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A randomised trial to assess the impact of an anti-thrombotic decision aid in patients with non-valvular atrial fibrillation

Submission date 20/04/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [X] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 21/04/2004	**Overall study status** Completed	
Last Edited 10/08/2007	**Condition category** Circulatory System	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
I-10

Study information

Scientific Title

Acronym
DAAFI

Study objectives
Not provided at time of registration

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Non-valvular atrial fibrillation

Interventions
A prospective, multi-centre, two-arm cluster randomised controlled trial to evaluate whether an evidence-based patient decision aid for patients with non-valvular atrial fibrillation can improve the appropriate use of antithrombotic therapy (as defined by the 2001 American College of Chest Physicians [ACCP] Recommendations) by patients and their family physicians compared to usual care.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/2004

Completion date
01/01/2005

Eligibility

Key inclusion criteria
Family physicians and their adult patients with non-valvular atrial fibrillation.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
1100

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/2004

Date of final enrolment
01/01/2005

Locations

Countries of recruitment
Canada

Study participating centre
2E3.24 Walter Mackenzie Centre
Edmonton
Canada
T6G 2R7

Sponsor information

Organisation
Canadian Stroke Network (Canada)

Sponsor details
451 Smyth Road
Ottawa
Canada
K1H 8M5

Sponsor type
Charity

Funder(s)

Funder type
Charity

Funder Name
Canadian Stroke Network (I-10) (Canada)

Funder Name
Alberta Heritage Foundation for Medical Research (Canada)

Alternative Name(s)
AHFMR

Funding Body Type
Government organisation

Funding Body Subtype
Local government

Location
Canada

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article	Protocol	05/05/2004		Yes	No
Results article	Results	30/08/2005		Yes	No