Enhancing patient communication in oncology consultations.

Gepubliceerd: 14-12-2012 Laatst bijgewerkt: 18-08-2022

N/A

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

Samenvatting

ID

NL-OMON23683

Bron
Nationaal Trial Register

Verkorte titel
PatientTIME

Aandoening

malignant lymphoma, patient-provider communication, e-health, web-based tailored information, participatory research,

Ondersteuning

Primaire sponsor: Nivel, UMC St Radboud
Overige ondersteuning: Dutch Cancer Society (The Netherlands)

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

Perceived and experienced efficacy for communication (QT0, QT3).

Toelichting onderzoek

Achtergrond van het onderzoek

OBJECTIVE:

The aim of the study is to test the effects of an online communication training to enhance cancer patients' utterance of questions, cues and concerns during oncology (follow-up) visits and their evaluation of the communication. The secondary aim of the study is to examine the feasibility of granting patients more control in the execution of a research project by asking them to collect audio recordings of their own follow-up visits.

STUDY DESIGN:

Using a randomised controlled trial, this study examines the effectiveness of an online communication training for patients diagnosed with malignant lymphoma. Patients will be randomly assigned to receive the online communication training or to receive access at the end of the study. Both intervention and control group patients will be asked to fill in questionnaires. Part of the participants will be provided with audio equipment to audiotape their consultations.

Doel van het onderzoek

N/A

Onderzoeksopzet

Patients will be asked to complete an online questionnaire at inclusion (QTi), before (QT0) and after (QT1) their first consultation during the year of data collection (with a maximum of 3 consultations). Subsequently, they will be asked to complete an online questionnaire after each following consultation (QT2) and three months after their last measured consultation (QT3).

Onderzoeksproduct en/of interventie

The intervention is an online communication training for patients, focused on communicating with their health care provider. The goal of the training is the support patients to prepare their consultations. The training is computer-tailored to patients’ efficacy for communication with health care providers, to their communication goals, to whether they attend their consultations alone or with a companion and to their health status. The training includes information, advice and video fragments of simulated consultations to model adequate communication behaviour. Additionally, the intervention includes a QPS (question prompt sheet). Patients can print this sheet to take it to the consultation. The intervention group

receives access to the online communication training one week before every consultation during the year of measurement. In this week the participants will be asked to visit the training before they enter the consultation. The participants can access the training from their own computer, at the time they prefer and as many times as they want during that week.

The control group will receive access to the program after the data-collection period has finished.

Contactpersonen

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

1. Diagnosed with malignant lymphoma;

2. Age 18 or older;

3. Having internet access at home;

4. At least one (follow up) consultation per year.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

1. Age under 18;

2. Not having internet access at home;

3. Less than one (follow up) consultation per year.

Onderzoeksopzet

Opzet

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

Deelname

Nederland

Status:	Werving gestart
(Verwachte) startdatum:	01-03-2013
Aantal proefpersonen:	100
Type:	Verwachte startdatum

Voornemen beschikbaar stellen Individuele Patiënten Data (IPD)

Wordt de data na het onderzoek gedeeld: Nog niet bepaald

Ethische beoordeling

Positief advies

Datum:	14-12-2012

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	NL3613
NTR-old	NTR3779
Ander register	KWF / METC St Radboud : NIVEL 2010-4747 / 38333.091.11
ISRCTN	ISRCTN wordt niet meer aangevraagd.

Resultaten

Samenvatting resultaten
N/A