

Induction of true remission in early poor prognosis (erosive) rheumatoid arthritis (RA) - a randomised study of methotrexate and intra-articular corticosteroids or placebo

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|--|---|--|
| Submission date 15/07/2002 | Recruitment status No longer recruiting | <input type="checkbox"/> Prospectively registered |
| Registration date 15/07/2002 | Overall study status Completed | <input type="checkbox"/> Protocol |
| Last Edited 03/10/2007 | Condition category Musculoskeletal Diseases | <input type="checkbox"/> Statistical analysis plan |
| | | <input checked="" type="checkbox"/> Results |
| | | <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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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

E0516

Study information

Scientific Title

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Early rheumatoid arthritis

Interventions

1. Receive combination therapy (n = 40)
2. Receive sulfasalazine (SSZ) alone (n = 42)

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

01/06/1995

Completion date

01/03/1997

Eligibility

Key inclusion criteria

1. All patients to meet the American College of Rheumatology criteria for RA
2. Poor prognosis disease according to criteria based on the presence of factors known to be predictive of poor long-term outcome
3. Adequate contraception a prerequisite

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

82

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/06/1995

Date of final enrolment

01/03/1997

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Rheumatology and Rehabilitation Research Unit
Leeds

United Kingdom
LS2 9NZ

Sponsor information

Organisation

Arthritis Research Campaign (ARC) (UK)

Sponsor details

Copeman House
St Mary's Court
St Mary's Gate
Chesterfield
Derbyshire
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S41 7TD

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info@arc.org.uk

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

Study outputs

| Output type | Details | Date created | Date added | Peer reviewed? | Patient-facing? |
|---------------------------------|---------|--------------|------------|----------------|-----------------|
| Results article | Results | 01/08/2000 | | Yes | No |