

Efficiency of immediate postoperative inpatient physical therapy (PT) following total knee arthroplasty: a randomised controlled trial (RCT)

Submission date 26/08/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 07/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 08/11/2022	Condition category Surgery	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

Efficiency of immediate postoperative inpatient physical therapy (PT) following total knee arthroplasty: a randomised controlled trial (RCT)

Study objectives

Does the intensity of PT treatment (once versus twice daily) in patients following TKA have an influence on short-term recovery, measured in range of motion and function?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Total knee arthroplasty in OA patients

Interventions

Intensive PT versus regular PT

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Primary outcome measures in this study were mobility and function

Secondary outcome measures

Secondary outcome measures were length of stay (LOS) following surgery, pain over the last 24 hours, satisfaction with treatment, satisfaction with the intermediate treatment results, number of PT sessions and Global perceived effect.

Overall study start date

15/01/2004

Completion date

01/08/2004

Eligibility

Key inclusion criteria

Patients who received a primary TKA at the University Hospital Maastricht between January 15 and June 1st, 2004. Subjects were considered eligible for the study if they were scheduled in the Joint Care program and signed an informed consent form.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

43

Total final enrolment

43

Key exclusion criteria

Patients undergoing knee revision surgery were excluded. Also patients over 85 years of age, patients with co-morbidity influencing gait and patient who did not understand or speak Dutch were excluded.

Date of first enrolment

15/01/2004

Date of final enrolment

01/08/2004

Locations

Countries of recruitment

Netherlands

Study participating centre
P.Debijelaan 25
Maastricht
Netherlands
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Sponsor information

Organisation

University Hospital Maastricht (The Netherlands)

Sponsor details

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Sponsor type

University/education

Website

<http://www.azm.nl>

ROR

<https://ror.org/02d9ce178>

Funder(s)

Funder type

University/education

Funder Name

University Hospital Maastricht (Netherlands)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

Not provided at time of registration

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article		31/08/2006		Yes	No