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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	[_] Prospectively registered [_] Protocol
Registration date 14/02/2006	**Overall study status** Stopped	[_] Statistical analysis plan [X] Results
Last Edited 25/10/2022	**Condition category** Respiratory	[_] 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

ClinicalTrials.gov (NCT)
NCT00294905

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

Primary study design
Interventional

Study design
Randomised controlled trial

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