Substitution of usual perioperative care by e-health & ICT in major abdominal surgery

Gepubliceerd: 04-02-2016 Laatst bijgewerkt: 18-08-2022

Considerable health efficiency gains can be achieved by the substitution and optimization of usual perioperative care by means of e-health and ICT.

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

Samenvatting

ID

NL-OMON24980

Bron
NTR

Verkorte titel
Ik Herstel 3.0 (second phase)

Aandoening

peri-operative care (perioperatieve zorg), eHealth, colonresection (colonresectie), hysterectomy (hysterectomie)

Ondersteuning

Primaire sponsor: VU University Medical Center,
EMGO+ Institute
Overige ondersteuning: ZonMw, The Netherlands Organization for Health Research and Development

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

- Return-to-normal activities (RNA)

Toelichting onderzoek

Achtergrond van het onderzoek

In the last decade the number of surgeries increased with 30% in the Netherlands. The increase of surgeries leads to rising hospital care costs. To reduce costs, in-hospital perioperative care is increasingly reduced due to one day hospitalisations and transferred to primary care. Guidance & monitoring on recovery and resumption of (work)activities are mostly not provided in secondary and primary care. Studies showed that due to the poor quality of usual perioperative care, return-to-normal-activities/work after surgery is hampered, leading to high productivity loss costs. We hypothesize that considerable health efficiency gains can be achieved by the substitution and optimization of usual perioperative care by means of e-health and ICT. In this trial we will study the (cost)effectiveness of a transmural, perioperative care program for patients undergoing abdominal surgery.

Doel van het onderzoek

Considerable health efficiency gains can be achieved by the substitution and optimization of usual perioperative care by means of e-health and ICT.

Onderzoeksopzet

1. Baseline;
2. 2 weeks;

3. 4 weeks

4. 6 weeks;

5. 3 months;

6. 6 months;

7. 9. months

8. 12 months

Onderzoeksproduct en/of interventie

Multidisciplinary perioperative care program including an interactive webportal. It aims to improve recovery and reduce costs by:

-SELFMANAGEMENT & EMPOWERMENT of patients during the perioperative period by supporting them with personalized pre- and postoperative recommendations to return to normal (work) activities. These recommendations are tailor made:
they are based on patient's own input of normal preoperative activities and the surgical technique applied (using algoritms).

-MONITORING OF POSTOPERATIVE CARE: With the webportal the patient as well as all involved physicians can monitor patient's recovery (bench mark information) and thus identify recovery problems.

-E-CONSULTATION is offered to patients to ask questions in case of recovery problems or to substitute standard postoperative consultation in outpatient clinics

Contactpersonen

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Wetenschappelijk

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

- Colonresection (left hemicolectomy, right hemicolectomy, extended version, transversumresection, sigmoidresection, segmentectomy, hartmann procedure)

- Hysterectomy (total laparoscopic hysterectomy or abdominal uterus extirpation)

- 18 - 75 years

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

In general:

- Concomitant health problems affecting daily activities

- Combination of surgery with other surgical procedures

- Severe comorbidity which might complicate the postoperative course

- Patient who are unable to understand the information belonging the research

- Insufficient understanding or ability to fill in (Dutch) questionnaires

Colonresection group:

- Surgery without a curative intention

- Neoadjuvant treatment

- Colectomy because of crohn’s disease or ulcerative colitis

Hysterectomy group:

- Deep infiltrating endometriosis

- Malignancies

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Toewijzing:	Gerandomiseerd
Blindering:	Open / niet geblindeerd
Controle:	Geneesmiddel

Deelname

Nederland

Status:	Werving gestopt
(Verwachte) startdatum:	01-02-2016
Aantal proefpersonen:	0
Type:	Werkelijke startdatum

Ethische beoordeling

Positief advies

Datum:	04-02-2016
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	NL5565
NTR-old	NTR5686
Ander register	ZonMW : 837002409

Resultaten

Samenvatting resultaten
N/A