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Comparing the Tolerability and Efficacy of Ostenil Mini in small joint osteoarthritis (OA) in the Carpus compared to a corticosteroid.

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [X] Results [ ] Individual participant data
Registration date 30/09/2004	**Overall study status** Completed	
Last Edited 09/09/2015	**Condition category** Musculoskeletal Diseases	

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

N0065120984

Study information

Scientific Title
Comparing the Tolerability and Efficacy of Ostenil Mini in small joint osteoarthritis (OA) in the Carpus compared to a corticosteroid.

Study objectives
To compare the tolerability and efficacy of Ostenil Mini in small joint osteoarthritis (OA) in the Carpus compared to a corticosteroid.

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)
Treatment

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
Small joint osteorathritis

Interventions
Randomised trial using two groups and questionnaires. Ostenil Mini in small joint OA in the Carpus compared to a corticosteroid

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
1. Ostenil Mini 2. Corticosteroid

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/03/2003

Completion date
01/12/2003

Eligibility

Key inclusion criteria
40 patients with OA of the carpus (20 in each group) will be recruited from patients seeking treatment from the rheumatology and upper limb orthopaedic departments.

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
40

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
01/03/2003

Date of final enrolment
01/12/2003

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Department of Orthopaedics
Sunderland Royal Hospital
Kayll Roa
Sunderland
United Kingdom
SR4 7TP

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
City Hospitals Sunderland NHS Trust (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:	01/09/2012		Yes	No