

Exploring the therapeutic potential of xanthine oxidase inhibitor allopurinol in angina

Submission date 21/02/2006	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 29/03/2006	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 08/11/2011	Condition category Circulatory System	☐ Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Narasimharajapura Rajendra

Contact details
Dept of Clinical Pharmacology
Level 7
Ninewells Hospital
Dundee
United Kingdom
DD1 9SY
+44 (0)1382 496355

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

Primary study design

Interventional

Study design

Randomised double blind placebo-controlled crossover trial

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?
Results article	results	16/08/2011		Yes	No