

A short course of low dose steroid tablets as a treatment for painful hand osteoarthritis- does it work?

Submission date 18/11/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 09/05/2011	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 10/09/2012	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

Protocol serial number
RR09/8877

Study information

Scientific Title

Treating painful hand osteoarthritis using low dose oral prednisolone: assessing short-term pain and imaging outcomes in a randomised, placebo-controlled, double-blind, 12 week, single centre study

Acronym

POLO

Study objectives

Current treatments for osteoarthritis (OA) have major limitations and other analgesic treatments are needed. Synovitis is prevalent in OA and previous studies have shown it to be a source of pain. Corticosteroids are used in routine practice at treating synovitis in both the inflammatory arthritides and in osteoarthritis and have been shown to be effective at reducing pain.

We propose that treating patients with moderate to severe OA hand symptoms with low dose oral prednisolone will be a practical and safe treatment to reduce synovitis and therefore reduce pain. This will potentially introduce a new treatment into the OA armamentarium which could be of particular use in the primary care setting.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Leeds (West) Research Ethics Committee approved on the 22nd May 2009 (ref: 09/H1307/53)

Study design

Randomised placebo-controlled double-blind single centre study

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Painful hand osteoarthritis

Interventions

Patients will be randomised to either one of the following regimens:

1. One capsule containing 5 milligrams of prednisolone daily for 28 days orally
2. Placebo capsule one daily for 28 days orally

The total trial duration will be 12 weeks; 4 weeks of treatment then a follow-up visit 8 weeks later.

Intervention Type

Drug

Phase

Phase IV

Drug/device/biological/vaccine name(s)

Prednisolone

Primary outcome(s)

Change in hand pain VAS, measured from 0 mm (no pain) to 100 mm (severe pain) from baseline to 4 weeks

Key secondary outcome(s)

1. Change in functional scores (including the Australian/Canadian Osteoarthritis Hand Index [AUSCAN] functional subscale) at 4 and 12 weeks. Measured using the VAS AUSCAN 0 - 100 mm whereby 0 mm = no difficulty and 100 mm = extreme difficulty
2. Change in pain scores (including VAS global disease activity score and AUSCAN pain subscale) at 4 and 12 weeks. Disease activity scores are measured from 0 mm (no activity) to 100 mm (extreme activity). Pain scores are measured from 0 mm (no pain) to 100 mm (extreme pain).
3. Change in Osteoarthritis Quality of Life questionnaire (OAQoL) scores at 12 weeks
4. Change in Hospital Anxiety and Depression Scale (HADS) scores at 4 and 12 weeks
5. Changes on extremity MRI scan at baseline and 4 weeks. MRIs will be scored for synovitis using a semi-quantitative scoring system.

Completion date

31/12/2010

Eligibility**Key inclusion criteria**

1. Patients must fulfil the American College of Rheumatology criteria for OA of the small joints of the hand
2. Moderately severe symptoms (greater than or equal to 40/100 on hand pain visual analogue scale [VAS] at screening)
3. Symptoms present on most days over the last 3 months
4. A previous radiograph of the hands with changes consistent with osteoarthritis
5. Stable analgesic requirements (including non-steroidal anti-inflammatory drugs [NSAIDs]) for at least 4 weeks
6. Stable doses of chondroitin or glucosamine for 4 months
7. No oral, intramuscular [IM], intra-arterial [IA], or intravenous [IV] steroids during the last 3 months
8. The patient must be able to adhere to the study visit schedule and other protocol requirements
9. The patient must be capable of giving informed consent and the consent must be obtained prior to any screening procedures
10. Aged 43 - 81 years, either sex

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. The presence of an inflammatory arthritis
2. Sensitivity, anaphylaxis or allergy to prednisolone
3. Patients who are pregnant, lactating or using hormonal birth control pills
4. Currently uncontrolled diabetes
5. Currently uncontrolled hypertension
6. Current active infection
7. Surgical procedure within 30 days of study initiation
8. Patients with osteoporosis or taking bisphosphonates
9. Patients will not be eligible if they have any contraindications to magnetic resonance imaging (MRI) scanning:
 - 9.1. Pacemakers
 - 9.2. Surgical clips within the head
 - 9.3. Certain inner ear implants
 - 9.4. Neuro-electrical stimulators
 - 9.5. Metal fragments within the eye or head
 - 9.6. Pregnant or breastfeeding women
10. Use of any investigational (unlicensed) drug within 1 month prior to screening or within 5 half-lives of the investigational agent, whichever is longer
11. Subjects with any uncontrolled, unstable or severe medical condition, which in the opinion of the investigator makes them unsuitable for the study

Date of first enrolment

17/08/2009

Date of final enrolment

31/12/2010

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre**Section of Musculoskeletal Disease**

Leeds

United Kingdom

LS7 4SA

Sponsor information

Organisation

University of Leeds (UK)

ROR

<https://ror.org/024mrx33>

Funder(s)

Funder type

University/education

Funder Name

University of Leeds (UK)

Alternative Name(s)

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

United Kingdom

Results and Publications

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	01/12/2012		Yes	No
HRA research summary			28/06/2023	No	No