

The implementation of a Dutch computer-generated tailored smoking cessation program AROM by pharmacists and general practitioners

Submission date
27/01/2006

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
27/01/2006

Overall study status
Completed

☐ Statistical analysis plan

☒ Results

Last Edited
04/11/2008

Condition category
Mental and Behavioural Disorders

☐ Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Study objectives

One tailored letter on smoking cessation will lead to more quit attempts at follow-up.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Received from the local medical ethics committee

Study design

Randomised, two-arm, controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Smoking cessation

Interventions

Adult smokers will receive either a tailored letter about smoking cessation (experimental group) or a thank you letter for participating (control group).

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Quitting activity between baseline and follow-up
2. Abstinence between baseline and follow-up

Secondary outcome measures

No secondary outcome measures

Overall study start date

01/01/1999

Completion date

31/12/2005

Eligibility

Key inclusion criteria

Having smoked in the last 7 days

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

1500

Key exclusion criteria

Younger than 17 years of age

Date of first enrolment

01/01/1999

Date of final enrolment

31/12/2005

Locations

Countries of recruitment

Netherlands

Study participating centre

Maastricht University

Maastricht

Netherlands

6200 MD

Sponsor information

Organisation

Sponsor not yet defined (The Netherlands)

Sponsor details

-
-
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Netherlands

Sponsor type

Not defined

Funder(s)

Funder type

Research organisation

Funder Name

The Netherlands Organisation for Health Research and Development (ZonMw) (The Netherlands)

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	01/02/2006		Yes	No