ISRCTN25275611 https://doi.org/10.1186/ISRCTN25275611

Prospective randomised double-blind placebo-controlled trial to establish the effectiveness of glucosamine sulphate and chondroitin sulphate in the treatment of non-specific mechanical low back pain

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 30/09/2004	**Overall study status** Completed	
Last Edited 14/11/2014	**Condition category** Musculoskeletal Diseases	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mr A Mohsen

Contact details
Orthopaedic Department
Hull Royal Infirmary
Anlaby Road
Hull
United Kingdom
HU3 2JZ
+44 (0)1482 328541

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0084089157

Study information

Scientific Title

Study objectives
Is glucosamine sulphate/chondroitin sulphate a useful tool in the armamentarium of the low back pain specialist in treating low back pain regardless of various combinations of biopsychosocial factors being present?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Prospective randomised double-blind placebo-controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Musculoskeletal Diseases: Low back pain

Interventions
Glucosamine sulphate and chondroitin sulphate vs placebo

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Glucosamine sulphate and chondroitin sulphate

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/08/2000

Completion date
01/01/2004

Eligibility

Key inclusion criteria
All eligible male and female patients with back pain would be invited to participate in the trial

Participant type(s)
Patient

Age group
Not Specified

Sex
Both

Target number of participants
200 patients, 100 in each arm of the study

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/08/2000

Date of final enrolment
01/01/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Orthopaedic Department
Hull
United Kingdom
HU3 2JZ

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
Government

Funder Name
The North and South Bank Research and Development Consortium (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