ISRCTN81818945 https://doi.org/10.1186/ISRCTN81818945

A prospective, multi-centre, randomised trial comparing low-dose aspirin with low-dose aspirin plus low-intensity oral anticoagulation in the primary prevention of thrombosis in patients positive for antiphospholipid antibodies

Submission date 04/02/2002	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 04/02/2002	**Overall study status** Completed	
Last Edited 08/10/2013	**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
K0554

Study information

Scientific Title

Acronym
ALIAPAS

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
Prevention of thrombosis

Interventions
Patients will be randomised to either low dose (75 mg) aspirin or low-dose aspirin and low international normalised ratio (INR) warfarin

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Aspirin, warfarin

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/09/2000

Completion date
01/01/2006

Eligibility

Key inclusion criteria
Patients positive for antiphospholipid antibodies

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/09/2000

Date of final enrolment
01/01/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Louise Coote Lupus Unit
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SE1 7EH

Sponsor information

Organisation
Arthritis Research Campaign (ARC) (UK)

Sponsor details
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Sponsor type
Charity

Website
http://www.arc.org.uk

ROR
https://ror.org/02jkpm469

Funder(s)

Funder type
Charity

Funder Name
Arthritis Research Campaign (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/02/2014		Yes	No