

Peak flow in heart failure

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 05/12/2014	Condition category Circulatory System	<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
N0159093858

Study information

Scientific Title

Peak flow in heart failure

Study objectives

Does treatment with ipratropium bromide (Atrovent) improve peak flow and associated symptoms?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised double blind 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

Cardiovascular: Heart failure

Interventions

Ipratropium bromide vs standard practice

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Ipratropium bromide

Primary outcome measure

Improvement in peak flow rates and other symptoms.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/02/2001

Completion date

01/04/2004

Eligibility

Key inclusion criteria

Patients with heart failure

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/02/2001

Date of final enrolment

01/04/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

North Tees & Hartlepool NHS Trust

Stockton-on-Tees

United Kingdom

TS19 8PE

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**

Hospital/treatment centre

Funder Name

North Tees and Hartlepool 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