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# A comparison of three methods for securing and dressing epidural catheters used for epidural infusion analgesia (EIA)

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| **Submission date**<br>12/09/2003 | **Recruitment status**<br>No longer recruiting | [ ] Prospectively registered<br>[ ] Protocol |
| **Registration date**<br>12/09/2003 | **Overall study status**<br>Completed | [ ] Statistical analysis plan<br>[ ] Results |
| **Last Edited**<br>28/02/2020 | **Condition category**<br>Signs and Symptoms | [ ] Individual participant data<br>[ ] 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

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**

N0192104704

# Study information

### Scientific Title
A comparison of three methods for securing and dressing epidural catheters used for epidural infusion analgesia (EIA)

### Study objectives
Is there a system of securing and dressing an epidural catheter for epidural infusion analgesia that is more efficacious than current practice?

### 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)
Treatment

### Participant information sheet

### Health condition(s) or problem(s) studied
Signs and Symptoms: Pain

### Interventions
System of securing and dressing an epidural catheter for epidural infusion analgesia vs current practice

### Intervention Type
Other

### Phase
Not Applicable

### Primary outcome measure

Ease of operator use, system adhesive and aseptic integrity, patient comfort measured using a 5-point Likert scale.

### Secondary outcome measures
Not provided at time of registration

### Overall study start date
11/07/2001

### Completion date
14/01/2002

## Eligibility

### Key inclusion criteria
Not provided at time of registration

### Participant type(s)
Patient

### Age group
Not Specified

### Sex
Not Specified

### Target number of participants
Total number of subjects = 180

### Key exclusion criteria
Not provided at time of registration

### Date of first enrolment
11/07/2001

### Date of final enrolment
14/01/2002

## Locations

### Countries of recruitment
England

United Kingdom

### Study participating centre

**Pain Management Centre**
Nottingham
United Kingdom
NG7 2UH

# Sponsor information

**Organisation**
Department of Health (UK)

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

**Sponsor type**
Government

**Website**
http://www.doh.gov.uk

# Funder(s)

**Funder type**
Government

**Funder Name**
Queen's Medical Centre University Hospital 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