

# Patient Reported Outcomes in Charnley's Hip Replacement Arthroplasty.

No registrations found.

<b>Ethical review</b>	Positive opinion
<b>Status</b>	Recruiting
<b>Health condition type</b>	-
<b>Study type</b>	Observational non invasive

## Summary

### ID

NL-OMON22815

### Source

NTR

### Health condition

Hip Replacement Arthroplasty  
Patient Reported Outcomes  
Charnley  
Quality of Life

## Sponsors and support

**Primary sponsor:** Leiden University Medical Center

**Source(s) of monetary or material Support:** Dutch Arthritis Association (Reumafonds)

## Intervention

## Outcome measures

### Primary outcome

Types of PROs used.

### Secondary outcome

Prevalence of PROs in time.

## Study description

### Background summary

Outcome assessment in Hip Replacement Arthroplasty (HRA) traditionally consists of the physicians' assessment of pain, range of motion, daily activities and radiographs. However, it is generally acknowledged that Patient Reported Outcomes (PROs) may reveal the patients' burden of disease more accurately than physicians' assessments. The objective of this study is to assess which PROs are used in follow-up studies of Charnley's HRA and to assess the prevalence of reporting PROs in time.

### Study objective

The objective of this study is to assess which Patient Reported Outcomes are used in follow-up studies of Charnley's Hip Replacement Arthroplasty and to assess the prevalence of reporting Patient Reporting Outcomes in time.

### Study design

N/A

### Intervention

Systematic review, using MEDLINE, the Cochrane Controlled Trials Register (1960-2009), EMBASE (1991-2009), Web of Knowledge and CINAHL. Additional studies will be identified by hand searches of journal databases and conference abstracts.

## Contacts

### Public

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## Eligibility criteria

### Inclusion criteria

Randomised trials or cohort studies of Charnley's HRA, which report outcomes after 12 months or longer.

### Exclusion criteria

Studies reporting multiple types of prostheses, which are not clearly defined.

## Study design

### Design

Study type:	Observational non invasive
Intervention model:	Parallel
Allocation:	Non controlled trial
Masking:	Open (masking not used)
Control:	N/A , unknown

### Recruitment

NL	
Recruitment status:	Recruiting
Start date (anticipated):	01-07-2009
Enrollment:	1885
Type:	Anticipated

## Ethics review

Positive opinion	
Date:	11-02-2010
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	NL2093
NTR-old	NTR2210
Other	:
ISRCTN	ISRCTN wordt niet meer aangevraagd.

## Study results

### Summary results

N/A