Assessment of pain and other benefits following ministernotomy for cardiac surgery: a prospective randomised trial

Submission date	Recruitment status	Prospectively registered
30/09/2004	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2004	Completed	Results
Last Edited	Condition category	Individual participant data
18/10/2017	Surgery	[] Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Mr U R Nair

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Assessment of pain and other benefits following ministernotomy for cardiac surgery: a prospective randomised trial

Study objectives

To determine the benefit obtained in patients who underwent ministernotomy for cardiac surgery.

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

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Surgery: Cardiovascular

Interventions

Random allocation to [a] treatment a [b] treatment b (Small chest incision vs full chest incision).

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Pain, total blood loss, analgesia requirement, lung function.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/05/2003

Completion date

01/12/2008

Eligibility

Key inclusion criteria

All adult cardiac surgical cases under the care of Mr R U Nair requiring sternotomy.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/05/2003

Date of final enrolment

01/12/2008

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Yorkshire Heart Centre

Leeds United Kingdom LS1 3EX

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

Leeds Teaching 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