

Sublingual immunotherapy for housedust mite

Submission date 12/09/2003	Recruitment status Stopped	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 27/09/2011	Condition category Respiratory	<input type="checkbox"/> Individual participant data <input type="checkbox"/> 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

Protocol serial number
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

Study type(s)

Not Specified

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(s)

Service outcome development.

Key secondary outcome(s)

Not provided at time of registration

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

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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**

United Kingdom

England

Study participating centre

Department of Rhinology, RNTNE

London

United Kingdom

WC1X 8DA

Sponsor information**Organisation**

Department of Health (UK)

Funder(s)**Funder type**

Government

Funder Name

The Royal Free Hampstead NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Not provided at time of registration