

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

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

### Contact name

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## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

NCT00294905

## Secondary identifying numbers

N/A

# Study information

## Scientific Title

-

## 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

## 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

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 measure**

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

**Secondary outcome measures**

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

**Overall study start date**

01/01/2006

**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

**Age group**

Not Specified

**Sex**

Not Specified

**Target number of participants**

2700 (open to recruitment)

**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)

**Sponsor details**

117 KWAZO

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**Sponsor type**

Not defined

**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

## Publication and dissemination plan

Not provided at time of registration

## Intention to publish date

## 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?
<a href="#">Results article</a>		01/11/2005		Yes	No