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Using a screening questionnaire to reduce non-attendances at first appointments for smoking cessation advice clinics in general practice: a clustered randomised controlled trial

Submission date
28/09/2007

Recruitment status
No longer recruiting

Registration date
28/09/2007

Overall study status
Completed

Last Edited
17/10/2016

Condition category
Mental and Behavioural Disorders

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] Individual participant data
[ ] 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
N0026186755

Study information

Scientific Title
Using a screening questionnaire to reduce non-attendances at first appointments for smoking cessation advice clinics in general practice: a clustered randomised controlled trial

Study objectives
1. Can a short pre-booking questionnaire be used to reduce wastage due to non-attendance of first appointments for smoking cessation advice?
2. Does screening for readiness by using a questionnaire have an impact on successful quit rate and waiting list?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Clustered randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
GP practice

Study type(s)
Treatment

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Smoking cessation

Interventions
The study will be a controlled trial that will take place in general practices in Bristol. In order to avoid contamination, randomisation will take place at the practice level. The intervention practices will use a short questionnaire which has been designed to assess readiness to quit, as a screening tool, while the smoking cessation advice clinics in control practices will continue to take referrals according to usual practice and supply their audit data at the end of a 6 month period.

The trial will take place over a six-month period.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
The primary outcome measure will be the percentage of missed first appointments compared with corresponding pre-trial periods. The waiting list and the audited quit rates will also be examined. Analysis will also compare the questionnaire scores with a successful quit attempt. It is anticipated that this will be used to contribute to the validation of the questionnaire.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/10/2006

Completion date
30/09/2007

Eligibility

Key inclusion criteria
Patients who think they may like to make an appointment for the smoking advice sessions in 15 Bristol general practices.

Participant type(s)
Patient

Age group
Adult

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
01/10/2006

Date of final enrolment
30/09/2007

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Air Balloon Surgery
Bristol
United Kingdom
BS5 7PD

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2007 Update - Department of Health

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.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Air Balloon Surgery

Funder Name
Avon Primary Care Research Collaborative

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