

Is phenoxybenzamine bathing improving the in situ flow and conduit size in harvested radial arteries for coronary surgery? A pilot clinical trial

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 18/07/2016	Condition category Surgery	<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

Mr Rex Stanbridge

Contact details

Cardiothoracic Surgery
St Mary's Hospital
Praed Street
London
United Kingdom
W2 1NY
+44 (0)20 7886 2212
rex.stanbridge@st-marys.nhs.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0241128292

Study information

Scientific Title

Is phenoxybenzamine bathing improving the in situ flow and conduit size in harvested radial arteries for coronary surgery? A pilot clinical trial

Study objectives

Does phenoxybenzamine protect or increase conduit blood flow in vivo during coronary bypass with radial artery?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Double-blind 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 contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Surgery: Cardiovascular

Interventions

Bathing with phenoxybenzamine during coronary bypass with radial artery

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome measure

Blood flow (ml/min)

Secondary outcome measures

Not provided at time of registration

Overall study start date

11/08/2003

Completion date

11/12/2003

Eligibility

Key inclusion criteria

24 patients undergoing coronary bypass surgery

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

24

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

11/08/2003

Date of final enrolment

11/12/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

St Mary's Hospital

London

United Kingdom
W2 1NY

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

Hospital/treatment centre

Funder Name

St Mary's 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