

United Kingdom

Study participating centre Centre for Health Sciences London United Kingdom E1 4NS

Sponsor information

Organisation St George's Hospital Medical School (UK)

Sponsor details Cranmer Terrace London England United Kingdom SW17 0RE

Sponsor type Hospital/treatment centre

Website http://www.sgul.ac.uk/

ROR https://ror.org/040f08y74

Funder(s)

Funder type

Charity

Funder Name Asthma UK (UK) (ref: 04/016)

Alternative Name(s) Asthma UK, Asthma + Lung UK

Funding Body Type Private sector organisation

Funding Body Subtype Other non-profit organizations

Location United Kingdom

Results and Publications

Publication and dissemination plan Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<u>Results article</u>	results	01/01/2010		Yes	No