

# Circumpatellar electrocautery denervation in total knee arthroplasty without patellar resurfacing: a prospective randomized study with 1-year follow-up

Published: 11-03-2008

Last updated: 15-05-2024

To determine the effect of circumpatellar electrocautery denervation in total knee arthroplasty without patellar resurfacing.

<b>Ethical review</b>	Approved WMO
<b>Status</b>	Recruiting
<b>Health condition type</b>	Joint disorders
<b>Study type</b>	Interventional

## Summary

### ID

NL-OMON32154

### Source

ToetsingOnline

### Brief title

Patellar denervation in total knee arthroplasty.

### Condition

- Joint disorders
- Bone and joint therapeutic procedures

### Synonym

anterior knee pain, patellofemoral pain

### Research involving

Human

## Sponsors and support

**Primary sponsor:** Deventer Ziekenhuis

**Source(s) of monetary or material Support:** geen

## Intervention

**Keyword:** arthroplasty, knee prosthesis, patella

## Outcome measures

### Primary outcome

12 months, 2 years and 4 years after total knee arthroplasty:

(1) Prevalence of anterior knee pain using the Clinical Anterior Knee Pain Rating.

### Secondary outcome

12 months, 2 years and 4 years after total knee arthroplasty:

(2) Pain on a Visual Analogue Scale (VAS)

Before surgery, and 12 months, 2 years and 4 years after total knee arthroplasty:

(3) WOMAC 3.1 Osteoarthritis Index

(4) Knee Society Score

## Study description

### Background summary

Anterior knee pain occurs in up to 31% of patients after total knee arthroplasty without patellar resurfacing, and in approximately 16% of patients with patellar resurfacing.

The cause of patellofemoral pain after total knee arthroplasty is unknown but may at least be partly related to postoperative malalignment and anatomic

variations resulting in high patellofemoral contact forces. Immunohistochemical studies on innervation of the anterior knee have found hyperinnervation in the peripatellar soft tissues. In theory, circumpatellar denervation could result in a lower prevalence of anterior knee pain.

To date, only one clinical study addressed the results of patellar denervation in total knee arthroplasty. Although the results showed better pain relief after patellar denervation, the number of patients randomized was small (40), and no blinding was used. Randomized controlled trials to clarify the role of patellar resurfacing during total knee arthroplasty used 200 patients or more to achieve power to detect a significant difference.

### **Study objective**

To determine the effect of circumpatellar electrocautery denervation in total knee arthroplasty without patellar resurfacing.

### **Study design**

Randomized prospective with intervention, blinded.

### **Intervention**

Circumpatellar electrocautery denervation in total knee arthroplasty without patellar resurfacing.

### **Study burden and risks**

None

## **Contacts**

### **Public**

Deventer Ziekenhuis

Postbus 5001  
7400 GC Deventer  
Nederland

### **Scientific**

Deventer Ziekenhuis

Postbus 5001  
7400 GC Deventer  
Nederland

## Trial sites

### Listed location countries

Netherlands

## Eligibility criteria

### Age

Adults (18-64 years)

Elderly (65 years and older)

### Inclusion criteria

Not applicable.

### Exclusion criteria

Isolated patellofemoral osteoarthritis

Contralateral patellar resurfacing (primary or as revision)

Insufficient Dutch language skills

Total hip replacement or contralateral total knee replacement within the study period or less than one year before entering the study

Inflammatory arthritis

History of patellar fracture

Prior patellectomy

Patellofemoral instability

Prior unicondylar knee replacement

Previous high tibial or distal femoral osteotomy

Any prior operation involving the extensor mechanism

A severe medical disability that limits the ability to walk

Disabling disease involving other joints of the lower extremities

## Study design

### Design

Study type: Interventional

Intervention model: Parallel  
Allocation: Randomized controlled trial  
Masking: Double blinded (masking used)

**Primary purpose:** Treatment

## Recruitment

NL  
Recruitment status: Recruiting  
Start date (anticipated): 08-04-2008  
Enrollment: 200  
Type: Actual

## Ethics review

Approved WMO  
Date: 11-03-2008  
Application type: First submission  
Review commission: METC Isala Klinieken (Zwolle)

## Study registrations

### Followed up by the following (possibly more current) registration

No registrations found.

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

ID: 24792  
Source: NTR  
Title:

### In other registers

Register	ID
CCMO	NL21569.075.08
OMON	NL-OMON24792