# Right ventricular function following coronary artery bypass graft

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
28/05/2010	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
01/10/2010	Completed	Results
Last Edited	Condition category	Individual participant data
30/09/2016	Nutritional, Metabolic, Endocrine	<ul><li>Record updated in last year</li></ul>

#### Plain English summary of protocol

Not provided at time of registration

## Contact information

#### Type(s)

Scientific

#### Contact name

Miss Beth Unsworth

#### Contact details

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# Additional identifiers

Protocol serial number

7353

# Study information

#### Scientific Title

Assessment of right ventricular function during and following cardiac surgery and evaluation of pericardial physiology in preserving right ventricular function

#### **Acronym**

**DRN 372** 

#### **Study objectives**

This is an observational study looking at the effects of cardiac surgery in right ventricular (RV) function in those patients referred for routine coronary artery bypass grafting (CABG) and aortic valve replacement (AVR) operations.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Sub-Committee of the REC, 19/01/2007, ref: 06/Q0403/163

#### Study design

Single-centre observational case-controlled study

#### Primary study design

Observational

#### Study type(s)

Treatment

#### Health condition(s) or problem(s) studied

Topic: Diabetes Research Network; Subtopic: Both; Disease: Cardiovascular disease

#### **Interventions**

Observational study using echocardiography to look at the immediate effects of cardiac surgery in RV function in those patients referred for routine CABG and AVR operations. Pre-, peri- and post-operative data collection.

Study entry: registration only

#### Intervention Type

Other

#### Phase

Not Applicable

#### Primary outcome(s)

Post-operative assessment of RV function; patients are last seen one month after surgery, but consented to return possibly at 6 months.

#### Key secondary outcome(s))

No secondary outcome measures

#### Completion date

30/09/2010

# **Eligibility**

#### Key inclusion criteria

- 1. Patients with chronic systolic heart failure confirmed by transthoracic echo
- 2. Male and female, aged 51 91 years

#### Participant type(s)

**Patient** 

#### Healthy volunteers allowed

No

#### Age group

Other

#### Sex

All

#### Key exclusion criteria

- 1. Patients with severe lung disease
- 2. Any condition that would preclude them from lying comfortably on a couch for the duration of the study

#### Date of first enrolment

01/05/2009

#### Date of final enrolment

30/09/2010

## Locations

#### Countries of recruitment

**United Kingdom** 

England

# Study participating centre Imperial College Healthcare NHS Trust

London United Kingdom W2 1LA

# Sponsor information

#### Organisation

Imperial College Healthcare NHS Trust (UK)

#### **ROR**

https://ror.org/056ffv270

# Funder(s)

#### Funder type

Charity

#### **Funder Name**

British Heart Foundation (BHF) (UK)

#### Alternative Name(s)

the\_bhf, The British Heart Foundation, BHF

#### **Funding Body Type**

Private sector organisation

#### **Funding Body Subtype**

Trusts, charities, foundations (both public and private)

#### Location

**United Kingdom** 

# **Results and Publications**

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?

Participant information sheet 11/11/2025 No Yes