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Randomised controlled trial of Skin Prick Testing

Submission date 25/04/2006	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 06/06/2006	**Overall study status** Completed	
Last Edited 08/02/2012	**Condition category** Respiratory	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
06/039/SMI

Study information

Scientific Title

Acronym
SPT

Study objectives
Does allergy assessment and appropriate advice in general practice enhance the care of patients with asthma and rhinitis and is it cost effective compared with routine medical care?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Ethics approval received from the Multicentre Research Ethics Committee on the 26th May 2003 (ref: MREC/02/10/13).

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
GP practice

Study type(s)
Treatment

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

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

Interventions
Allergy intervention - structured allergy history and skin prick testing and appropriate advice on allergy avoidance versus routine medical care

Intervention Type
Other

Phase

Not Specified

Primary outcome measure
1. Symptom scores (validated scale for asthma and rhinoconjunctivitis)
2. Health-related quality of life (using child or adult specific instruments)

Secondary outcome measures
1. Patients' subjective assessment of symptomatic improvement
2. Change in patient management (introduction or withdrawal of allergen avoidance)

Overall study start date
01/05/2006

Completion date
01/05/2008

Eligibility

Key inclusion criteria
Patients with a working diagnosis of asthma and/or rhinoconjunctivitis

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
528 patients

Key exclusion criteria
1. Less than 5 years or greater than 50 years
2. Terminal illness
3. Confusional state
4. History of anaphylaxis
5. Tested in the preceding two years

Date of first enrolment
01/05/2006

Date of final enrolment
01/05/2008

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Brighton and Sussex Medical School
Brighton
United Kingdom
BN1 9PH

Sponsor information

Organisation
Brighton and Sussex University Hospitals NHS Trust (UK)

Sponsor details
Royal Sussex County Hospital
Eastern Road
Brighton
England
United Kingdom
BN2 5BE
+44 (0)1273 696955
scott.harfield@bsuh.nhs.uk

Sponsor type
Hospital/treatment centre

Website
http://www.bsuh.nhs.uk/

Funder(s)

Funder type
Industry

Funder Name
Brighton and Sussex Medical School (UK) - originally held by from Southampton University

Funder Name
Alk-Abello Ltd (UK) (ref: D105 R107)

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?
Results article	results	01/03/2009		Yes	No