An evaluation of the cost effectiveness of single test screening spirometry in the early diagnosis and management of Chronic Obstructive Pulmonary Disease (COPD) in primary care

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year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0375126454

Study information

Scientific Title

Study objectives

Does knowledge of an abnormal spirometry result significantly improve smoke stop rates for smokers aged 35-75?

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: Chronic obstructive pulmonary disease (COPD)

Interventions

Subjects will be recruited opportunistically in primary care. On average each practice is expected to recruit between 80-90 individuals. Subjects who accept will then be randomised to receive or not receive spirometry at the start of the HTQ programme. Both groups will be assessed on generic quality of life (QoL) and anxiety. Practice nurses will be trained and administer all questionnaires and tests, for which they will receive a payment. At 12 months subjects in both groups will be reviewed and receive spirometry and repeat QoL and anxiety questionnaires.

Please note, this trial was stopped due to lack of funding.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/06/2002

Completion date

01/06/2004

Reason abandoned (if study stopped)

Lack of funding

Eligibility

Key inclusion criteria

All smokers with 20 pack years plus, aged between 35 and 75, presenting in primary care to the HTQ Programme will be eligible for entry into the trial. All subjects recruited to the HTQ programme who fulfill the eligibility criteria will be invited to take part in the study.

Participant type(s)

Patient

Age group

Adult

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

- 1. Eg breathlessness
- 2. Terminally ill
- 3. Subjects with prior knowledge of spirometry status

Date of first enrolment

01/06/2002

Date of final enrolment

01/06/2004

Locations

Countries of recruitment

England

TF3 1FB

United Kingdom

Study participating centre Stirchley Medical Practice Telford United Kingdom

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Government

Funder Name

Shrewsbury and Telford Research and Development Consortium (UK)

Results and Publications

Publication and dissemination planNot provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summaryNot provided at time of registration