

Randomised controlled trial of Skin Prick Testing

Submission date 25/04/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 06/06/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 08/02/2012	Condition category Respiratory	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof Helen Smith

Contact details
Brighton and Sussex Medical School
Mayfield House
University of Brighton
Falmer
Brighton
United Kingdom
BN1 9PH
+44 (0)1273 644563
h.e.smith@bsms.ac.uk

Additional identifiers

Protocol serial number
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

Study type(s)

Treatment

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(s)

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

Key secondary outcome(s))

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

Completion date

01/05/2008

Eligibility**Key inclusion criteria**

Patients with a working diagnosis of asthma and/or rhinoconjunctivitis

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

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

United Kingdom

England

Study participating centre

Brighton and Sussex Medical School

Brighton

United Kingdom

BN1 9PH

Sponsor information

Organisation

Brighton and Sussex University Hospitals NHS Trust (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

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
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes