

Informing Patients about Symptom Distress

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1a. Providing information about expected symptom distress due to chemotherapy in a structured manner improves patients' information recall, satisfaction, and trust in the physician. 1b. Communicating information about expected symptom...

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

Samenvatting

ID

NL-OMON22417

Bron

NTR

Verkorte titel

INSTRUCT

Aandoening

Patient-caregiver communication, information structuring, tailoring, empathy, hematology, malignant lymphoma, experimental video-vignettes

Ondersteuning

Primaire sponsor: Department of Medical Psychology
Academic Medical Center | University of Amsterdam

Overige ondersteuning: Dutch Cancer Society

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

Primary outcome of the study is participants' recall (active memory and recognition) of medical information.

Toelichting onderzoek

Achtergrond van het onderzoek

Treatment-related symptoms cause patients suffering from cancer significant distress. This impairs patients' everyday functioning, sometimes even leading them to discontinue their treatment. Providing patients with information that is congruent with their needs can help them to prepare for treatment, increase their adherence, reduce their distress, and ultimately promote their recovery. Adequate information provision thus forms a crucial component of optimal cancer care. Yet, while it has been consistently shown that patients indeed value information about symptoms highly, their information needs are often not met. Hence, improving oncologists' information-giving skills is imperative. While much has been written about the importance of information provision to ensure better health outcomes, the exact pathways underlying this process remain largely unclear. To advance oncologists' information-giving skills, it must thus first be determined what optimal provision of information in fact entails.

The aim of the overall project, therefore, is twofold. First, through a series of experimental studies, the essential components of oncologists' – and, more specifically, hematologists' – adequate information provision are explored. Doing so, particular attention is paid to the ways in which doctors provide structure and tailor information to the patient's needs. Moreover, the role of doctors' display of empathy is investigated. The effects on patients' recall of information, information satisfaction, and perceived distress are of prime interest. Second, on the basis of the experimental findings, this project ultimately seeks to practically contribute to the improvement of oncologists' information-giving competence by designing and evaluating different prototypes of innovative training modules (e.g., mobile apps, e-learning programs, and serious games). This is done in close collaboration with a panel of hematologists, to ensure their learning needs are met. Led by a team of psychologists, health communication researchers, education specialists, and doctors, this project thereby combines insights from medical practice with a solid, theory-driven approach to ultimately contribute to the reduction of patients' treatment-related symptom distress.

Doel van het onderzoek

1a. Providing information about expected symptom distress due to chemotherapy in a structured manner improves patients' information recall, satisfaction, and trust in the physician.

1b. Communicating information about expected symptom distress due to chemotherapy in an emphatic manner improves patients' information recall, satisfaction, and trust in the physician.

1c. The hypothesized effect in 1a is strengthened when the physician communicates information about expected symptom distress due to chemotherapy in an emphatic manner (cf. 1b).

2a. Tailoring information about expected symptom distress due to chemotherapy to the patient's preferences improves patients' information recall, satisfaction, and trust in the physician.

2b. The hypothesized effect in 2a is strengthened when the physician communicates information about expected symptom distress due to chemotherapy in an emphatic manner (cf. 1b).

Onderzoeksopzet

Participants will be asked to complete a short series of questions before the intervention (Q0) as well as after having viewed the video (Q1). Q0, video viewing, and Q1 have to be completed in one sitting.

Onderzoeksproduct en/of interventie

This experimental study consists of twelve experimental conditions, including two main streams: 4 conditions pertaining to H1 (information structuring) and 8 conditions belonging to H2 (information tailoring). Each participant is randomly assigned to one of the twelve conditions. The intervention consists of a role-played video-vignette of a consultation between a hematologist and a patient. Upon having seen the video, participants are asked to complete a series of questions concerning their perceptions, knowledge, and thoughts about the video.

In the first experimental stream, participants receive a video in which information structuring is varied (high/low) and crossed with physician empathy (high/low) in a 2x2 factorial design.

In the second experimental stream patient information preferences (high/low) are combined with information tailoring (preferences met/unmet) and subsequently crossed with physician empathy in a 2x2x2 design.

Contactpersonen

Publiek

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Wetenschappelijk

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

1. Age 18 or older;
2. Dutch speaking;
3. Having a stable internet connection

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

1. Age under 18
2. Non-Dutch speaking

- 3. Internet connection problems
- 4. Reading/comprehension issues

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Toewijzing:	Gerandomiseerd
Blinding:	Enkelblind
Controle:	Geneesmiddel

Deelname

Nederland	
Status:	Werving gestart
(Verwachte) startdatum:	17-10-2016
Aantal proefpersonen:	420
Type:	Verwachte startdatum

Ethische beoordeling

Positief advies	
Datum:	03-11-2016
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	NL6022
NTR-old	NTR6153
Ander register	KWF AMCUVA 2014-6777 : AMR CJ451014

Resultaten

Samenvatting resultaten

N/A