# 'AQURO', a new, potentially automatable approach for quatifying mitral regurgitation: technology development and validation through collaboration between cardiovascular science and bioengineering

Submission date	Recruitment status No longer recruiting	<ul><li>Prospectively registered</li></ul>		
18/06/2010		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
18/06/2010	Completed	[X] Results		
<b>Last Edited</b> 27/08/2014	<b>Condition category</b> Circulatory System	Individual participant data		

# Plain English summary of protocol

Not provided at time of registration

#### Contact information

# Type(s)

Scientific

#### Contact name

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#### Contact details

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# Additional identifiers

#### Protocol serial number

7355

# Study information

#### Scientific Title

#### **Acronym**

**DRN 373 (AQURO)** 

#### Study objectives

Stable subjects with mitral regurgitation previously diagnosed on echocardiography will be recruited from the outpatient service of Imperial College Healthcare NHS Trust. We will follow the standard British Society of Echocardiography protocol, and then add additional views. The standard protocol involves ultrasound recordings in different positions around the heart. Each complete echocardiography appointment will take approximately 40 minutes. All calculations made from the recorded images will occur later, after the patient has left.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

MREC approved (ref: 08/H0707/71)

#### Study design

Single-centre non-randomised interventional diagnosis trial

#### Primary study design

Interventional

#### Study type(s)

Diagnostic

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

Topic: Diabetes Research Network; Subtopic: Type 2; Disease: Cardiovascular disease

#### **Interventions**

Patients are recruited and ultrasound scans are performed for about 40 minutes to be analysed and compared with the standard technique.

Study entry: registration only

#### Intervention Type

Other

#### Phase

Not Applicable

#### Primary outcome(s)

Effective regurgitant orifice area (EROA), measured during 40 minutes of acquisition

# Key secondary outcome(s))

Not provided at time of registration

#### Completion date

23/03/2012

# **Eligibility**

#### Key inclusion criteria

Stable subjects with mitral regurgitation previously diagnosed on echocardiography

#### Participant type(s)

**Patient** 

#### Healthy volunteers allowed

No

#### Age group

**Not Specified** 

#### Sex

**Not Specified** 

#### Key exclusion criteria

- 1. Atrial fibrillation
- 2. Moderate or severe disease of tricuspid or pulmonary valves
- 3. Any aortic valve disease graded mild or higher
- 4. Prosthetic aortic valve and body habitus or coexistent disease that precludes satisfactory imaging quality

#### Date of first enrolment

23/03/2009

#### Date of final enrolment

23/03/2012

# Locations

#### Countries of recruitment

United Kingdom

England

# Study participating centre St Mary's Hospital

London United Kingdom W2 1NY

# 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?
Results article	results	01/07/2013		Yes	No