

Forefoot arthroplasty in rheumatoid arthritis

Submission date 11/12/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 11/12/2002	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 02/02/2017	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

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

A0524

Study information

Scientific Title

Forefoot arthroplasty in rheumatoid arthritis

Study objectives

Not provided at time of registration

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)

Hospital

Study type(s)

Treatment

Participant information sheet**Health condition(s) or problem(s) studied**

Rheumatoid arthritis

Interventions

Randomised to have one of two types of forefoot arthroplasty:

1. Modified Kates Kessel Kay type (excision of metatarsal heads through plantar incision)
2. Modified Stainsby procedure (excision base of proximal phalanges through dorsal incisions).

All patients underwent arthrodesis of the 1st metatarsophalangeal joint.

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

01/04/1997

Completion date

31/03/2000

Eligibility

Key inclusion criteria

Patients with rheumatoid arthritis with forefoot involvement

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/04/1997

Date of final enrolment

31/03/2000

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Avon Orthopaedic Centre

Bristol

United Kingdom

BS10 5NB

Sponsor information

Organisation

Arthritis Research Campaign (ARC) (UK)

Sponsor details

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Sponsor type

Charity

Website

<http://www.arc.org.uk>

ROR

<https://ror.org/02jkpm469>

Funder(s)**Funder type**

Charity

Funder Name

Arthritis Research Campaign (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

