ISRCTN85634342 https://doi.org/10.1186/ISRCTN85634342

The effect of prostanoid precursors on platelet angiotensin II binding

Submission date	Recruitment status	[_] Prospectively registered
12/09/2003	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
12/09/2003	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
15/01/2009	Circulatory System	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr PMS O'Brien

Contact details
Obstetrics and Gynaecology Academic Department
Maternity Block
City General
Newcastle Road
Stoke-on-Trent
United Kingdom
ST4 6QG
+44 (0)1782 554998

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0158024607

Study information

Scientific Title

Study objectives
To determine the effect of fish oil dietary supplements and aspirin on platelet angiotensin II binding.

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)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Cardiovascular: Hypertension

Interventions
Normotensive and hypertensive women will be randomly allocated in a double blind manner to one of the four groups: fish oil, aspirin, fish oil and aspirin and placebo.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
fish oil, aspirin

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/06/1994

Completion date
01/06/2004

Eligibility

Key inclusion criteria
128 women in total will be recruited, 32 patients to each of the four treatment arms. Half the women will be hypertensive and the other half normotensive.

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
128

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/06/1994

Date of final enrolment
01/06/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Obstetrics and Gynaecology Academic Department
Stoke-on-Trent
United Kingdom
ST4 6QG

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
Government

Funder Name
North Staffordshire Research and Development Consortium (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	01/01/1999		Yes	No