# Anterior knee pain after total knee replacement: A systematic review.

No registrations found.

Ethical review	Not applicable
Status	Recruiting
Health condition type	-
Study type	Observational non invasive

## **Summary**

## ID

NL-OMON20661

Source NTR

**Brief title** N/A

#### **Health condition**

Total knee replacement.

## **Sponsors and support**

**Primary sponsor:** Deventer Ziekenhuis, Department of Orthopaedic Surgery **Source(s) of monetary or material Support:** Deventer Ziekenhuis, Department of Orthopaedic Surgery

## Intervention

#### **Outcome measures**

#### **Primary outcome**

Factors associated with anterior knee pain after primary total knee replacement.

#### Secondary outcome

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Outcome instruments used to assess the presence of anterior knee pain after total knee replacement.

# **Study description**

#### **Background summary**

Anterior knee pain is reported to occur in 4–49% of patients following primary total knee replacement. The cause is unknown, but may be related to patient characteristics, degree of patellar cartilage wear, prosthetic design, operative technique, and the use of patellar resurfacing.

#### **Study objective**

The objective of the study is to identify factors associated with anterior knee pain after primary total knee replacement.

#### Study design

N/A

#### Intervention

Systematic literature review using MEDLINE, the Cochrane Database of Systematic Reviews, the Cochrane Central Register of Controlled Trials, and EMBASE. Additional studies will be identified by reviews of the bibliographies of eligible articles.

# Contacts

#### Public

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# **Eligibility criteria**

## **Inclusion criteria**

- 1. Studies involving patients undergoing primary total knee replacement;
- 2. RCTs or cohort studies including a minimum of 10 patients;
- 3. Studies relating the prevalence of anterior knee pain with one or more variables;
- 4. English-language studies.

## **Exclusion criteria**

1. Case reports;

2. RCTs or cohort studies with incompletely described patient populations, less than 10 included patients, and insufficient descriptions of treatment.

# Study design

## Design

Study type:Observational non invasiveIntervention model:ParallelAllocation:Non controlled trialControl: N/A , unknownVariable

## Recruitment

NL	
Recruitment status:	Recruiting
Start date (anticipated):	01-01-2011

Enrollment:	
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Type:

0 Anticipated

# **Ethics review**

Not applicable Application type:

Not applicable

# **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	NL2552
NTR-old	NTR2670
Other	: N/A
ISRCTN	ISRCTN wordt niet meer aangevraagd.

# **Study results**

Summary results N/A