

Epidural and Nerve Root Corticosteroid and Local Anaesthetic Injections for Lumbar Nerve Root Compression

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 14/03/2014	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0199129906

Study information

Scientific Title

Study objectives

To compare efficacy in pain relief

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)

Hospital

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Musculoskeletal Diseases: Spinal stenosis

Interventions

Randomised, double blind. 50 patients with proplapsed disc, 50 patients with spinal stenosis. Each group randomised into 2 treatment arms (25 in each):

1. Nerve root injection of local anaesthetic and corticosteroid
2. Epidural injection of local anaesthetic and corticosteroid

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Epidural and Nerve Root Corticosteroid and Local Anaesthetic Injections

Primary outcome measure

Pain relief on a visual analogue scale

Secondary outcome measures

Not provided at time of registration

Overall study start date

15/01/2003

Completion date

31/12/2007

Eligibility

Key inclusion criteria

100 patients with diagnosis of prolapsed disc or spinal stenosis of the lumbar-sacral vertebrae. Unilateral pain radiating from back to below knee lasting 6-28 weeks, with leg pain being greater than back pain.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

100

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

15/01/2003

Date of final enrolment

31/12/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Orthopaedic Department
Reading
United Kingdom
RG1 5AN

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
Royal Berkshire and Battle Hospitals 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