# Patient specific biological responses to total joint replacements

| Submission date   | Recruitment status                                 | <ul><li>Prospectively registered</li></ul> |
|-------------------|--|--|
| 24/06/2010        | No longer recruiting                               | ☐ Protocol                                 |
| Registration date | Overall study status                               | Statistical analysis plan                  |
| 24/06/2010        | Completed  | Results                                    |
| Last Edited       | <b>Condition category</b> Musculoskeletal Diseases | Individual participant data                |
| 06/12/2019        |  | Record updated in last year                |

#### Plain English summary of protocol

Not provided at time of registration

# Contact information

#### Type(s)

Scientific

#### Contact name

Mr MH Stone

#### Contact details

Great George Street Leeds United Kingdom LS1 3EX +44 113 392 4769 martin.stone@leedsth.nhs.uk

# Additional identifiers

Protocol serial number 3298

# Study information

#### Scientific Title

Patient specific biological responses to total joint replacements

#### **Acronym**

Patient specific biological responses to total joint replacements

#### **Study objectives**

This research will be carried out at the University of Leeds and is a collaborative research programme between the Institute of Medical and Biological Engineering and Consultant Orthopaedic Surgeons, Mr Martin Stone (Chapel Allerton) and Mr David Shaw (Bradford Royal Infirmary). Patients will be recruited from two groups, patients awaiting hip replacement operations and patients with long-term polyethylene implants. Approximately 100 patients from each group will be studied.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

MREC approved (ref: CA03/008)

#### Study design

Single centre non-randomised interventional treatment trial

#### Primary study design

Interventional

#### Study type(s)

Treatment

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

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

#### **Interventions**

Patients awaiting hip replacement will have an additional 30 ml of blood collection preoperatively. Inclusion in the study will not affect patient care in any way, and the surgical procedure will be the same whether the patient chooses to participate in the study or not. Patients with long-term implants will be invited by letter to attend the clinic to volunteer the additional 30 ml blood sample.

The blood samples will be processed by the researchers at the University of Leeds, where the white blood cells will be isolated and used in culture with real wear particles generated by articulation. Cytokine production will be measured by enzyme-linked immunospecific assay (ELISA). Approximately 1 x 10^6 cells will be stored for genetic analysis to determine specific base changes in the DNA. The clinical results of the patients will be followed in follow-up clinics.

The control group will comprise approximately 30 healthy volunteers from the research group (iMBE, Faculty of Biological Sciences & School of Mechanical Engineering). A single 30 ml blood sample will be collected from each control subject by trained personel. Collection of blood samples for research purposes is performed routinely and approval has been awarded from the Faculty of Biological Sciences Research Ethics Committee.

#### **Intervention Type**

Other

#### Phase

Not Applicable

#### Primary outcome(s)

Cytokine released by primary human macrophages from individual patients in response to 100 cubic micrometers of polyethylene.

### Key secondary outcome(s))

- 1. Correlation of genotype with clinical outcome of THR surgery
- 2. Correlation of phenotypic response to cytokine promoter polymorphism genotype
- 3. Number of patients that are low responders, intermediate responders and aggressive responders to wear particles

#### Completion date

30/06/2010

# **Eligibility**

#### Key inclusion criteria

Patients with a Charnley hip prosthesis that has been in situ for 10 years

#### Participant type(s)

**Patient** 

#### Healthy volunteers allowed

No

#### Age group

Not Specified

#### Sex

#### Key exclusion criteria

Not provided at time of registration

#### Date of first enrolment

01/09/2009

#### Date of final enrolment

30/06/2010

# Locations

#### Countries of recruitment

United Kingdom

England

# Study participating centre Great George Street

Leeds

# Sponsor information

#### Organisation

University of Leeds (UK)

#### **ROR**

https://ror.org/024mrxd33

# Funder(s)

#### Funder type

Research council

#### **Funder Name**

Engineering and Physical Sciences Research Council (EPSRC) (UK)

## Alternative Name(s)

EPSRC Engineering & Physical Sciences Research Council, UKRI Engineering and Physical Sciences Research Council, Engineering and Physical Sciences Research Council - UKRI, Engineering & Physical Sciences Research Council, The Engineering and Physical Sciences Research Council (EPSRC), EPSRC

#### Funding Body Type

Government organisation

## Funding Body Subtype

National government

#### Location

United Kingdom

# **Results and Publications**

Individual participant data (IPD) sharing plan

IPD sharing plan summary