

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	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 14/11/2014	Condition category Musculoskeletal Diseases	<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

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Additional identifiers

Protocol serial number

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

Study type(s)

Treatment

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(s)

Not provided at time of registration

Key secondary outcome(s))

Not provided at time of registration

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

Healthy volunteers allowed

No

Age group

Not Specified

Sex

All

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

United Kingdom

England

Study participating centre

Orthopaedic Department

Hull

United Kingdom

HU3 2JZ

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Government

Funder Name

The North and South Bank Research and Development Consortium (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration