# Prospective randomised single blind trial comparing the effectiveness of combined arthroscopic washout and intra articular hyaluronan injection to intra articular hyaluronan injection and arthroscopic washout in isolation, for osteoarthritis of knee

Recruitment status	<ul><li>Prospectively registered</li></ul>
Stopped	☐ Protocol
Overall study status	Statistical analysis plan
Stopped	Results
Condition category	Individual participant data
Musculoskeletal Diseases	Record updated in last year
	Overall study status Stopped Condition category

# Plain English summary of protocol

Not provided at time of registration

# **Contact information**

Type(s)

Scientific

#### Contact name

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#### Contact details

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## Additional identifiers

**EudraCT/CTIS** number

**IRAS** number

## ClinicalTrials.gov number

## Secondary identifying numbers

N0084105632

# Study information

#### Scientific Title

## **Study objectives**

Comparing the results of arthroscopic knee washout followed by intra articular hyaluronan (HA) injection to that of arthroscopic washout and intra articular HA injection in isolation.

## 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)

**Not Specified** 

## Participant information sheet

## Health condition(s) or problem(s) studied

Musculoskeletal Diseases: Osteoarthritis of knee

## **Interventions**

- 1. Arthroscopic washout
- 2. Intra articular hyaluronan injection
- 3. Arthroscopic washout and intra articular hyaluronan injection combined

This trial was stopped due to a lack of funding.

## Intervention Type

Other

## **Phase**

**Not Specified** 

## Primary outcome measure

Not provided at time of registration

## Secondary outcome measures

Not provided at time of registration

## Overall study start date

01/01/2002

## Completion date

31/03/2005

## Reason abandoned (if study stopped)

Lack of funding

# **Eligibility**

## Key inclusion criteria

Not provided at time of registration

## Participant type(s)

**Patient** 

## Age group

**Not Specified** 

#### Sex

**Not Specified** 

## Target number of participants

219 patients are required.

## Key exclusion criteria

Not provided at time of registration

## Date of first enrolment

01/01/2002

## Date of final enrolment

31/03/2005

# **Locations**

## Countries of recruitment

England

## **United Kingdom**

Study participating centre
Orthopaedic Department
Hull
United Kingdom
HU3 2JZ

# Sponsor information

## Organisation

Department of Health (UK)

## Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL

## Sponsor type

Government

## Website

http://www.doh.gov.uk

# Funder(s)

## Funder type

Government

## Funder Name

The North and South Bank Research and Development Consortium (UK)

# **Results and Publications**

## Publication and dissemination plan

Not provided at time of registration

## Intention to publish date

# Individual participant data (IPD) sharing plan

**IPD sharing plan summary**Not provided at time of registration