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

Gepubliceerd: 18-01-2006 Laatst bijgewerkt: 18-08-2022

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

Ethische beoordeling	Positief advies
Status	Werving gestart
Type aandoening	-
Onderzoekstype	Interventie onderzoek

Samenvatting

ID

NL-OMON28475

Bron
Nationaal Trial Register

Verkorte titel
N/A

Aandoening

COPD, smoking

Ondersteuning

Primaire sponsor: ZONMW

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

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

Toelichting onderzoek

Achtergrond van het onderzoek

COPD is an increasing cause of death and morbidity and smoking is its primary cause. Professional smoking cessation treatment is very cost-effective and therefore recommended by national guidelines. A controlled study demonstrated that a smoking cessation protocol in routine primary care, specifically targeted at patients with COPD (SMOCC), doubled the quit rates. The protocol was tested under optimal trial conditions, but it is unclear if a large-scale implementation strategy is (cost-)effective. Therefore the present study investigates an large scale implementation strategy in a 2-armed community intervention trial. The research question is how (cost-)effective this implementation strategy is compared to usual implementation procedures.

Doel van het onderzoek

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

Onderzoeksproduct en/of interventie

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.

Contactpersonen

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

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

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

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

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Blindering:	Open / niet geblindeerd
Controle:	N.v.t. / onbekend

Deelname

Nederland

Status:	Werving gestart
(Verwachte) startdatum:	01-01-2006
Aantal proefpersonen:	2700

Type:	Verwachte startdatum

Ethische beoordeling

Positief advies

Datum:	18-01-2006
Soort:	Eerste indiening

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

Geen registraties gevonden.

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register	ID
NTR-new	NL524
NTR-old	NTR568
Ander register	: N/A
ISRCTN	ISRCTN52757029

Resultaten

Samenvatting resultaten
N/A