

Implementation of a smoking cessation protocol in dental practices

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

Protocol serial number
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

Study type(s)

Treatment

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(s)

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

Key secondary outcome(s))

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.

Completion date

01/11/2008

Eligibility

Key inclusion criteria

Smoking patients visiting dental practices.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

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)

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, Pfizer Inc

Funding Body Type

Government organisation

Funding Body Subtype

For-profit companies (industry)

Location

United States of America

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary