

Do self management plans for asthma based on symptoms confer advantage in reducing asthma morbidity in addition to regular attendance at a nurse run asthma clinic?

Submission date 23/01/2004	Recruitment status Stopped	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 23/01/2004	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 18/01/2010	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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

HUGHES1X

Study information

Scientific Title

Study objectives

Do self management plans for asthma based on symptoms confer advantage in reducing asthma morbidity in addition to regular attendance at a nurse run asthma clinic?

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)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Asthma

Interventions

1. Self management' group increasing inhaled preventer treatment in response to asthma symptoms
2. Control group - no self-management

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Between group comparison of asthma morbidity and medical involvement at end of study.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/1993

Completion date

31/12/1993

Reason abandoned (if study stopped)

Not specified

Eligibility

Key inclusion criteria

Patients attending asthma clinic

Participant type(s)

Patient

Age group

Not Specified

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not taking regular inhaled steroids

Date of first enrolment

01/01/1993

Date of final enrolment

31/12/1993

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

The Kirkbymoorside Surgery

York

United Kingdom

YO6 6AR

Sponsor information

Organisation

NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details

The Department of Health

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

+44 (0)20 7307 2622

dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

NHS Executive Northern and Yorkshire (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