

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	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 04/02/2002	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 08/10/2013	Condition category Circulatory System	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Munther Khamashta

Contact details
Louise Coote Lupus Unit
St Thomas' s Hospital
The Rayne Institute
London
United Kingdom
SE1 7EH
+44 (0)20 7928 9292
106404.2325@compuserve.com

Additional identifiers

Protocol serial number

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

Study type(s)

Treatment

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(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/01/2006

Eligibility

Key inclusion criteria

Patients positive for antiphospholipid antibodies

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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

United Kingdom

England

Study participating centre

Louise Coote Lupus Unit

London

United Kingdom

SE1 7EH

Sponsor information

Organisation

Arthritis Research Campaign (ARC) (UK)

ROR

<https://ror.org/02jkpm469>

Funder(s)

Funder type

Charity

Funder Name

Arthritis Research Campaign (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	01/02/2014		Yes	No