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Sublingual immunotherapy for housedust mite

Submission date	Recruitment status	[ ] Prospectively registered
12/09/2003	Stopped	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
12/09/2003	Stopped	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
27/09/2011	Respiratory	[ ] Record updated in last year

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
N0256107987

Study information

Scientific Title

Study objectives
To see whether sublingual immunotherapy and probiotics given in combination will provide a safe effective means of modifying allergic disease expression and whether this will provide a safe and effective mode of treatment for common use.

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)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Respiratory: Housedust allergy

Interventions
Randomised controlled trial comparing sublingual immunotherapy and probiotics with standard treatment.

July 2008: the trial was stopped in 2003

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Service outcome development.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/09/2002

Completion date
30/09/2004

Reason abandoned (if study stopped)
Product to be used became unusable.

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
80 patients

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/09/2002

Date of final enrolment
30/09/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Department of Rhinology, RNTNE
London
United Kingdom
WC1X 8DA

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
The Royal Free Hampstead NHS Trust (UK)

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