

Kwaliteit van communicatie in de spreekkamer.

Gepubliceerd: 05-11-2014 Laatste bijgewerkt: 18-08-2022

With a multi-faceted strategy we are able to implement shared decision making so that it meets the needs and demands of both professionals and patients, without interfering in daily practice.

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

Samenvatting

ID

NL-OMON23927

Bron

NTR

Aandoening

- breast cancer
- shared decision making
- quality improvement
- patient preferences
- patient participation

Ondersteuning

Primaire sponsor: Universiteit Maastricht

Overige ondersteuning: Pink Ribbon

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

- Knowledge on treatment

- Perceived shared decision making

- Decisional conflict

Toelichting onderzoek

Achtergrond van het onderzoek

Background and aim

The majority of patients diagnosed with early stage breast cancer are eligible to decide between having a mastectomy or lumpectomy with radiation (breast conserving therapy). The long term survival for mastectomy versus lumpectomy and radiotherapy is comparable, therefore patients' informed preferences are important in the decision making. Although most clinicians believe they involve patients in the decision making process, the information that women with breast cancer receive regarding the surgical options in breast cancer is often colored. Shared decision making can help patients to clarify their preferences. Patient decision aids support shared decision making.

The aim of this study is to pilot-test and optimize strategies for implementation of shared decision making on the treatment of early-stage breast cancer in the true clinical setting.

Methods/Design

This study concerns a pre-post pilot implementation study, carried out from October 2014 to June 2015. The intervention consists of implementing shared decision making (SDM) using a patient decision aid. The intervention will be evaluated using qualitative and quantitative measures, acquired prior to, during and after the implementation of SDM. Outcome measures are knowledge on treatment, perceived shared decision making and decisional conflict. We also will also conduct face-to-face interviews with a purposeful sample of these patients and their caregivers to assess their experiences on implementation of shared decision making and the patient decision aid.

Discussion

The outcomes and findings of this study will be used as a basis to finalise a multi-faceted implementation strategy with the intention to test the implementation of shared decision making and a patient decision aid on cost-effectiveness in a multicentre cluster RCT.

Country of recruitment: The Netherlands.

Doel van het onderzoek

With a multi-faceted strategy we are able to implement shared decision making so that it meets the needs and demands of both professionals and patients, without interfering in daily practice.

Onderzoeksopzet

- Knowledge on treatment and perceived shared decision making are measured immediately after intervention.
- Decisional conflict is measured two months after treatment.
- Patients and caregivers experiences are measured within three months after the implementation.

Onderzoeksproduct en/of interventie

Implementing shared decision making (SDM) using a patient decision aid.

Contactpersonen

Publiek

Postbus 616
W. Savelberg
Maastricht 6200 MD
The Netherlands
043-3882336

Wetenschappelijk

Postbus 616
W. Savelberg
Maastricht 6200 MD
The Netherlands
043-3882336

Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

Patients diagnosed with stage I or II breast cancer, provided the two treatment options, mastectomy or breast conserving surgery with radiotherapy, are applicable. Eligible patient should speak and understand the Dutch language.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

-Men diagnosed with breast cancer

-Women not eligible to decide between having a mastectomy or lumpectomy with radiation.

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Anders
Toewijzing:	N.v.t. / één studie arm
Blinding:	Open / niet geblindeerd
Controle:	Geneesmiddel

Deelname

Nederland	
Status:	Werving gestart
(Verwachte) startdatum:	01-10-2014
Aantal proefpersonen:	40
Type:	Verwachte startdatum

Ethische beoordeling

Positief advies	
Datum:	05-11-2014

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	NL4751
NTR-old	NTR4879
Ander register	Pink Ribbon : 2012.PS23

Resultaten