

# Exploring the therapeutic potential of xanthine oxidase inhibitor allopurinol in angina

<b>Submission date</b> 21/02/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 29/03/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 08/11/2011	<b>Condition category</b> Circulatory System	<input type="checkbox"/> 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

**Protocol serial number**  
JUS002

## Study information

**Scientific Title**

**Study objectives**

Xanthine oxidase inhibition in chronic stable angina improves endothelial function.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Approved by the Regional Ethics Committee on 08/11/2005, reference number: 05/S1401/101

**Study design**

Randomised double blind placebo-controlled crossover trial

**Primary study design**

Interventional

**Study type(s)**

Treatment

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

Chronic stable angina

**Interventions**

Allopurinol versus placebo

**Intervention Type**

Drug

**Phase**

Not Specified

**Drug/device/biological/vaccine name(s)**

Allopurinol

**Primary outcome(s)**

Change in endothelial function as assessed by:

1. Flow-mediated dilatation
2. Forearm venous occlusion plethysmography

**Key secondary outcome(s)**

Changes in brain natriuretic peptide (BNP)

**Completion date**

02/04/2008

## **Eligibility**

**Key inclusion criteria**

1. Documented coronary artery disease
2. Chronic stable angina

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**Key exclusion criteria**

1. Left ventricular systolic dysfunction
2. Renal failure
3. Concomitant warfarin therapy
4. Allergy to allopurinol

**Date of first enrolment**

05/01/2006

**Date of final enrolment**

02/04/2008

## Locations

**Countries of recruitment**

United Kingdom

Scotland

**Study participating centre**

**Dept of Clinical Pharmacology**

Dundee

United Kingdom

DD1 9SY

## Sponsor information

**Organisation**

University of Dundee (UK)

**ROR**

<https://ror.org/03h2bxq36>

# Funder(s)

## Funder type

Charity

## Funder Name

The British Heart Foundation (UK)

# 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?
<a href="#">Results article</a>	results	16/08/2011		Yes	No