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# Exploring the value for patients of audiotaping the revascularisation consultation. A pilot study.

**Submission date**
30/09/2004

**Recruitment status**
No longer recruiting

**Registration date**
30/09/2004

**Overall study status**
Completed

**Last Edited**
15/04/2010

**Condition category**
Surgery

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

**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**
N0071122002

# Study information

### Scientific Title

### Study objectives
What is the value for patients of receiving an audiotape of their revascularisation consultation?

### 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)
Other

### Participant information sheet

### Health condition(s) or problem(s) studied
Surgery: Myocardial revascularisation

### Interventions
The pilot study will involve a randomised controlled trial, comparing the experiences and outcomes among patients receiving an audiotape of their revascularisation consultation with patients who do not receive an audiotape of their consultation.

### Intervention Type
Procedure/Surgery

### Phase
Not Specified

### Primary outcome measure
Assessment of spontaneous recall through use of open-ended questions to ask the patient to report what the surgeon said during the consultation.

### Secondary outcome measures

Not provided at time of registration

### Overall study start date
01/05/2002

### Completion date
01/10/2003

## Eligibility

### Key inclusion criteria
Patients on the waiting list for an appointment with the named cardiothoracic surgeon, will be identified by the research nurse. On average, five patients are referred to the Consultant Surgeon for consultation regarding revascularisation surgery, each week. To allow for patients who decline to participate in the study, the study will aim to recruit 20 patients over an eight-week period. Mapping postcodes on to available deprivation indicators will identify the socio economic background of the sample. A subsequent full-scale study would be stratified to ensure enhanced representation from patients living in areas of socio economic deprivation.

### Participant type(s)
Patient

### Age group
Adult

### Sex
Both

### Target number of participants
20

### Key exclusion criteria
Does not match inclusion criteria

### Date of first enrolment
01/05/2002

### Date of final enrolment
01/10/2003

## Locations

### Countries of recruitment
England

United Kingdom

### Study participating centre

**City Wide Initiative for Reducing Cardiovascular Disease (CIRC/CHS)**
Sheffield
United Kingdom
S10 3TH

# 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**
Research organisation

**Funder Name**
Sheffield Health and Social Research 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

## Study outputs

| Output type | Details | Date created | Date added | Peer reviewed? | Patient-facing? |
|---|---|---|---|---|---|
| Results article | results | 01/11/2005 | | Yes | No |