

The arcOGEN study: a genome wide association study of osteoarthritis

Submission date 23/04/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 27/05/2010	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
Last Edited 22/11/2013	Condition category Musculoskeletal Diseases	<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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NE1 4LP

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
4628

Study information

Scientific Title

A non-randomised observational trial in participants with primary osteoarthritis who will be used for mapping osteoarthritis susceptibility genes by genome-wide association scan

Acronym

arcOGEN

Study objectives

A collection of participants with primary osteoarthritis who will be used for mapping osteoarthritis susceptibility genes by genome-wide association scan and by other genetic approaches.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Oxfordshire Research Ethics Committee (REC) C approved on the 3rd January 2008 (ref: 07/H0606/150)

Study design

Non-randomised observational trial in primary, secondary and tertiary centres

Primary study design

Observational

Secondary study design

Multi-centre

Study setting(s)

Other

Study type(s)

Screening

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Topic: Musculoskeletal; Subtopic: Musculoskeletal (all Subtopics); Disease: Musculoskeletal

Interventions

There is no intervention in this trial. Recruited participants have a blood sample taken for a DNA test and a questionnaire will be answered. No treatments are given.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Questionnaire, completed at time of blood sampling

Secondary outcome measures

No secondary outcome measures

Overall study start date

01/03/2008

Completion date

31/12/2010

Eligibility

Key inclusion criteria

1. Primary osteoarthritis ascertained by joint replacement of the hip or the knee
2. Male and female, lower age limit of 40 years

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Planned Sample Size: 8000; UK Sample Size: 8000

Key exclusion criteria

Secondary forms of osteoarthritis or non-osteoarthritic diseases that result in joint replacement surgery

Date of first enrolment

01/03/2008

Date of final enrolment

31/12/2010

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

The Medical School
Newcastle
United Kingdom
NE1 4LP

Sponsor information

Organisation

Newcastle upon Tyne Hospitals NHS Foundation Trust (UK)

Sponsor details

Wolfson Unit of Clinical Pharmacology
Institute of Cellular Medicine
Framlington Place
Newcastle
England
United Kingdom
NE2 4HH

Sponsor type

Hospital/treatment centre

Website

<http://www.newcastle-hospitals.org.uk/>

ROR

<https://ror.org/05p40t847>

Funder(s)

Funder type

Charity

Funder Name

Arthritis Research Campaign (ARC) (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/05/2011		Yes	No
Results article	results	01/09/2012		Yes	No