Patients’ preferences for esophageal cancer screening strategies

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patient preferences

Ethische beoordeling	Positief advies
Status	Werving gestopt
Type aandoening	-
Onderzoekstype	Observationeel onderzoek, zonder invasieve metingen

Samenvatting

ID

NL-OMON25816

Bron
Nationaal Trial Register

Aandoening

esophageal cancer
Barrett's esophagus

Ondersteuning

Primaire sponsor: Radboud university medical center
Overige ondersteuning: fund = initiator = sponsor

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

The main endpoint of this study is the relative importance of the selected attributes in the CBC analysis, expressed in the utility of each individual attribute.
Utilities indicate how much the difference between the levels of one attribute affects the decision of a subject to take part in a screening program.

Toelichting onderzoek

Achtergrond van het onderzoek

To explore preferences for EAC screening tests attributes and screening test, a survey with a choice-based conjoint analysis will be conducted. Subjects will be asked on their preferences for aspects of EAC screening methods. Each individual will answer 12 discrete choice questions presenting two hypothetical tests comprised of 5 attributes.

Doel van het onderzoek

patient preferences

Onderzoeksopzet

time frame: 8 months

Onderzoeksproduct en/of interventie

Choice-based conjoint analysis (CBC)

CBC analyses involve surveys in which respondents are asked to choose between hypothetical alternatives defined by a set of differing attributes. The method is based on the idea that goods and (health care) services can be described by their characteristics, also called attributes, and each attribute is assigned a range of predefined dimensions called attribute-levels. The levels of attributes will be varied systematically in a series of questions and respondents will choose the option that they prefer for each question.

Ultimately, CBC analysis can determine which attributes are driving patients preferences, the trade-offs people make between attributes and how changes in attributes can lead to changes in preferences and screening uptake.

Contactpersonen

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

In order to be eligible to participate in this study, a subject must meet all of the following criteria:

- The subjects is aged 50-74 years old.

- The subject is randomly selected from the municipal database of Nijmegen

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

- The subject is illiterate.

- The subject has been previous diagnosed with Barrett’s esophagus or esophageal cancer.

- The subjects has a history of esophageal or gastric resection or surgery which has changed the esophageal anatomy.

- Patients with prior history of ablation (photodynamic therapy, radiofrequency ablation, cryotherapy, argon plasma coagulation) or endoscopic mucosal resection.

- The subject is unable to provide informed consent.

Onderzoeksopzet

Opzet

Type:	Observationeel onderzoek, zonder invasieve metingen
Onderzoeksmodel:	Anders
Toewijzing:	N.v.t. / één studie arm

Controle: N.v.t. / onbekend

Deelname

Nederland

Status:	Werving gestopt
(Verwachte) startdatum:	01-04-2018
Aantal proefpersonen:	400
Type:	Werkelijke startdatum

Voornemen beschikbaar stellen Individuele Patiënten Data (IPD)

Wordt de data na het onderzoek gedeeld: Nog niet bepaald

Ethische beoordeling

Positief advies

Datum:	26-02-2018
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	NL6869
NTR-old	NTR7047
Ander register	2018-4079 : 2018-4079

Resultaten