

Implementation of an evidence based smoking cessation strategy (SMOCC) for patients with COPD in primary care

Submission date 14/02/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 14/02/2006	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 25/10/2022	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

ClinicalTrials.gov (NCT)
NCT00294905

Protocol serial number
N/A

Study information

Scientific Title

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Study objectives

The large implementation of SMOCC will be more (cost-)effective than the usually applied basic dissemination strategies for guidelines.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval received from the local medical ethics committee

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Chronic Obstructive Pulmonary Disease (COPD), smoking.

Interventions

Large scale implementation of a combined strategy, aimed at the complete GP practice team (education by consultant at the practice, help with detecting smoking COPD patients, supplying materials for patient education, helpdesk/website, reminders by e-mail and phone) versus usual care.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Primary outcome measures will be biochemically validated smoking abstinence at 12 and 18 months.

Key secondary outcome(s)

Secondary outcome measures will be counseling contacts and counseling behaviour of professionals and cessation attempts of patients.

Completion date

01/11/2008

Reason abandoned (if study stopped)

Participant recruitment issue

Eligibility

Key inclusion criteria

1. COPD
2. Smoking
3. Age 40 or more

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

1. Under control of lung specialist
2. Not Dutch-speaking
3. Serious physical or psychiatric comorbidity
4. Age under 40

Date of first enrolment

01/01/2006

Date of final enrolment

01/11/2008

Locations

Countries of recruitment

Netherlands

Study participating centre

University Medical Centre St. Radboud
Nijmegen
Netherlands
6500 HB

Sponsor information

Organisation

University Medical Center St. Radboud, Centre for Quality of Care Research (WOK) (The Netherlands)

ROR

<https://ror.org/05wg1m734>

Funder(s)**Funder type**

Research organisation

Funder Name

Netherlands Organisation for Health Research and Development (ZonMw)

Alternative Name(s)

Netherlands Organisation for Health Research and Development

Funding Body Type

Private sector organisation

Funding Body Subtype

Other non-profit organizations

Location

Netherlands

Results and Publications**Individual participant data (IPD) sharing plan**

Not provided at time of registration

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		01/11/2005		Yes	No