

Atrial pacing in paroxymal Atrial Fibrillation (AF) (ELA)

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 25/07/2012	Condition category Circulatory System	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr J Pitts-Crick

Contact details
c/o Research and Development Office
Level 1, Old Building
Bristol Royal Infirmary
Marlborough Street
Bristol
United Kingdom
BS2 8HW
+44 (0)117 928 3473

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0264058807

Study information

Scientific Title

Study objectives

Does atrial pacing at higher than spontaneous sinus rhythm (SR) prevent episodes of paroxymal AF?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised single blind controlled crossover group trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Prevention

Participant information sheet

Health condition(s) or problem(s) studied

Atrial fibrillation (AF)

Interventions

Implant diagnostic pacemakers and randomise cross over single blind programme to higher or lower rate.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Frequency and duration of episodes of AF.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/08/2000

Completion date

25/12/2005

Eligibility

Key inclusion criteria

Patients undergoing pacing and ablation for AF.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/08/2000

Date of final enrolment

25/12/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

c/o Research and Development Office

Bristol

United Kingdom

BS2 8HW

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)**Funder type**

Hospital/treatment centre

Funder Name

United Bristol Healthcare NHS Trust (UK)

Results and Publications**Publication and dissemination plan**

Not provided at time of registration

Intention to publish date**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration