

Ultra low dose anticoagulation in atrial fibrillation

Submission date 23/01/2004	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 23/01/2004	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 30/08/2012	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
Prof Gareth Beevers

Contact details
University Department of Medicine
City Hospital
Dudley Rd
Birmingham
United Kingdom
B18 7QH
+44 (0)121 507 5080

Additional identifiers

Protocol serial number
MC7

Study information

Scientific Title

Study objectives

Previous studies have demonstrated increased markers of thrombogenesis in patients with atrial fibrillation (AF), suggesting the presence of a hypercoaguable or prothrombotic state. The objective of this study was to determine the effects of introducing ultra-low-dose warfarin (1 mg), conventional warfarin, and aspirin (300 mg) therapy on thrombogenesis and platelet activation in AF.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Heart disease

Interventions

3 treatment arms:

1. Ultra-low-dose warfarin (1 mg)
2. Conventional warfarin
3. Aspirin (300 mg)

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Warfarin, aspirin

Primary outcome(s)

Platelet activation

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/04/1996

Eligibility

Key inclusion criteria

Patients with atrial fibrillation

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/04/1994

Date of final enrolment

01/04/1996

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

University Department of Medicine

Birmingham

United Kingdom

B18 7QH

Sponsor information**Organisation**

Record Provided by the NHS R&D 'Time-Limited' National Programme Register - Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

NHS Cardiovascular Disease and Stroke National Research and Development Programme (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	01/08/1996		Yes	No