patient participation in 10 projects in palliative care

Gepubliceerd: 08-06-2016 Laatst bijgewerkt: 18-08-2022

The main objective of this study is to study the impact of patient participation from the perspective of patient(representatives) and researchers in 10 projects in palliative care. And to sustainably strengthen patient participation in the context...

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

Samenvatting

ID

NL-OMON22964

Bron
NTR

Aandoening

palliative care, terminal care

Ondersteuning

Primaire sponsor: Zorgbelang Limburg, Hogeschool Zuyd, Huis voor de Zorg
Overige ondersteuning: ZON-MW, The Netherlands Organization for Health Research and Development

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

The impact of patient participation from the perspective of patient(representatives) and researchers. Additionally, the underlying processes, how the impact was created, will be studies by using the context-mechanism-outcome classification.

Toelichting onderzoek

Achtergrond van het onderzoek

Participation of patients in palliative care is a relatively new development. Professionals often want to do more in patient participation. However, they lack the knowledge and experience to put the mechanisms of participation into meaningful practice. The main objective of this study is to sustainably strengthen patient participation in the context of palliative care in education, research and care practice projects.

This study consists of participatory action research with multiple case studies. The study will be carried out within a national research programme on palliative care. Ten out of 18 projects will be involved covering a range of different palliative care contexts such as education, research and practice.

The participants are patient (representatives) and senior and junior researchers.

The first part, implementation of patient participation, consists of several implementation activities: a)training and support for patient(representatives); b) training and coaching for research; c) working with the participation matric; d) setting up a participation community of practice among all 10 participating projects; e) development-oriented evaluation and creating sustainable conditions.

The second part focus on researching the impact of patient participation from the perspective of patient(representatives) and researchers. Additionally, the underlying processes, how the impact was created, will be studies by using the context-mechanism-outcome classification.

During the implementation data is collected through field notes, observations, informal conversations and video recordings. In the evaluation of the impact the data will be collected by in-depth interviews and focus group discussions. Data will be analyzed using content analysis.

Doel van het onderzoek

The main objective of this study is to study the impact of patient participation from the perspective of patient(representatives) and researchers in 10 projects in palliative care. And to sustainably strengthen patient participation in the context of palliative care in education, research and care practice projects.

Onderzoeksopzet

During the implementation data is collected through field notes, observations, informal conversations and video recordings. In the evaluation of the impact the data will be collected by in-depth interviews and focus group discussions.

Onderzoeksproduct en/of interventie

the implementation of patient participation, consists of several implementation activities: a)training and support for patient(representatives); b) training and coaching for research; c) working with the participation matric; d) setting up a participation community of practice among all 10 participating projects; e) development-oriented evaluation and creating sustainable conditions.

Contactpersonen

Publiek

Mercator 1
E. Stoffers
Sittard 6135 KW
The Netherlands
+31 (0)6 40160551

Wetenschappelijk

Mercator 1
E. Stoffers
Sittard 6135 KW
The Netherlands
+31 (0)6 40160551

Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

The participants are patient (representatives) and senior and junior researchers.

Belangrijkste redenen om niet deel te kunnen nemen

(Exclusiecriteria)

N/A

Onderzoeksopzet

Opzet

Onderzoeksmodel: Anders
Controle: N.v.t. / onbekend

Deelname

Nederland

Status:	Werving gestart
(Verwachte) startdatum:	15-06-2016
Aantal proefpersonen:	30
Type:	Verwachte startdatum

Ethische beoordeling

Positief advies

Datum:	08-06-2016
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	NL5474
NTR-old	NTR5891
Ander register	METC Zuyderland-Zuyd : 16-N-108

Resultaten

Samenvatting resultaten
N/A