

Development of a home-based exercise program delivered by means of a tablet-PC for patients after a Total Hip operation.

No registrations found.

Ethical review	Positive opinion
Status	Other
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON24746

Source

NTR

Health condition

Total Hip Arthroplasty, rehabilitation, home based exercise program

Sponsors and support

Primary sponsor: UMC Groningen

Source(s) of monetary or material Support: AnnaFonds & SPRINT

Intervention

Outcome measures

Primary outcome

The primary outcome measure is feasibility determined by an evaluation questionnaire and adherence to the program

Secondary outcome

The secondary outcome measure is clinical effectiveness measured by the HOOS, SF-36, EQ5D, Timed Up & Go Test and the Five Times Sit-to Stand Test.

Study description

Background summary

Background

Due to the projected growth of the older population and obesity epidemic, the number of Total Hip Arthroplasties (THA) is expected to increase dramatically in the coming decades. In the Netherlands an increasing number of THA patients are undergoing fast track surgery, after which people leave the hospital within a few days. This quick transfer leads to suboptimal rehabilitation. In addition postoperative physiotherapy is not regularly covered by the health insurance. To avoid suboptimal rehabilitation a home based telemonitored rehabilitation program is recommended.

Objectives

The first aim is to determine the feasibility of a home based rehabilitation program delivered by means of a tablet-pc for patients after THA. Second aim is to get a first impression of the clinical effectiveness of the rehabilitation program.

Methods

The home based exercise program consists of strengthening, balance and walking exercises. Participants have to perform the exercises 5 times a week for 12 weeks. They will have weekly telephone contact with a coach. The exercise program is delivered by means of a tablet-pc. During the intervention and measurements participants wear a motion sensor.

Measurements

Measurements will be done preoperative (T0), postoperative at 4 weeks (T1), 12 weeks (T2) and 26 weeks (T3). Primary outcome measure is feasibility determined by an evaluation questionnaire and adherence to the program. Secondary outcome measure is effectiveness measured by the HOOS, SF-36, EQ5D, Timed Up & Go Test and the Five Times Sit-to Stand Test.

Study objective

To determine the feasibility of a home based rehabilitation program delivered by means of a tablet-pc for patients after THA

To get a first impression of the clinical effectiveness of the rehabilitation program

Study design

Measurements will be done one week before the operation and 4 and 12 weeks postoperative. A follow up will be done 24 weeks postoperative.

Intervention

The home based exercise program consists of strengthening, balance and walking exercises. Participants have to perform the exercises 5 times a week for 12 weeks. They will have weekly telephone contact with a coach. The exercise program will be delivered by means of a tablet-pc. During the intervention patients wear a motion sensor.

Contacts

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Eligibility criteria

Inclusion criteria

Patients between 18 and 70 years, who live independently and are on a waiting list for a Total Hip Arthroplasty

Exclusion criteria

Patients with a revision surgery, cognitive problems, severe comorbidity and/or patients who do not master the Dutch language.

Study design

Design

Study type:	Interventional
Intervention model:	Parallel
Allocation:	Non controlled trial
Masking:	Open (masking not used)
Control:	Active

Recruitment

NL	
Recruitment status:	Other
Start date (anticipated):	01-06-2015
Enrollment:	30
Type:	Unknown

Ethics review

Positive opinion	
Date:	15-04-2015
Application type:	First submission

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
NTR-new	NL4879
NTR-old	NTR5150
Other	ABR: NL50372.042.14 : METc 2014/399

Study results