Implementation of a smoking cessation protocol in dental practices

Submission date	Recruitment status	Prospectively registered
04/04/2006	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
04/04/2006	Completed	Results
Last Edited	Condition category	Individual participant data
18/08/2009	Mental and Behavioural Disorders	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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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

NTR614

Study information

Scientific Title

Acronym

TAPS

Study objectives

The intervention increases advice given by dental professionals, the number of patients having quit smoking and the number of quit attempts.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Tobacco addiction

Interventions

Provision of a 4 hour central course for whole practice teams, patient-mediated monitoring and feedback of the provision of cessation advice, patient education materials and a task organisation protocol for the various members of the dental team.

The dental teams in the control group deliver usual care.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Patient-reported number of smoking cessation advice given by the dental team, patients' smoking status and quit attempts.

Secondary outcome measures

Professionals' attitudes towards smoking and smoking cessation support, their knowledge of health consequences of smoking tobacco (especially dental health), patients' perspectives of smoking cessation.

Support by dentists will be measured as well as self-efficacy to provide smoking cessation support.

Overall study start date

01/01/2006

Completion date

01/11/2008

Eligibility

Key inclusion criteria

Smoking patients visiting dental practices.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

1300

Key exclusion criteria

Patients under the age of 18.

Date of first enrolment

01/01/2006

Date of final enrolment

01/11/2008

Locations

Countries of recruitment

Netherlands

Study participating centre University Medical Center St Radboud Nijmegen Netherlands 6500 HB

Sponsor information

Organisation

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

Sponsor details

117 KWAZO P.O. Box 9101 Nijmegen Netherlands 6500 HB

Sponsor type

Not defined

ROR

https://ror.org/05wg1m734

Funder(s)

Funder type

Industry

Funder Name

Pfizer (Netherlands)

Alternative Name(s)

Pfizer Inc., Pfizer Consumer Healthcare, Davis, Charles Pfizer & Company, Warner-Lambert, King Pharmaceuticals, Wyeth Pharmaceuticals, Seagen

Funding Body Type

Government organisation

Funding Body Subtype

For-profit companies (industry)

Location

Results and Publications

Publication and dissemination planNot provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summaryNot provided at time of registration